

Structure–Property Relationships in Unsymmetric Bis(antiaromatics): Who Wins the Battle between Pentalene and Benzocyclobutadiene?[†]

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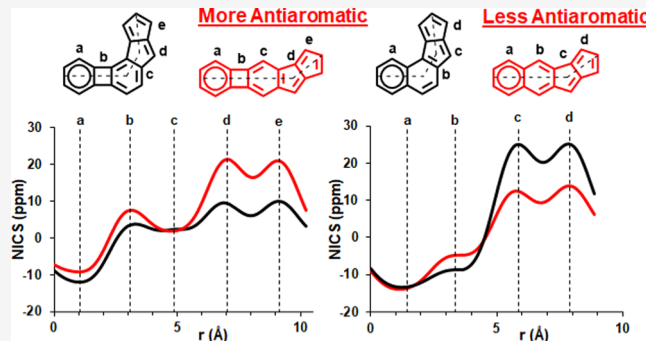
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ABSTRACT: According to the currently accepted structure–property relationships, aceno-pentalenes with an angular shape (fused to the 1,2-bond of the acene) exhibit higher antiaromaticity than those with a linear shape (fused to the 2,3-bond of the acene). To explore and expand the current view, we designed and synthesized molecules where two isomeric, yet, different, 8 π antiaromatic subunits, a benzocyclobutadiene (BCB) and a pentalene, are combined into, respectively, an angular and a linear topology via an unsaturated six-membered ring. The antiaromatic character of the molecules is supported experimentally by ¹H NMR, UV–vis, and cyclic voltammetry measurements and X-ray crystallography. The experimental results are further confirmed by theoretical studies including the calculation of several aromaticity indices (NICS, ACID, HOMA, FLU, MCI). In the case of the angular molecule, double bond-localization within the connecting six-membered ring resulted in reduced antiaromaticity of both the BCB and pentalene subunits, while the linear structure provided a competitive situation for the two unequal [4n] π subunits. We found that in the latter case the BCB unit alleviated its unfavorable antiaromaticity more efficiently, leaving the pentalene with strong antiaromaticity. Thus, a reversed structure–antiaromaticity relationship when compared to aceno-pentalenes was achieved.



INTRODUCTION

Polycyclic conjugated systems that incorporate [4n] π antiaromatic^{1–3} subunits are of increasing interest in contemporary organic material design for small-molecule semiconductors. The rationale behind the application of antiaromatic motifs is their ability to efficiently reduce the aromaticity of acene-type systems while maintaining extended π -conjugation, thus ensuring high charge mobilities with reasonable chemical stability.⁴ In fact, there are recent reports on the successful realization of organic electronic devices (OFETs, solar cells) based on this principle.^{5–11} Antiaromaticity is also considered a design element in the development of supramolecular systems^{12–16} and molecular wires.^{17–21}

Among antiaromatic carbocyclic structures, synthetic efforts are focused mainly on indacene,^{22–31} cyclobutadiene,^{32–40} and pentalene^{41–50} derivatives. Pentalene,^{51,52} as an 8 π molecule, is one of the simplest antiaromatic polycyclic conjugated hydrocarbons, which provides several opportunities for modification and hence different degrees of stabilization that is affecting molecular conductivity, crystallization, or thin-film formation properties.

There are distinct approaches to tune the antiaromaticity of the pentalene unit within its π -extended derivatives (Figure 1).

These include the fusion of heteroaromatic rings^{53–57} on the pentalene core, introduction of substituents^{58–62} with electron-donating or -withdrawing character, and different degrees of π -extension^{63–68} by the fusion of acene-type ring systems. A particularly important aspect within this latter approach is the topology of the fusion to the antiaromatic subunits.

A general pattern that can be recognized from the structure–property studies of these molecules, regardless of if they are monoaryl^{69,70} or diaryl systems,^{71–73} is that the more linear the structure (fusion to the 2,3-bond of the acene), the less antiaromatic becomes the pentalene core. This not only is true for pentalenes but also applies to π -extended cyclobutadienes,³⁹ indacenes,⁷⁴ and pentalene derivatives having N-heteroatoms within the five-membered rings.⁷⁵ Haley and co-workers rationalized this effect in a combined

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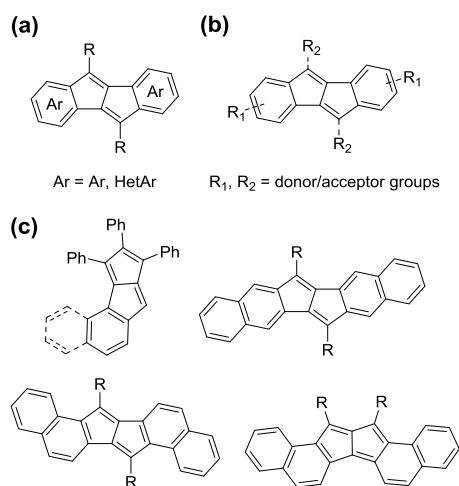


Figure 1. General strategies to tune the antiaromaticity of pentalenes. (a) Fusion of aryl/heteroaryl rings; (b) introduction of donor/acceptor substituents either on the pentalene core or on the fused aryl ring; and (c) variation of the fusion pattern around the pentalene core in aceno-pentalene derivatives.

experimental and theoretical study of these compounds.⁷⁴ They found a correlation between the bond order of the fused aromatics and the antiaromaticity of the resulting π -extended $[4n]\pi$ structure. Fusion to the bond with a higher bond order led to increased antiaromaticity and vice versa (Figure 2a).

Based on the above analysis, it was predicted that this correlation may be general to all diareno-fused antiaromatic systems.⁷⁴ With regard to the geometric structure of the resulting molecules, this implies that—in the case of the fusion

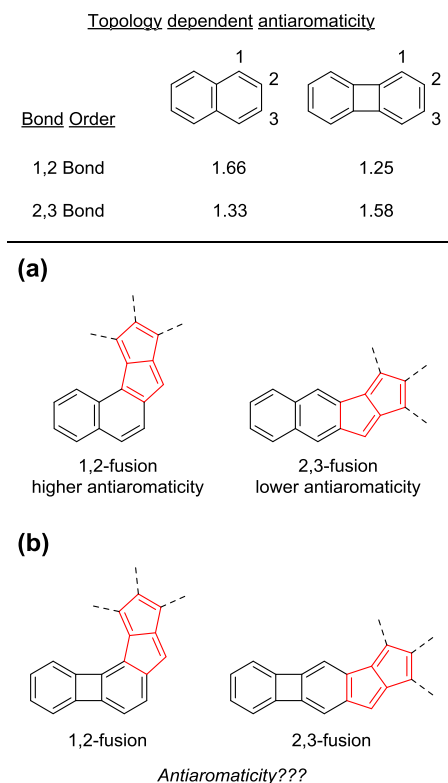


Figure 2. Role of bond order in the topology-dependent antiaromaticity of fused pentalenes. (a) Naphtho-pentalenes; (b) biphenyleno-pentalenes.

of regular acenes to antiaromatic units—the molecules that are less antiaromatic will be more linear (2,3-fusion), while those with higher antiaromaticity will be more angular (1,2-fusion). This pattern leads to a serious limitation among aceno-antiaromatics as it restricts distinct electronic properties to distinct molecular topologies. It is well-known that molecular topology greatly affects the solid-state properties of the molecules, and this strongly influences their efficiency in device applications.^{4,76,77} With regard to the correlation between the acene bond-order and antiaromaticity, this could mean that the electronically more interesting highly antiaromatic systems could be of limited use in device applications due to their disadvantageous geometric structures. Thus, the question arises whether the correlation between the topology of a molecule and its antiaromaticity can be reversed by a different approach to the molecular design.

As approaches to reverse the relationship between molecular shape and antiaromaticity have not been explored previously, our goal was to deviate from the expected low antiaromaticity in the case of linearly fused acenes and to construct π -systems with a linear shape that maintain high antiaromaticity. A key feature of our design was the identification of a system to be fused with the pentalene unit having reversed bond orders of the 1,2- and 2,3-bonds when compared to the previously used acenes. We turned to biphenylene, a 12π cyclobutadiene derivative, that has been described as antiaromatic, although the description of its aromaticity and the destabilizing effect of its cyclobutadiene subunit are not straightforward.^{78–81} In this molecule, to decrease the cyclobutadiene character, there is considerable double-bond localization that leads to lower bond order at the 1,2-bond compared to that of the 2,3-bond (Figure 2b).⁸² This is the opposite trend to what is present in acenes such as naphthalene or anthracene. Hence, we argue that π -extended pentalenes with a linear shape and preserved antiaromaticity could be prepared by the fusion with biphenylene. From a structural point, the proposed molecules could be considered as the linking of isomeric benzocyclobutadiene (BCB) and pentalene, having different levels of antiaromaticity, through an unsaturated six-membered ring in different topologies (Figure 3).

In this regard, they are isoelectronic to the recently reported bis-pentalenes, where two pentalene moieties were fused to a central benzene ring.⁸³ These latter systems were dominated by the two pentalene subunits, which exhibited an antiaromatic character. Thus, the reversed structure–property relationship was expected by the fusion of biphenylene instead of acenes to pentalene. The key question addressed herein is which one of the two unequally antiaromatic subunits alleviates its antiaromaticity to the most extent; in other words, who wins the battle, BCB or pentalene?⁸⁴

RESULTS AND DISCUSSION

We have synthesized two monoannulated biphenyleno-pentalenes **9** and **14** having an angular and a linear topology, respectively. These molecules were studied experimentally by ¹H NMR spectroscopy, X-ray crystallography, UV–vis spectroscopy, and cyclic voltammetry, and the computational analysis of their aromaticity was performed (NICS-XY scan, ACID, HOMA, FLU, MCI). Throughout the manuscript, we compared the properties of the newly prepared molecules **9** and **14** to those of the previously reported naphtho-pentalene derivatives **15** and **16** with similar topologies.⁶⁹

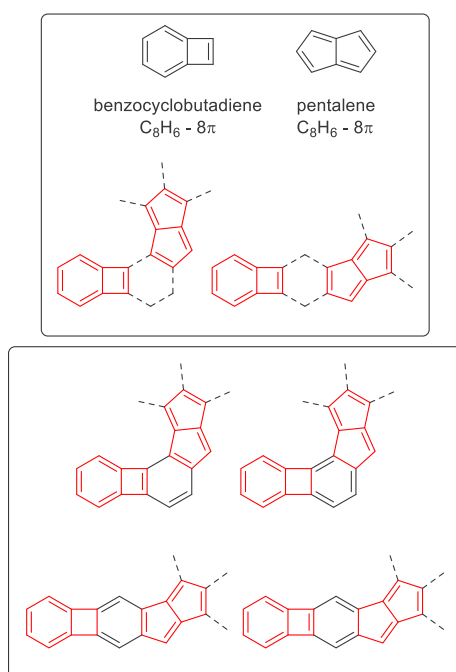


Figure 3. Combination of benzocyclobutadiene (BCB) and pentalene with different topologies.

Synthesis. We have synthesized two biphenylene fused monoannulated pentalenes with an angular (**9**) (Scheme 1) and a linear (**14**) (Scheme 2) topology. In the angular structure, the pentalene moiety is fused to the 1,2-bond of biphenylene, while in the linear case, it is fused to the 2,3-bond. Both synthesis sequences relied on the modifications of biphenylene (**2**) earlier explored by McOmie and co-workers.^{85,86}

Biphenylene itself was prepared from anthranilic acid (**1**) via a benzyne intermediate with varying yields (27–32%), due to the instability of the diazonium salt generated from **1**. Formylation of **2** was performed selectively using dichloro-

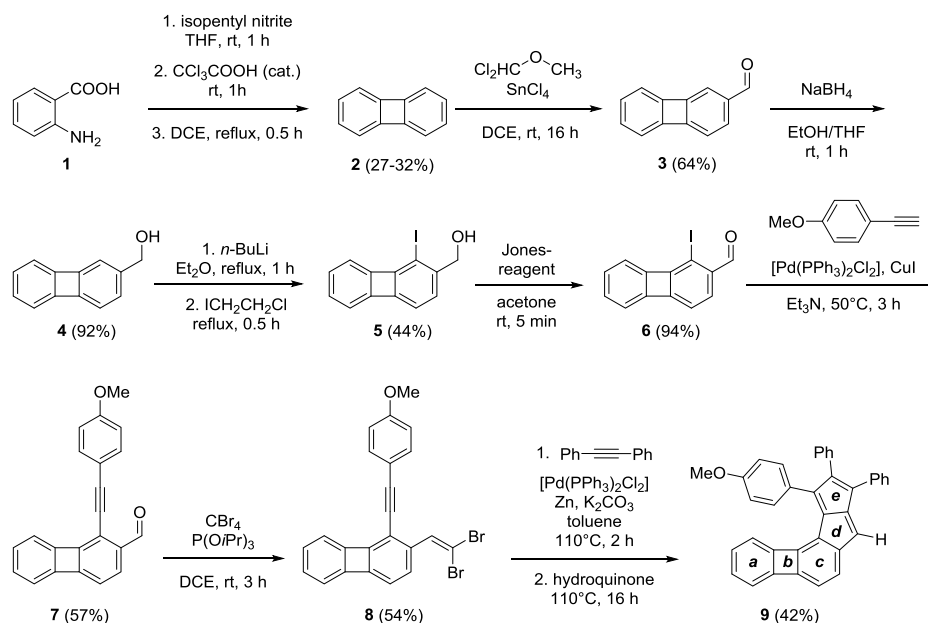
omethyl methyl ether in the presence of SnCl_4 . The reduction of the formyl group of compound **3** to the corresponding alcohol **4** by NaBH_4 was necessary for the selective iodination in the following step via the *ortho*-lithiation/iodination sequence. The benzylic alcohol group of iodinated derivative **5** was oxidized back to the corresponding aldehyde **6** using the Jones reagent. Subsequent Sonogashira coupling with 4-methoxyphenylacetylene, yielding **7**, and *gem*-dibromoolefin formation provided the key intermediate **8** for the pentalene formation. The key step for the synthesis of the pentalene unit was a Pd-catalyzed cascade carbopalladation reaction between alkynes and *gem*-dibromoolefins pioneered by Diederich and co-workers.^{43,69} The carbopalladation cascade provided the angular biphenylene-pentalene **9** in 42% yield.

For the synthesis of linear biphenylene-pentalene (Scheme 2), compound **2** was first brominated using *N*-bromosuccinimide (NBS) in *N,N*-dimethylformamide (DMF), yielding bromobiphenylene **10**, which was formylated using the dichloromethyl methyl ether/ SnCl_4 system to obtain aldehyde **11**. The low yield obtained for compound **11** is due to the low selectivity of the reaction. Subsequent Sonogashira coupling with 4-methoxyphenylacetylene, yielding **12**, and *gem*-dibromoolefin formation provided the key intermediate **13** for the pentalene formation. The carbopalladation cascade provided the linear biphenylene-pentalene **14** in 27% yield.

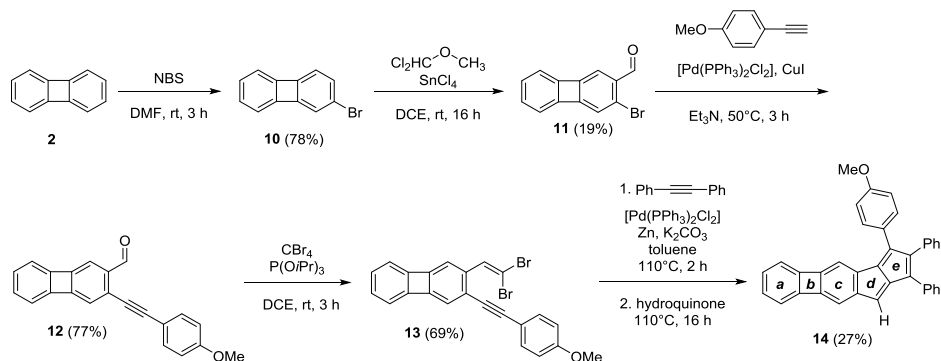
Based on previous examples,⁸³ the introduction of a methoxy group in the dibromoolefins is advantageous as it increased the yield of the cascade reaction and facilitated the purification of the products. Compound **9** is a deep-purple bench-stable compound, while the orange-brown compound **14** was found to degrade under slightly acidic conditions (during silica column chromatography and to some extent in CDCl_3 ; for measurements, solvents treated with basic alumina were used).

¹H NMR Spectroscopy. The ability to sustain diatropic and paratropic ring currents is a characteristic of aromaticity and antiaromaticity, respectively.^{2,87–89} Such ring current effects are reflected in the proton chemical shifts of aromatic

Scheme 1. Synthesis of Biphenylene-pentalene **9** with an Angular Topology



Scheme 2. Synthesis of Biphenyleno-pentalene 14 with a Linear Topology



and antiaromatic compounds, which could be used as indicators of the extent of their aromaticity or antiaromaticity. It has been shown recently that $[4n]\pi$ subunits can sustain paratropicity within larger conjugated frameworks.⁸³ Because the molecules presented here also comprise two different 8π antiaromatic subunits, a BCB and a pentalene, their ^1H NMR spectra are a good approximation of whether antiaromaticity is preserved in their structure (Figure 4a).

The pentalene protons in both molecules showed the characteristic upfield shift that has been observed for monoaryl-pentalenes earlier.^{43,69} The absorption of the pentalene proton (H_1 , Figure 4a) of the angular compound **9** appears at 6.60 ppm, while in the linear compound **14**, this signal appears at 6.02 ppm in CDCl_3 (protons were assigned by the 1D-NOESY technique; see Section S6, Supporting Information; for spectra recorded in CD_2Cl_2 , see Section S6, Supporting Information). As the potential shielding effect by the proximal pendant phenyl substituents is expected to be the same in both cases, the upfield shifts originate from the interplay of the remaining diatropic ring current in ring *c* and the paratropic ring current in the pentalene subunit. The results suggest that the antiaromaticity of the pentalene subunit in the linear structure is more preserved.

Comparing these chemical shifts to the pentalene protons of the corresponding naphtho-pentalenes **15** and **16** that have been reported earlier,⁶⁹ the opposite trend was found (Figure 4b). In these latter cases, the pentalene proton of the angular structure **15** appears at 6.11 ppm, while in the linear structure **16**, it is at 6.71 ppm, showing lower antiaromaticity of the linear structure in this case. As a further comparison, the chemical shift of the pentalene proton in the related monobenzo-pentalene structure **17** is 6.28 ppm (Figure 4b).⁴³ Note that the effect of the substituent on the peripheral phenyl groups on the chemical shifts is negligible, as described earlier.⁸³ The chemical shifts of the protons on the six-membered rings between the two antiaromatic subunits also appear in the alkene region, which points toward the decreased aromaticity of this ring in both cases (H^2 and H^3 , Figure 4a).

An extensive comparison of the ^1H NMR shifts is difficult as only a few examples of diareno-pentalenes with unsubstituted pentalene rings have been reported. Still, the pentalene protons of dibenzo[*a,e*]pentalene appear at 6.40 ppm,⁹⁰ while this value for diareno-pentalenes with further π -extension increases up to around 7 ppm, showing diminishing antiaromaticity in these cases.⁶⁷

X-ray Crystallography. The topology of the aryl-fusion around the pentalene core has been shown to greatly influence the antiaromaticity and hence the optical and electronic

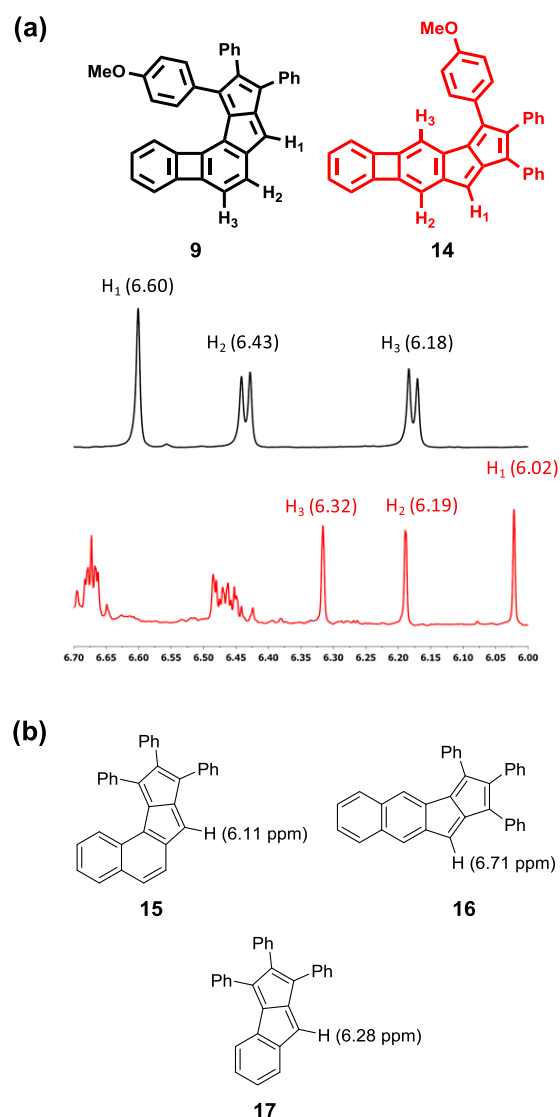


Figure 4. (a) Partial ^1H NMR spectra of compounds **9** and **14** (CDCl_3 , 500 MHz, room temperature (rt)). Protons were assigned by the one-dimensional-nuclear over-Hauser effect spectroscopy (1D-NOESY) technique. (b) Reported monoaryl pentalene structures for comparison of the chemical shifts of the pentalene proton (all measured in CDCl_3).

properties of the molecules in the case of both monoareno- and diareno-pentalenes.^{69–73} The general observation for naphthalene- and anthracene-fused systems is that the 1,2-

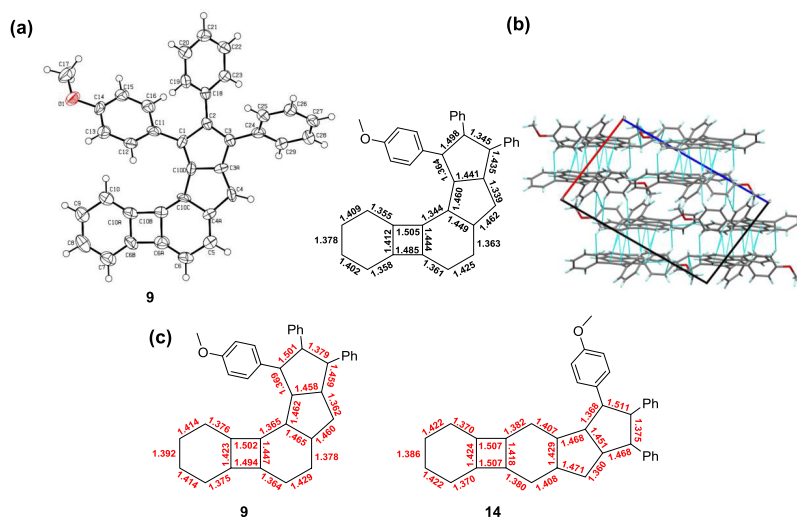


Figure 5. (a) X-ray structure of compound **9** and the corresponding bond lengths; ORTEP representation of **9** is drawn at the 50% probability level; (b) layered structure of compound **9** in the crystalline state presented from the view of the crystallographic *b*-axis. Blue lines represent intermolecular short contacts (\leq sum of van der Waals radii + 0.1 Å). (c) Calculated bond lengths of compounds **9** and **14**.

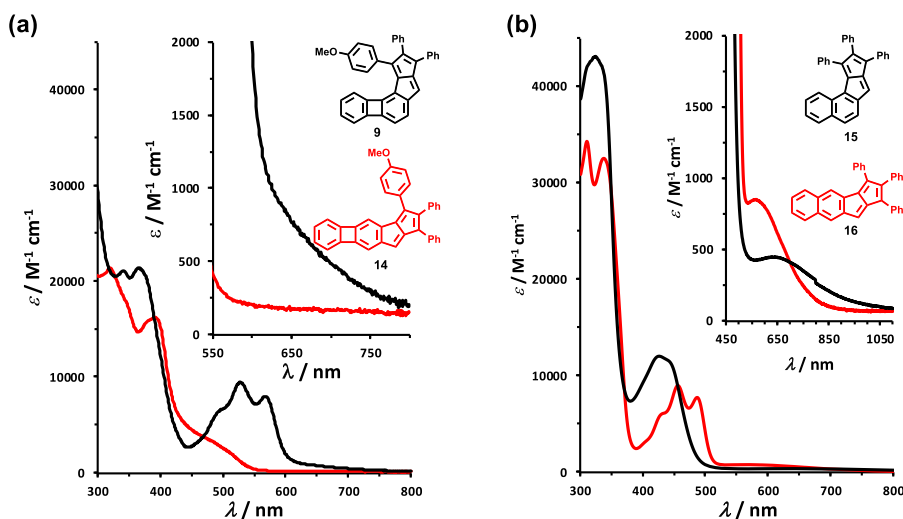


Figure 6. (a) UV–vis spectra of compounds **9** and **14** (in CHCl_3); (b) UV–vis spectra of naphtho-pentalenes **15** and **16** (in CHCl_3).

fused (angular) areno-pentalenes are better preserving their antiaromaticity compared to the 2,3-fused (linear) systems. We have examined the synthesized biphenyleno-pentalene molecules in this context, as the behavior of the six-membered ring that is confined between two antiaromatic units (BCB and pentalene) was expected to be particularly interesting and decisive for the extent of antiaromaticity of the subunits.

We could successfully grow single crystals of the angular compound **9** suitable for X-ray crystallographic analysis (Figure S1a) (for further details, see Section S1, Supporting Information). Significant bond-length alteration was found through the periphery of the conjugated core. The calculated bond lengths (Figure S1c) are in agreement with those determined experimentally. It is interesting to note that the polycyclic core of the molecule is not completely planar, and it is slightly out of plane around the four-membered ring (C10–C10A–C10B–C10C dihedral angle is 9.86°). In the calculated structure of compound **9**, the same dihedral angle is about 1.5°, while the structure without the phenyl groups is planar. Hence, the observed effect could be attributed partially to intramolecular crowding and partially to forces in the crystal.

The crystal structure reveals that double bond-localization occurs within the six-membered ring between the two antiaromatic subunits in such a manner that decreases the bond order at the four- and five-membered ring fusion. By this localization, the antiaromatic characters of both the BCB and pentalene subunits are decreased. Molecules in the crystals are held together by multiple weak secondary interactions that lead to a layered structure (Figure 5b).

Unfortunately, we were not able to crystallize the linear derivative **14**, but for comparison, we calculated the bond lengths within the molecule (Figure S5c). Calculations predict bond length alteration in this case as well. Importantly, the predicted bond length at the pentalene/six-membered ring fusion is shorter than that in compound **9** (1.429 and 1.465 nm, respectively), indicating higher bond order at this bond in **14**.

Opto-electronic Properties. UV–vis spectra of compounds **9** and **14** were recorded in CHCl₃ (Figure 6a) and compared to those of compounds **15** and **16** (Figure 6b).⁶⁹ Spectra recorded in alternative solvents (CH₂Cl₂ and tetrahydrofuran (THF)) showed no pronounced differences

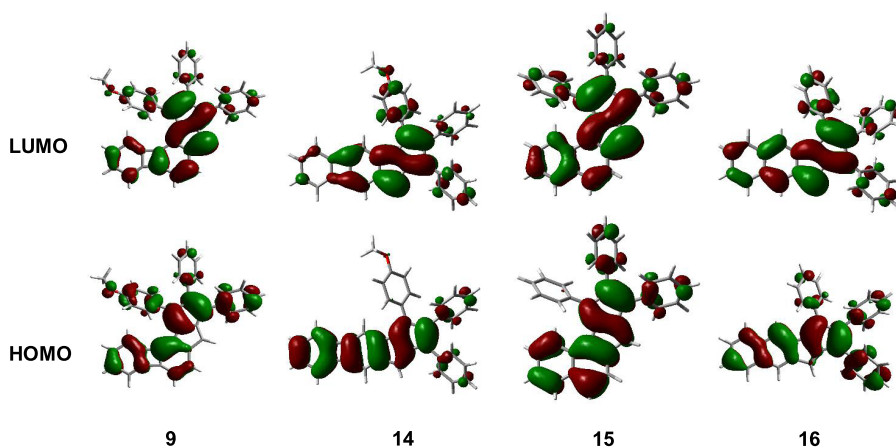


Figure 7. Calculated HOMO and LUMO orbitals of biphenyleno-pentalenes **9** and **14** and naphtho-pentalenes **15** and **16**.

(see Section S2, Supporting Information). It is immediately apparent that the shape of the absorption bands between 400 and 600 nm is reversed within the two compound pairs. In the case of the more antiaromatic angular naphtho-pentalene **15**, a broad absorption band appears with a maximum at 426 nm, while in the case of the less antiaromatic linear compound **16**, there are two distinct maxima (at 456 and 486 nm) giving a more structured absorption profile in the same region. In the case of the biphenyleno-pentalenes, however, structure **9** having an angular topology exhibits the absorption profile that resembles that of linear naphtho-pentalene **16**. The two distinct absorption maxima in the case of compound **9** are bathochromically shifted to 527 and 568 nm. The spectrum of compound **14** is characterized by a maximum at 391 nm and a broad shoulder between 430 and 550 nm. In the spectra of compounds **15** and **16**, low-energy long-wavelength absorptions that are characteristic of the symmetry-forbidden highest occupied molecular orbital (HOMO) \rightarrow lowest unoccupied molecular orbital (LUMO) transitions for pentalenes are clearly present ($\lambda_{\text{max}} = 660$ nm for **15** and 566 nm for **16**). These absorptions correspond to HOMO \rightarrow LUMO transitions of 1.84 and 2.19 eV, respectively. As no clear low-energy absorptions could be identified in the spectra of **9** and **14**, we performed time-dependent density functional theory (TD-DFT) calculations (B3LYP/6-311+G(d,p) level of theory) to gain more insight into the electronic properties and transition energies of these compounds.

In the case of angular compound **9**, the computed HOMO–LUMO gap was found to be 2.11 eV ($\lambda = 590$ nm, $f = 0.016$), which suggests that the corresponding absorption is overlapping with the shoulder in the region above the 568 nm maximum. In the case of linear compound **14**, the transition energy is calculated to 1.61 eV ($\lambda = 770$ nm, $f = 0.0004$). These results could be rationalized by the analysis of the frontier molecular orbitals of the compounds (Figure 7). In the case of compound **14**, both the HOMO and the LUMO possess an S-shaped geometry that is responsible for the symmetry-forbidden nature of this transition and explains the lack of the corresponding absorption in its UV–vis spectrum. Compared to the S-shaped geometry of the HOMO in compound **14**, the HOMO of compound **9** is distorted; hence, there is a comparably increased absorption in the region where the HOMO \rightarrow LUMO transition is expected to occur.

Similar to the trend in ^1H NMR shifts and UV–vis absorptions, in the case of the orbital features, a reversal of the

patterns can be observed when comparing naphtho-pentalenes and biphenyleno-pentalenes. While the S-shape of the frontier orbitals and hence the lower absorption intensity that corresponds to the HOMO \rightarrow LUMO transition are characteristic of the angular compound **15**, in the case of biphenyleno-pentalenes, these features are associated with the linear structure of **14**. On the other hand, distorted symmetry and thus higher-intensity UV–vis absorptions that correspond to HOMO \rightarrow LUMO transitions are characteristic of both the linear naphtho-pentalene **16** and the angular biphenyleno-pentalene **9**.

The frontier orbitals were also calculated for the antiaromatic subunits BCB and pentalene and for those units having exo-double bonds (Figure 8). These substructures are expected to have a contribution to the overall structures of molecules **9** and **14**.

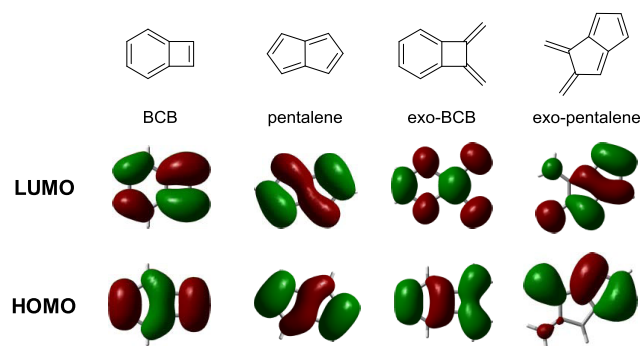


Figure 8. Calculated HOMO and LUMO orbitals of BCB, pentalene, exo-BCB, and exo-pentalene.

In line with the experimentally determined bond lengths from the crystal structure of compound **9**, the fulvene-like exo-pentalene structure was found to have a strong contribution to the HOMO of this molecule. The exo-double bonds are part of the adjacent six-membered ring that exhibits pronounced double bond-localization. Contrary to compound **9**, where this bond localization led to the formation of an exo-pentalene substructure, in the HOMO of compound **14**, the contribution of BCB and pentalene is clearly present. Furthermore, the LUMO of both substructures can be recognized in the LUMO of **14**. These results suggest that in the linear compound **4**, antiaromaticity is more significantly present than in the angular molecule **9**.

Table 1. Summary of Electrochemical, Optical, and Computational Data for Compounds 9, 14, 15, and 16

entry	compound	E_{1ox} [V] ^a	E_{1red} [V] ^a	HOMO [eV] ^b	LUMO [eV] ^b	ΔE_{redox} [eV] ^c	ΔE_{opt} [eV]	ΔE_{calc} [eV] ^d
1	9	0.55 ^e	−1.47 ^f	−5.35	−3.33	2.02		2.11
2	14	0.25 ^e	−1.32 ^f	−5.05	−3.48	1.57		1.61
3	15 ⁶⁹	0.44 ^e	−1.44 ^e	−5.24	−3.36	1.88	1.84	1.85
4	16 ⁶⁹	0.60 ^e	−1.53 ^f	−5.46	−3.27	2.13	2.19	2.15

^aElectrochemical measurements were carried out in 0.1 M Bu₄NPF₆ in dichloromethane (DCM) at a scan rate of 0.1 V s^{−1} on a platinum wire working electrode. All potentials are given versus the Fc/Fc⁺ couple used as the internal standard. ^bHOMO and LUMO energy levels in electron volt were approximated using the equation HOMO = −(4.80 + E_{1ox}), LUMO = −(4.80 + E_{1red}).^{91,92} ^c ΔE_{redox} = LUMO − HOMO. ^dCalculations were performed on the B3LYP/6-311+G(d,p) level of theory. ^eReversible first reduction or oxidation wave. ^fIrreversible first reduction or oxidation wave.

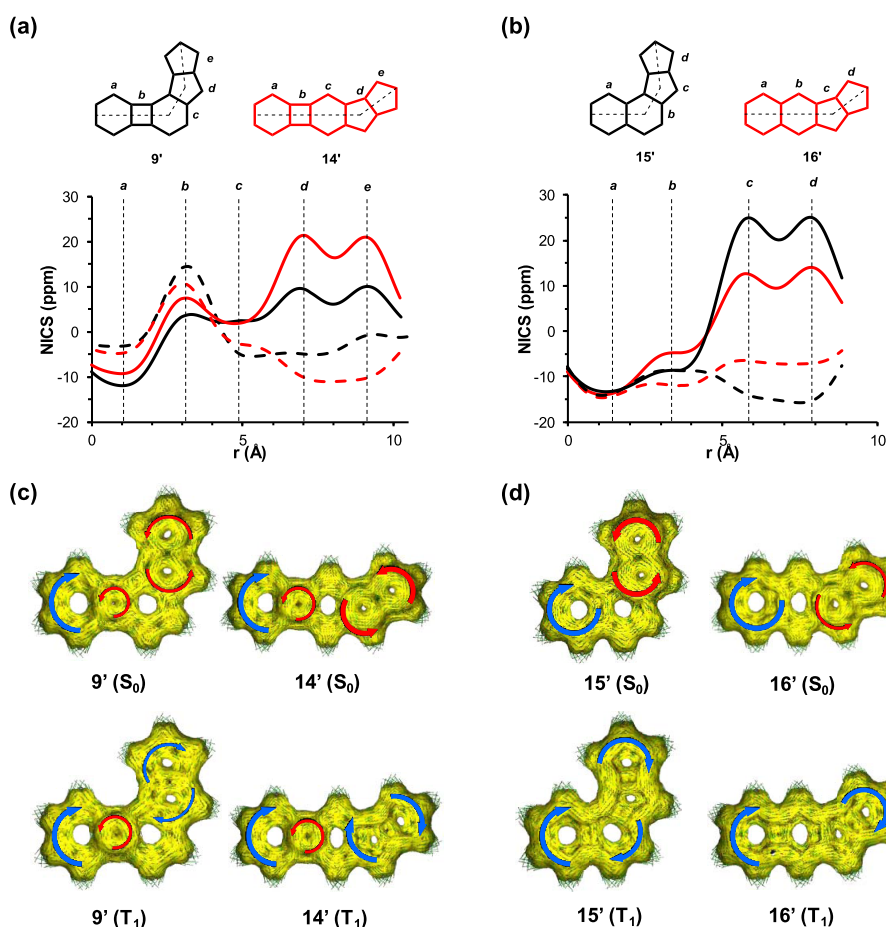


Figure 9. (a) NICS-XY scans of biphenyleno-pentalenes 9' and 14' in the S₀ (solid line) and T₁ (dashed line) states. (b) NICS-XY scans of naphtho-pentalenes 15' and 16' in the S₀ (solid line) and T₁ (dashed line) states. (c) ACID plots of biphenyleno-pentalenes 9' and 14' in the S₀ and T₁ states. (d) ACID plots of naphtho-pentalenes 15' and 16' in the S₀ and T₁ states. (For higher-resolution images, see Section S4.2.1, Supporting Information.) Blue and red arrows correspond to diatropicity and paratropicity, respectively. The width of the arrow denotes the strength of the ring current.

Although the optical HOMO–LUMO gaps could not be determined experimentally from the UV–vis absorptions, we could obtain experimental support for the calculated energy differences by cyclic voltammetry (CV) measurements (Table 1) (see Section S3, Supporting Information). Both compounds 9 and 14 exhibited first reversible oxidation at 0.55 and 0.25 V and irreversible reduction at −1.47 and −1.32 V, respectively. The electrochemical gap is approximative due to the irreversible nature of the reductions; however, the obtained values of 2.02 V for compound 9 and 1.57 V for compound 14 are in good agreement with the theoretically predicted HOMO–LUMO gaps.

The calculated SOMO features of the radical cation and anion for both biphenyleno-pentalene isomers suggest that the unpaired electrons in the cationic and anionic species are delocalized over the system; however, in the anion, a pronounced pentalene character is present (see Section S4.5, Supporting Information). That may lead to further reactions, resulting in the irreversibility of the reduction process.

Aromaticity Analyses. As described above, substantial differences among compounds 9, 14, 15, and 16 were observed both in their HOMO–LUMO transitions obtained from UV–vis absorption spectroscopy and in their HOMO–LUMO energy gaps provided by electrochemical measurements. Low-intensity absorptions that correspond to the symmetry-

forbidden HOMO \rightarrow LUMO transitions are found in the UV–vis spectra of **14** and **15**, while absorptions with increased intensity for the same transitions are detected for **9** and **16** (Figure 6). Below, we show that the associated changes in energies can be linked to aromaticity changes between the singlet (S_0) ground state and the first excited states (T_1 , S_1). At this point, it should be noted that the S_1 and T_1 states for all four compounds are similar in terms of configurations as the excitations according to TD-DFT calculations mainly are described by singly excited HOMO to LUMO ($\pi\pi^*$) configurations.⁹³ As the two states only differ in multiplicity, we explored the T_1 state as this state is more straightforward computationally than the S_1 state (for comparative MCI and FLU values for the S_1 state, see Tables S9 and S11, Supporting Information). Additionally, the trend in HOMO–LUMO gaps matches that of vertical excitations to the T_1 state. To explore if the experimentally observed HOMO–LUMO gap variations for **9**, **14**, **15**, and **16** can be linked to (anti)aromaticity changes upon excitation, we first analyzed their (anti)aromatic character in the S_0 states and then compared with the T_1 states.

Simplified structures where the Ph substituents of all molecules were replaced by H-atoms (denoted **9'**, **14'**, **15'**, and **16'**) were computed. This simplification does not considerably alter the findings (for further details, see Section S4.2, Supporting Information). All geometry optimizations were made with the Gaussian 16 package using the B3LYP hybrid functional and the 6-311+G(d,p) basis set.^{94–96}

The aromatic character has been evaluated by means of magnetic (ACID plots and NICS-XY scans)^{97–102} and geometric (HOMA)^{103–105} indices^{106,107} in their ground (S_0) and excited (T_1) states, computed at the (U)B3LYP/6-311+G(d,p) level (for further details, see Section S4.1, Supporting Information). Aromatic rings are characterized by (i) clockwise ring currents revealed through ACID plots, (ii) diatropic ring currents detected by negative NICS values, and (iii) HOMA values in the range 0.5–1.0. On the other hand, anticlockwise ring currents, positive NICS, and negative HOMA values are indicative of antiaromaticity.

Here, we briefly point out some critical aspects of the usage of different indices to assess the (anti)aromatic character of a molecule. The usage of NICS alone, without analysis of the ring currents, is not recommendable.¹⁰⁸ With regard to HOMA, there is recent criticism regarding the usage of this index for polycyclic aromatic hydrocarbons as it was found that HOMA values of internal six-membered rings in such hydrocarbons are overestimated.¹⁰⁹ Still, the broad applicability of the HOMA index also for polycyclic systems is well documented.¹⁰⁵ We focus the aromaticity analysis on the BCB and pentalene subunits, realizing that this is a simplification as other circuits in the polycyclic compounds also will contribute to the (anti)aromatic character of the molecules. Yet, the BCB and pentalene substructures play significant roles in determining the electronic character of the molecules as shown in the previous Experimental Section on the properties of **9** and **14**. To obtain a qualitative view, we therefore focus on the BCB and pentalene subunits.

The NICS-XY scans of molecules **9'** and **14'** in their S_0 state (Figure 9a) highlight that within these structures, the patterns that are characteristic of BCB and pentalene (see Figure S9, Supporting Information) to a good extent are preserved, even though the antiaromatic characters of the substructures that correspond to BCB (circuit *a+b*) and pentalene (circuit *d+e*) are clearly attenuated. Yet, it is important to note that the

NICS values of **9'** throughout the scan are lower than those of **14'**, which agrees with the lower antiaromaticity of the angular topology determined experimentally by ¹H NMR spectroscopy. It is particularly noteworthy that the NICS values in the *d* + *e* subunit of **9'** are significantly lower than those within the same substructure of **14'**. The ACID plots of **9'** and **14'** in their S_0 states (Figure 9c) further support this observation. The opposite trend is obtained for molecules **15'** and **16'** in their NICS values and ACID plots regarding their pentalene (*c+d*) subunits (Figure 9b). In these cases, structure **15'** having an angular topology exhibits higher antiaromaticity in its *c+d* subunit compared to that of **16'** with a linear topology. We also calculated the relative energies of BCB and pentalene and found that the latter is more stable by approximately 6 kcal/mol. Complementary to this, as expected, the calculation of the relative energies of the corresponding exo-BCB and exo-pentalene structures shows the opposite, exo-BCB being more stable by about 4.6 kcal/mol. This supports the preferred rearrangement of the BCB unit to the exo-BCB structure in molecule **14**. Furthermore, the interaction of BCB and pentalene subunits in **9'** and **14'** is clear from the comparison of the NICS-XY scans of **9'**, **14'**, and benzopentalene (**17'**) (see Section S4.4, Supporting Information). The alleviated antiaromaticity of BCB in **14'** enforces a stronger pentalene antiaromaticity in **14'** compared to that in benzopentalene. Furthermore, the remaining aromaticity of the six-membered ring *c* is higher in benzopentalene as it lacks the BCB subunit, which significantly decreases the aromaticity of ring *c* in molecules **9'** and **14'**. Similarly, the comparison of **9'**, **14'**, and the corresponding compounds that lack the terminal benzene rings (“cyclobutadieno-benzopentalenes”, structures **S4** and **S5**, Section S4.4, Supporting Information) suggests that the electronic structures of **9'** and **14'** are dominated by the interaction of the BCB and pentalene subunits rather than the cyclobutadiene and pentalene subunits.

The geometry-based HOMA values for the BCB and pentalene substructures in **9'** and **14'** (Table 2) are in line

Table 2. Geometric (HOMA) Aromaticity Indices Calculated for the BCB and Pentalene Subunits in Their S_0 and T_1 States

compound/subunit	HOMA- S_0	HOMA- T_1
BCB	−0.460	0.568
pentalene	−0.388	0.820
<i>a+b</i> (9')	0.073	0.085
<i>d+e</i> (9')	0.004	0.339
<i>a+b</i> (14')	−0.050	0.169
<i>d+e</i> (14')	−0.123	0.606
<i>c+d</i> (15')	−0.150	0.673
<i>c+d</i> (16')	−0.036	0.379

with the magnetic indices. The HOMA values of both the BCB and the pentalene unit in **9'** reflect its lower antiaromaticity when compared to **14'**, and this agrees with the crystallographically as well as the computationally determined bond-length alterations (see Figures S7 and S8, Supporting Information). That the pentalene subunits of **9'** and **14'** are the main antiaromatic subunits, rather than the BCB subunit, becomes clear when analyzing the difference of aromaticity going from the two units as separate molecules to subunits in **9'** and **14'**. The perimeter of BCB is slightly more antiaromatic (HOMA = −0.460) than that of pentalene (HOMA =

−0.388). When the two molecules are combined to form compounds **14'** and **9'**, HOMA reveals a reduction of the antiaromatic character of both subunits, but the alleviation is larger in the BCB than in the pentalene subunit ($\Delta\text{HOMA} = 0.53$ and 0.41 versus 0.38 and 0.26 , respectively). Moreover, the contribution of BCB in **14'** to LUMO is very small compared to **9'** (Figure 7) as the BCB subunit efficiently alleviates its antiaromaticity contrary to the pentalene subunit (see Tables S14 and S15, Supporting Information). In agreement with the magnetic indices, HOMA values calculated for the *c+d* subunits of structures **15'** and **16'** (see Figure S6, Supporting Information) show the opposite structure–antiaromaticity relationship, as molecule **15'** with an angular topology maintains a higher antiaromaticity when compared to the linear structure **16'**.

Despite the fact that the bicyclic *a+b* and *d+e* moieties are substructures within molecules **9'** and **14'**, it is interesting to compare their HOMA values with those of BCB and pentalene alone (Table 2). In their S_0 states, the HOMA values of perimeters of BCB and pentalene are −0.460 and −0.388, respectively, revealing clear antiaromaticity. Upon connecting these units in an angular fashion, as in **9'**, these values change to 0.073 (*a+b*) and 0.004 (*d+e*), indicating nonaromatic circuits. Such a decrease in the antiaromatic character can be explained as a consequence of the strong double bond localization in ring *c* as seen in the crystal structure of **9**, leading to simultaneous antiaromaticity alleviations in both the BCB and pentalene units. In contrast, when BCB and pentalene are fused in a linear fashion, as in **14'**, similar simultaneous antiaromaticity alleviations of the BCB and pentalene units are impossible. According to the HOMA values of **14'**, the pentalene unit retains a slightly more antiaromatic character than the BCB unit and the change is larger for the perimeter of the *a+b* moiety ($\Delta\text{HOMA} = 0.410$) than for *d+e* ($\Delta\text{HOMA} = 0.265$). As a consequence, the linear compound **14'** displays an antiaromatic character in the pentalene moiety.

As noted above, **9**, **14**, **15**, and **16** display marked differences in their HOMO–LUMO transitions determined via UV–vis absorption spectroscopy and electrochemical measurements. We now explored if there is a link between the HOMO–LUMO gaps and the (anti)aromaticity changes between the singlet ground state and the first excited state (T_1).^{110–115} We analyzed the T_1 state instead of the S_1 state for reasons described above. Moreover, we present NICS-XY, ACID, and HOMA results for the T_1 states as these are the most readily calculated, while FLU and MCI data for the S_1 state are given in the Supporting Information (Tables S9 and S11, respectively). The aromaticity indices corresponding to the BCB and pentalene subunits in **9'** and **14'** in their T_1 state (Table 2) can potentially be connected to the ground-state characteristics of these molecules.

Upon excitation of **9'** and **14'** to their T_1 states, there is a marked change in the (anti)aromaticity character of the pentalene subunit of **14'** as it clearly becomes Baird-aromatic according to NICS-XY, ACID, as well as HOMA (Figure 9 and Table 2). Thus, the strong antiaromatic character localized to the pentalene subunit in the S_0 state of **14'** switches into a considerable aromatic character in the T_1 state. In this latter case, both experimental data and computational analyses in the S_0 state confirm higher antiaromaticity, especially in the pentalene subunit. Hence, preserved pentalene antiaromaticity is the price for an alleviated BCB antiaromaticity. This ground-state situation is well reflected in the T_1 features of **14'**. Similar

to that in **9'**, the nonaromatic BCB subunit has a low contribution to T_1 aromaticity of compound **14'**.

Overall, compound **9'** exhibits a relatively low antiaromaticity in the S_0 state and low aromaticity in the T_1 state, while the pentalene unit in compound **14'** has an antiaromatic character in the S_0 state and an aromatic character in the T_1 state. Hence, the relative (anti)aromaticities in the S_0 and T_1 states of these compounds, revealed by NICS-XY, ACID, and HOMA, indicate a larger ground-state destabilization and an enhanced aromatic stabilization in the T_1 state of **14'**, leading to a $\Delta E(T_1-S_0)$ of 0.51 eV for **14'** and 0.85 eV for **9'**. This is also in agreement with the experimentally determined as well as the calculated lower HOMO–LUMO gap of **14'** compared to **9'** (Table 1).

Both the previously reported experimental data⁶⁹ and the calculated aromaticity indices (Figure 9 and Table 2) confirm the stronger antiaromaticity of the pentalene subunit in angular naphtho-pentalene **15'** compared to linear **16'**, where the double bond pattern in the fused five-membered rings rather defines a fulvene-like system (just as in **9'**). Importantly, in line with the experimental findings, all of the indices confirm a similarly high antiaromaticity and Baird aromaticity of the pentalene subunit within the linear biphenylene-pentalene **14'** and the angular naphtho-pentalene **15'**.

CONCLUSIONS

We have shown that the generally accepted correlation between fusion pattern and antiaromaticity among π -extended pentalenes (aceno-pentalenes) could be altered by extending the molecular design to unsymmetric bis(antiaromatic) systems. There are two key features in the design of the newly synthesized molecules, which are composed of a biphenylene and a pentalene substructure. On the one hand, the bond-order values of the 1,2- and 2,3-bonds in biphenylene, to which the pentalene unit is fused, are reversed compared to those in acenes. On the other hand, the molecules can be considered as the mergers of two different subunits with an unequal antiaromatic character: a benzocyclobutadiene (BCB), contained in the biphenylene substructure, and a pentalene. When these subunits were fused in an angular fashion (through the 1,2-bond of biphenylene, as in compound **9**), bond localization in the confined six-membered ring minimized both the BCB and the pentalene character in the molecule as supported by ¹H NMR studies, X-ray crystal structure, and computational aromaticity analyses (NICS, ACID, HOMA). However, when BCB and pentalene were combined in a linear fashion (through the 2,3-bond of biphenylene, as in compound **14**), there was no possibility for both subunits to alleviate their aromaticity. In this case, the BCB subunit alleviated its antiaromaticity to a greater extent than the pentalene subunit. Theoretical calculations supported the thermodynamically favored electronic rearrangement of the BCB unit and the strong antiaromaticity of the pentalene structure. The combined experimental (¹H NMR, UV–vis, CV, and crystallography) and computational study of the synthesized molecules proved the reversal of the structure–antiaromaticity correlation compared to aceno-pentalenes. It has to be noted that our results do not violate the proposed correlation between bond order and antiaromaticity⁷⁴ but expand the design to unsymmetric bis(antiaromatic) systems, where the bond orders favor the construction of linear and highly antiaromatic compounds.

The work presented here provides the basis of the syntheses of linear π -systems having strongly antiaromatic subunits and could aid in the design of solid-state structures beneficial for organic electronic device applications. Research on the extension of this design to different antiaromatic subunits and the construction of more extended π -systems is in progress in our laboratories.

EXPERIMENTAL SECTION

General Information. Commercial reagents, solvents, and catalysts (Aldrich, Fluorochem, VWR) were purchased as reagent-grade and used without further purification. Solvents for extraction or column chromatography were of technical quality. Organic solutions were concentrated by rotary evaporation at 25–40 °C. Thin-layer chromatography was carried out on SiO₂-layered aluminum plates (60778-2SEA, Fluka). Column chromatography was performed using SiO₂-60 (230–400-mesh ASTM, 0.040–0.063 mm from Merck) at 25 °C or using a Teledyne Isco CombiFlash Rf+ automated flash chromatographer with silica gel (25–40 μ m, Zeochem). Room temperature refers to 25(\pm 1) °C. NMR spectra were acquired on a Varian 500 NMR spectrometer, running at 500 and 126 MHz for ¹H and ¹³C, respectively. The residual solvent peaks were used as the internal reference. Chemical shifts (δ) are reported in ppm. The following abbreviations are used to indicate the multiplicity in ¹H NMR spectra: s, singlet; d, doublet; t, triplet; q, quartet; h, heptet; and m, multiplet. ¹³C NMR spectra were acquired on a broad-band decoupled mode. Gas chromatography–mass spectrometry (GC–MS) analysis was performed on a Shimadzu GCMS-QP2010 Ultra System operated in the electron impact ionization (EI) mode. Mass spectrometric measurements were performed using a Q-TOF Premier mass spectrometer (Waters Corporation, Milford, MA) in the positive electrospray ionization mode.

General Procedures. *General Procedure for the Sonogashira Reactions (GP1).* Orthohalo-formylbiphenylene (1 equiv) and 4-ethynylanisole (1.05 equiv) were dissolved in triethylamine (0.1 M). The solution was added to a vial that contained Pd(PPh₃)₂Cl₂ (0.05 equiv) and CuI (0.05 equiv) under an inert atmosphere (N₂). The mixture was heated to 50 °C in an aluminum heating block and stirred for 3–12 h. After the reaction was completed, the mixture was diluted with EtOAc and washed once with 10% HCl and twice with brine. The organic phase was dried over MgSO₄. The solvent was evaporated in vacuo, and the crude product was further purified with column chromatography (SiO₂, *n*-hexane/EtOAc).

General Procedure for Gem-dibromoolefination (GP2). The product of GP1 (1 equiv) and CBr₄ (1.5 equiv) was dissolved in 1,2-dichloroethane (DCE, 0.1 M). After the solution was purged with N₂ for 10 min, P(OiPr)₃ (3 equiv) was added. Following 3 h of stirring, the solution was diluted with DCE and was washed with water. The organic phase was separated and then dried over MgSO₄. The solvent was evaporated in vacuo, and the crude product was further purified with column chromatography (SiO₂, *n*-hexane/EtOAc).

General Procedure for the Carbopalladation Cascade Reaction (GP3). The product of GP2 (1 equiv) and diphenylacetilene (5 equiv) was dissolved in toluene (0.1 M). The solution was added to a vial, which contained Pd(PPh₃)₂Cl₂ (0.1 equiv), Zn (0.1 equiv), and K₂CO₃ (2 equiv) under an inert atmosphere (N₂). The mixture was heated up to 110 °C in an aluminum heating block and stirred for 2 h. Subsequently, the reaction was cooled to rt, hydroquinone (1 equiv) was added to the mixture, and it was purged again with N₂. The reaction was heated up to 110 °C and stirred for 16 h at this temperature. This was followed by cooling to rt, and the mixture was diluted with EtOAc and washed twice with water and once with brine. The organic phase was separated and then dried over MgSO₄. The solvent was evaporated under reduced pressure, and the crude product was further purified with column chromatography (SiO₂, *n*-hexane/EtOAc).

Synthetic Procedures. *Synthesis of Biphenylene (2).*¹¹⁶ Isopentyl nitrite (2.4 mL, 18.0 mmol) was added to a solution of anthranilic acid (2 g, 14.6 mmol) in THF (30 mL). This mixture was

stirred for 1 h upon which the formation of a red precipitate was observed. Subsequently, a catalytic amount of trichloroacetic acid (~20 mg) was added to the mixture and stirred until a brown precipitate was observed (approximately 1 h). The brown precipitate was filtered (Caution! Always keep the precipitate wet by solvent! The dried precipitate is highly explosive!) and washed with THF until the solvent was colorless. Subsequently, the residue was washed with 1,2-dichloroethane (3 \times 10 mL) and then suspended in 1,2-dichloroethane. This suspension was carefully added to gently boiling 1,2-dichloroethane (60 mL, heated in an oil bath), yielding a dark-brown solution, which was stirred and boiled for another 15 min. The resulting solution was allowed to cool to room temperature and was washed with brine. The organic phase was separated and dried over MgSO₄. The solvent was evaporated in vacuo, giving a brown solid as a crude product, which was purified further with column chromatography (SiO₂, *n*-hexane) to give the product as yellowish-white crystals. Yield: 303 mg, 27%. ¹H NMR (500 MHz, CDCl₃) δ = 6.74 (dd, *J* = 4.8, 2.9 Hz, 1H), 6.63 (dd, *J* = 4.8, 2.9 Hz, 1H) ppm; ¹³C{¹H} NMR (126 MHz, CDCl₃) δ = 151.5 (4), 128.4 (4), 117.5 (4) ppm.

Synthesis of 2-Formylbiphenylene (3). SnCl₄ (4 mL, 34.2 mmol) was added to a stirred solution of 2 (300 mg, 1.97 mmol) and dichloromethyl methyl ether (1 mL, 11.1 mmol) in 1,2-dichloroethane (30 mL) under a N₂ atmosphere. The solution was stirred for 16 h at rt, and then, ice-cold HCl solution (3 M, 50 mL) was added. Following 1 h of vigorous stirring at rt, the mixture was extracted with CH₂Cl₂. The organic phase was washed with water and brine and then dried over MgSO₄. The solvent was removed under reduced pressure, and the crude mixture was purified by flash column chromatography (SiO₂, *n*-hexane \rightarrow *n*-hexane/EtOAc (15%)). Evaporation of the solvent gave the product as yellow crystals. Yield: 225 mg, 64%. ¹H NMR (500 MHz, CDCl₃) δ = 9.68 (s, 1H), 7.27 (d, *J* = 7.1 Hz, 1H), 7.07 (s, 1H), 6.86 (dt, *J* = 17.1, 7.4 Hz, 2H), 6.76 (dd, *J* = 12.3, 6.4 Hz, 3H) ppm; ¹³C{¹H} NMR (126 MHz, CDCl₃) δ = 191.0, 158.3, 152.1, 149.9, 149.5, 137.2, 136.7, 130.2, 129.2, 119.2, 118.7, 116.7, 114.1 ppm; HRMS (ESI): *m/z*: [M + H]⁺, calcd for C₁₂H₁₀O⁺: 181.0653; found 181.0651.

Synthesis of 2-Hydroxymethylbiphenylene (4). A suspension of NaBH₄ (66.1 mg, 1.75 mmol) in ethanol (5.25 mL) was added in one portion to a stirred solution of 3 (210 mg, 1.17 mmol) in THF (585 μ L). The mixture was stirred for 1 h at rt, and then water (~50 mL) was added. The solution was extracted with EtOAc (2 \times 20 mL). The organic phase was washed with water and brine and dried over MgSO₄. Evaporation of the solvent under reduced pressure gave the product as bright-yellow crystals. Yield: 196 mg, 92%. ¹H NMR (500 MHz, CDCl₃) δ = 6.78–6.72 (m, 2H), 6.71 (d, *J* = 7.0 Hz, 1H), 6.67 (s, 1H), 6.64 (dd, *J* = 5.8, 3.7 Hz, 2H), 6.59 (d, *J* = 7.0 Hz, 1H), 4.44 (s, 2H), 1.89 (s, 1H) ppm; ¹³C{¹H} NMR (126 MHz, CDCl₃) δ = 151.9, 151.0, 150.9, 150.8, 141.2, 128.5, 128.4, 126.6, 117.6, 117.5, 117.2, 116.8, 65.8 ppm; HRMS (ESI): *m/z*: [M]⁺, calcd for C₁₂H₁₁O⁺: 182.0732; found 182.0726.

Synthesis of 2-Hydroxymethylbiphenylene 2-Hydroxymethyl-1-iodobiphenylene (5). *n*-BuLi (2.07 mL, 5.2 mmol, 2.5 M solution in hexane) was added slowly to a stirred solution of 4 (445 mg, 2.5 mmol) in diethyl ether (40 mL) at rt. The mixture was heated to reflux in an oil bath and stirred for 1 h. Subsequently, the solution was cooled to room temperature, and 1-chloro-2-iodoethane (340.6 mg, 3.1 mmol) in diethyl ether (10 mL) was added. The resulting solution was refluxed for another 30 min, and then it was cooled to rt and washed with water and brine. The ether solution was dried over MgSO₄. The solvent was evaporated under reduced pressure, giving a dark-brown oil. The crude oily product was purified by column chromatography (SiO₂, CH₂Cl₂/MeOH (2%)). The solvent was removed under reduced pressure, yielding the product as bright-yellow crystals. Yield: 335 mg, 44%. ¹H NMR (500 MHz, CDCl₃) δ = 6.89–6.84 (m, 1H), 6.84–6.78 (m, 2H), 6.75 (d, *J* = 6.8 Hz, 1H), 6.67–6.61 (m, 1H), 6.54 (d, *J* = 6.8 Hz, 1H), 4.45 (s, 2H), 1.95 (s, 1H) ppm; ¹³C{¹H} NMR (126 MHz, CDCl₃) δ = 157.8, 151.7, 151.0, 149.0, 141.1, 129.4, 128.5, 128.0, 117.8, 116.6, 116.5, 83.8, 68.2 ppm;

HRMS (ESI): m/z : $[M]^+$, calcd for $C_{12}H_9OI^+$: 307.9701; found 307.9701.

Synthesis of 1-Iodo-2-formylbiphenylene (6). Jones reagent (80 μ L) was added to the solution of **5** (80 mg, 0.26 mmol) in acetone (1 mL). The mixture was stirred for 10 min, and a green solid precipitate was formed. The precipitate was filtered, and acetone was removed in vacuo. The remaining residue was dissolved in EtOAc (10 mL) and washed with water. The organic phase was separated and dried over $MgSO_4$. The solvent was evaporated under reduced pressure, giving an orange oil. The oil was purified by column chromatography (SiO_2 , n -hexane/EtOAc (10%)). Evaporation of the solvent under reduced pressure gave the product as yellow crystals. Yield: 75 mg, 94%. 1H NMR (500 MHz, $CDCl_3$) δ = 9.80 (s, 1H), 7.31 (d, J = 7.0 Hz, 1H), 6.96 (d, J = 6.4 Hz, 1H), 6.95–6.86 (m, 2H), 6.76 (d, J = 6.4 Hz, 1H), 6.66 (d, J = 7.0 Hz, 1H) ppm; $^{13}C\{^1H\}$ NMR (126 MHz, $CDCl_3$) δ = 192.7, 158.7, 158.6, 150.6, 148.0, 135.0, 133.6, 130.5, 130.1, 119.4, 117.5, 116.6, 82.2 ppm; HRMS (ESI): m/z : $[M + H]^+$, calcd for $C_{12}H_8OI^+$: 306.9620; found 306.9618.

Synthesis of 1-((4-Ethoxyphenyl)ethynyl)biphenylene-2-carbaldehyde (7). This compound was prepared from compound **6** according to GP1. The crude product was purified by flash column chromatography (SiO_2 , n -hexane \rightarrow n -hexane/EtOAc (10%)), giving the product as yellow crystals. Yield: 34 mg, 57%. 1H NMR (500 MHz, $CDCl_3$) δ = 10.25 (s, 1H), 7.47 (dd, J = 13.1, 7.9 Hz, 3H), 6.96–6.81 (m, 5H), 6.75 (d, J = 6.7 Hz, 1H), 6.67 (d, J = 7.1 Hz, 1H), 3.84 (s, 3H) ppm; $^{13}C\{^1H\}$ NMR (126 MHz, $CDCl_3$) δ = 190.2, 160.4, 157.4, 154.3, 149.6, 149.1, 135.1, 133.4 (2), 132.1, 130.4, 129.6, 119.2, 118.8, 116.4, 114.7, 114.3 (2), 114.1, 98.1, 81.2, 55.5 ppm; HRMS (ESI): m/z : $[M + H]^+$, calcd for $C_{22}H_{15}O_2^+$: 311.1072; found 311.1072.

Synthesis of 2-(2,2-Dibromovinyl)-1-((4-methoxyphenyl)ethynyl)biphenylene (8). This compound was prepared according to GP2. The crude product was purified by flash column chromatography (SiO_2 , n -hexane \rightarrow n -hexane/EtOAc (10%)), giving the product as orange-yellow crystals. Yield: 28 mg, 54%. 1H NMR (500 MHz, $CDCl_3$) δ = 7.63 (s, 1H), 7.47 (d, J = 8.9 Hz, 2H), 7.20 (dd, J = 7.3, 0.8 Hz, 1H), 6.90 (d, J = 8.9 Hz, 2H), 6.84–6.78 (m, 3H), 6.70–6.66 (m, 1H), 6.59 (d, J = 7.2 Hz, 1H), 3.84 (s, 3H) ppm; $^{13}C\{^1H\}$ NMR (126 MHz, $CDCl_3$) δ = 160.2, 153.3, 151.2, 150.1, 150.1, 136.0, 135.7, 133.3 (2), 129.3, 129.1, 128.6, 118.3, 118.1, 116.2, 115.1, 114.3 (2), 112.8, 97.4, 90.7, 82.7, 55.5 ppm; HRMS (ESI): m/z : $[M]^+$, calcd for $C_{23}H_{14}OBr_2^+$: 463.9411; found 463.9416.

Synthesis of 1-(4-Methoxyphenyl)-2,3-diphenylpentaleno[1,2-*a*]-biphenylene (9). This compound was prepared according to GP3. The crude product was purified by flash column chromatography (SiO_2 , n -hexane \rightarrow n -hexane/EtOAc (5%)), giving the product as purple crystals. Yield: 12 mg, 42%. 1H NMR (500 MHz, $CDCl_3$) δ = 7.22–7.11 (m, 8H), 7.01 (d, J = 8.4 Hz, 2H), 6.90 (d, J = 7.2 Hz, 2H), 6.88–6.84 (m, 1H), 6.84–6.78 (m, 3H), 6.73 (t, J = 7.5 Hz, 1H), 6.60 (s, 1H), 6.43 (d, J = 6.7 Hz, 1H), 6.18 (d, J = 6.7 Hz, 1H), 5.76 (d, J = 7.0 Hz, 1H), 3.82 (s, 3H) ppm; 1H NMR (500 MHz, CD_2Cl_2) δ = 7.25–7.10 (m, 8H), 7.01 (d, J = 8.5 Hz, 2H), 6.92 (d, J = 7.3 Hz, 2H), 6.88 (t, J = 7.5 Hz, 1H), 6.82 (d, J = 8.6 Hz, 3H), 6.75 (t, J = 7.5 Hz, 1H), 6.62 (s, 1H), 6.47 (d, J = 6.7 Hz, 1H), 6.21 (d, J = 6.7 Hz, 1H), 5.76 (d, J = 7.1 Hz, 1H), 3.82 (s, 3H) ppm; $^{13}C\{^1H\}$ NMR (126 MHz, CD_2Cl_2) δ = 159.9, 154.0, 151.1, 150.8, 150.3, 149.4, 141.1, 136.2, 135.3, 134.0, 132.3 (2), 131.9, 130.4 (2), 129.2 (2), 129.1, 129.1, 128.7 (2), 128.4 (2), 128.2, 127.7, 127.3, 125.3, 120.9, 118.2, 113.9, 113.9 (2), 55.9 ppm; HRMS (ESI): m/z : $[M + H]^+$, calcd for $C_{37}H_{25}O^+$: 485.1905; found 485.1902.

Synthesis of 2-Bromobiphenylene (10). Compound **2** (700 mg, 4.6 mmol) and NBS (930 mg, 5.2 mmol) were dissolved in dried DMF (12 mL). The mixture was stirred for 2–3 h while the reaction was followed with GC–MS and TLC (SiO_2 , n -hexane). After the reaction was completed, the solution was diluted with DCM (40 mL) and washed with water (3 \times 20 mL). The organic phase was separated and dried over $MgSO_4$, and the solvents were evaporated in vacuo, giving an orange residue. The crude product was further purified with column chromatography (SiO_2 , n -hexane). Evaporation of the eluent under reduced pressure gave the product as bright-yellow crystals.

Yield: 828 mg, 78%. 1H NMR (500 MHz, $CDCl_3$) δ 6.90 (d, J = 7.3 Hz, 1H), 6.79–6.76 (m, 2H), 6.75 (s, 1H), 6.68–6.63 (m, 2H), 6.48 (d, J = 7.3 Hz, 1H) ppm; $^{13}C\{^1H\}$ NMR (126 MHz, $CDCl_3$) δ = 152.8, 150.4, 150.0, 149.9, 130.6, 129.1, 128.7, 121.6, 121.3, 118.7, 118.2, 118.0; HRMS (ESI): m/z : $[M + H]^+$, calcd for $C_{12}H_7Br^+$: 229.9731; found 229.9731.

Synthesis of 3-Bromo-2-formylbiphenylene (11). $SnCl_4$ (6 mL, 51.3 mmol) was added to a stirred solution of **10** (708 mg, 3.06 mmol) and dichloromethyl methyl ether (1.5 mL, 16.6 mmol) in 1,2-dichloroethane (45 mL) under a N_2 atmosphere. The solution was stirred for 16 h at rt, and then, ice-cold HCl solution (3 M, 75 mL) was added. Following 1 h of vigorous stirring at room temperature, the mixture was extracted with CH_2Cl_2 . The organic phase was washed with water and brine and dried over $MgSO_4$. The solvent was removed under reduced pressure, and the crude mixture was purified by flash column chromatography (SiO_2 , n -hexane \rightarrow n -hexane/EtOAc (10%)). Evaporation of the solvent gave the product as yellow crystals. Yield: 149 mg, 19%. 1H NMR (500 MHz, $CDCl_3$) δ = 10.16 (s, 1H), 7.09 (s, 1H), 6.96–6.84 (m, 3H), 6.81 (d, J = 7.3 Hz, 2H) ppm; $^{13}C\{^1H\}$ NMR (126 MHz, $CDCl_3$) δ = 191.2, 158.6, 150.7, 149.7, 148.6, 133.5, 131.3, 131.0, 129.6, 122.5, 119.9, 119.3, 116.4 ppm; HRMS (ESI): m/z : $[M + H]^+$, calcd for $C_{13}H_8OBr^+$: 258.9759; found 258.9759.

Synthesis of 3-((4-Methoxyphenyl)ethynyl)biphenylene-2-carbaldehyde (12). This compound was prepared according to GP1. The crude product was purified by flash column chromatography (SiO_2 , n -hexane \rightarrow n -hexane/EtOAc (10%)), giving the product as yellow crystals. Yield: 191.5 mg, 77%. 1H NMR (500 MHz, $CDCl_3$) δ = 10.44 (s, 3H), 7.46 (d, J = 8.7 Hz, 7H), 7.13 (s, 3H), 6.94–6.87 (m, 13H), 6.81 (dd, J = 8.3, 4.7 Hz, 10H), 3.84 (s, 11H) ppm; $^{13}C\{^1H\}$ NMR (126 MHz, $CDCl_3$) δ = 190.9, 160.5, 156.9, 150.8, 150.0, 149.2, 137.2, 133.4 (2), 131.3, 130.6, 129.6, 120.3, 119.6, 119.2, 114.5, 114.4 (2), 113.8, 98.0, 85.1, 55.5 ppm; HRMS (ESI): m/z : $[M + H]^+$, calcd for $C_{22}H_{15}O_2^+$: 311.1072; found 311.1076.

Synthesis of 2-(2,2-Dibromovinyl)-3-((4-methoxyphenyl)ethynyl)biphenylene (13). This compound was prepared according to GP2. The crude product was purified by flash column chromatography (SiO_2 , n -hexane \rightarrow n -hexane/EtOAc (10%)), giving the product as orange-yellow crystals. Yield: 112 mg, 69%. 1H NMR (500 MHz, $CDCl_3$) δ = 7.77 (s, 1H), 7.45 (d, J = 8.8 Hz, 2H), 7.11 (s, 1H), 6.89 (d, J = 8.8 Hz, 2H), 6.86–6.79 (m, 2H), 6.75 (s, 1H), 6.72 (ddd, J = 6.7, 4.4, 1.5 Hz, 2H), 3.84 (s, 3H) ppm; $^{13}C\{^1H\}$ NMR (126 MHz, $CDCl_3$) δ = 160.1, 150.9, 150.4, 150.2, 137.8, 136.5, 133.2 (2), 129.3, 129.2, 123.3, 119.9, 118.4, 118.4, 116.6, 115.2, 114.3 (2), 96.8, 90.6, 87.2, 55.5 ppm; HRMS (ESI): m/z : $[M]^+$, calcd for $C_{23}H_{14}OBr_2^+$: 463.9411; found 463.9409.

Synthesis of 1-(4-Methoxyphenyl)-2,3-diphenylpentaleno[1,2-*b*]-biphenylene (14). This compound was prepared according to GP3. The crude product was purified by flash column chromatography (SiO_2 , n -hexane \rightarrow n -hexane/EtOAc (5%)), giving the product as a brown solid. Yield: 15 mg, 27%. 1H NMR (500 MHz, $CDCl_3$) δ = 7.24–7.09 (m, 6H), 7.05 (dd, J = 6.6, 3.1 Hz, 2H), 6.96 (d, J = 8.8 Hz, 2H), 6.86 (d, J = 6.9 Hz, 2H), 6.77 (d, J = 8.8 Hz, 2H), 6.69–6.65 (m, 2H), 6.49–6.44 (m, 2H), 6.32 (s, 1H), 6.19 (s, 1H), 6.02 (s, 1H), 3.80 (s, 3H) ppm; 1H NMR (500 MHz, CD_2Cl_2) δ = 7.22–7.12 (m, 6H), 7.05 (dd, J = 6.6, 3.0 Hz, 2H), 6.97 (d, J = 8.7 Hz, 2H), 6.88 (d, J = 7.0 Hz, 2H), 6.77 (d, J = 8.7 Hz, 2H), 6.70–6.66 (m, 2H), 6.49 (dd, J = 12.9, 5.2 Hz, 2H), 6.32 (s, 1H), 6.23 (s, 1H), 6.04 (s, 1H), 3.79 (s, 3H) ppm; $^{13}C\{^1H\}$ NMR (126 MHz, CD_2Cl_2) δ = 160.8, 153.5, 152.2, 150.4, 150.2, 150.1, 149.3, 147.9, 147.6, 140.3, 137.8, 135.5, 134.9, 134.8, 133.6, 131.1 (2), 130.1 (2), 128.8 (2), 128.6 (2), 128.4 (2), 128.3, 128.1, 127.8, 127.7, 126.7, 116.5, 116.3, 115.5, 113.8 (2), 112.6, 55.8 ppm; HRMS (ESI): m/z : $[M + H]^+$, calcd for $C_{37}H_{25}O^+$: 485.1905; found 485.1902.

■ ASSOCIATED CONTENT

Supporting Information


The Supporting Information is available free of charge at <https://pubs.acs.org/doi/10.1021/acs.joc.9b03119>.

X-ray crystallography details for compound **9**; electrochemical measurement details and cyclic voltammograms for compounds **9** and **14**; details of computational studies of molecules **9'**, **14'**, **15'**, and **16'**; ^1H and ^{13}C NMR and HRMS spectra for all new compounds (PDF)
Crystal structure (CIF)

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
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Notes

The authors declare no competing financial interest.

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DEDICATION

[†]This article is dedicated to Professor François Diederich.

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