9,10-Dihydrophenanthrene with Two Spiro(dibenzocycloheptatriene) Units: A Highly Strained Caged Hydrocarbon Exhibiting Reversible Electrochromic Behavior

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Abstract: The title dispiro hydrocarbon 1 was designed as a new electrochromic material. This multiply clamped hexaphenylethane-type electron donor was prepared from 2,2′-diiodobiphenyl via biphenyl-2,2′-diylbis(dibenzotropylium) 2⁺ salt. X-ray analysis of 1 revealed a highly strained structure as reflected by an elongated “ethane” bond [bond length: 1.6665(17) Å] and nearly eclipsed conformation. The weakened bond was cleaved upon two-electron oxidation to regenerate the deeply colored dication 2⁺. The reversible interconversion between 1 and 2⁺ is accompanied not only by a drastic color change but also by C–C bond formation/cleavage. Thus, the voltammogram showed a pair of well-separated redox waves, which is characteristic of “dynamic redox (dyrex)” behavior. The tetrahydro derivative of 1 with two units of spiro(dibenzocycloheptadiene), which suffers from more severe steric congestion, was also prepared. The crystallographically determined bond length for the central C–C bond [1.705(4) Å] is greatest among the values reported for 9,9,10,10-tetraaryl-9,10-dihydrophenanthrene derivatives.

Keywords: polycyclic aromatic hydrocarbons; long bond; dynamic redox system; dication; dibenzotropylium; electrochromism; through-bond interaction

1. Introduction

π-Conjugated systems have attracted much attention due to the facile modification of their properties by unique molecular design, as shown in several reports on sterically-congested and/or curved polycyclic aromatic hydrocarbons (PAHs) [1–4]. Some of them have been shown to have unusual geometric features, such as extremely twisted aromatic rings [5–12] or an extraordinarily long C–C bond [13–20], which result in intriguing physical properties.

9,10-Dihydrophenanthrene (DHP) with four aryl groups at the 9,9,10,10-positions is the clamped derivative of hexaphenylethane (HPE). In contrast to the easy dissociation of unclamped HPEs into two trityl radicals upon cleavage of the “ethane” bond [21], a series of tetraaryl-DHPs (Ar₄DHPs) are considered to be thermally stable [22]. The C₉–C₁₀ bond fission in Ar₄DHP is no longer entropically favored, so that even an elongated bond with a bond length far greater than the standard (1.54 Å) is intact when incorporated in the Ar₄DHP skeleton [23]. Thus, the DHP skeleton is one of the most useful scaffolds for examining highly strained structures and their special properties.

Hexabenzo[4.4.4]propellane A [24], which was first reported in 1971, is a multiply clamped HPE in which the central Csp³–Csp³ bond [bond length: 1.563(2) Å] is not greatly expanded [25]. In this
nonspiro-type clamped HPE, the steric repulsion among the six benzene rings is effectively reduced by adopting a twisted conformation with a torsion angle (α) of about 60° around the central C–C bond (Figure 1). On the other hand, dibenzodinaphtho[4.3.3]propellane B [26] has a longer bond [1.612(4) Å] because it has a less-skewed geometry due to the rigid naphthalene planes. This difference in bond length as well as its correlation with torsion/twisting angles (α, θ) could be supported by DFT calculations. On the other hand, Ar₄DHPs were predicted to have much longer C–C bonds than those in B since the steric hindrance regarding the C₉–C₁₀ bond is greater for two unfused benzene rings than for a naphthalene nucleus.

We previously studied the structures and properties of a series of Ar₄DHPs [27–32]. When we conducted an X-ray analysis of (4-CH₃OC₆H₄)₄DHP (C), we could not obtain accurate values for the bond length or torsion/twisting angles due to the positional disorder of the ethane unit [27]. Such disorder has often been observed in globular ethanes (e.g., unclamped HPEs [33] or hexachloroethane [34]), but was not present in the spiro-type Ar₄DHPs (D and E). Their C₉–C₁₀ bond lengths could be accurately determined to be 1.646(4) and 1.635(2) Å by X-ray analyses [29,30]. These values are greater than those for other DHPs, and, thus, the spiro ring can be considered to be the key structure for observing a very long bond in Ar₄DHPs.

Another interesting point regarding Ar₄DHP is the electrochromic response with “dynamic redox (dyrex)” behavior. Two-electron oxidation induces the formation of cationic chromophores accompanied by fission of the elongated C₉–C₁₀ bond. Heterocyclic units (acridan/xanthene) or alkoxy/amino groups have often been incorporated into Ar₄DHP to raise the HOMO level and to stabilize the corresponding dicationic species. However, under an appropriate molecular design, we envisaged that reversible electrochromic systems could be constructed without the aid of heteratoms. Herein we report the details of the title spiro-type Ar₄DHP 1, which is a pure hydrocarbon that can still exhibit reversible electrochromic behavior thanks to the raised HOMO level of 1 as well as the stability of the 14π–dibenzotropylium unit in the bond-dissociated dication 2²⁺.

![Figure 1](image_url)

**Figure 1.** Reported structural parameters of Ar₄DHPs determined by X-ray analyses. The values obtained by DFT calculations at the B3LYP/6-31G* level in this study are shown in brackets.

## 2. Results and Discussion

### 2.1. Design and Theoretical Study

The through-bond interaction (TBI) [35] in PAHs is an important factor which modifies the MO levels that determine their electron-donating properties. As postulated based on a simplified s-cis diphenylethane model (Figure 2), the σ–orbital of the ethane bond would be lowered whereas the π⁺–orbital should be raised through TBI. Greater perturbation is expected when the two energy levels are closer. The parallel arrangement of these orbitals is also important for realizing effective TBI.
For Ar₄DHP, TBI would be maximized by elongation of the C₉–C₁₀ bond, since a longer bond has a higher σ-orbital level to narrow the energy gap toward the π⁺-orbital. At the same time, an eclipsed conformation is desirable to ensure that the orbitals are parallel. The spiro(dibenzocycloheptatriene) units are the ideal skeleton to be incorporated into Ar₄DHP. In general, Ar₄DHPs tend to adopt a skewed geometry to reduce the “front strain” [21] among the four aryl groups over the C₉–C₁₀ bond. The spiro(xanthene or 10-methylacridane) with the central six-membered ring can force the DHP skeleton to adopt a less-skewed geometry (Figure 1). Here, we propose that spiro(dibenzocycloheptatriene) with a central seven-membered ring is more favorable since it can make the DHP skeleton become nearly eclipsed by the greater steric hindrance between the inner protons on the benzo groups and the DHP plane (Figure 3). The nearly eclipsed conformation would also cause elongation of the C₉–C₁₀ bond due to the greater steric repulsion than in the skewed conformation, and, thus, the HOMO level of Ar₄DHP with two spiro(dibenzocycloheptatriene) 1 would be effectively raised through TBI.

![Figure 2. Schematic view of through-bond interaction in the diphenylethane model.](image)

![Figure 3. Molecular design of Ar₄DHP 1 with two spiro(dibenzocycloheptatriene) units.](image)

The tetrahydro derivative 3 with two units of spiro(dibenzocycloheptadiene) was also included in this study as a reference compound; it cannot adopt a similar eclipsed conformation due to severe steric repulsion between the CH₂CH₂ units in the cycloheptadiene rings.

Density functional theory (DFT) calculations [36] at the B3LYP/6-31G* level predicted that the optimized structure of 1 adopts a nearly eclipsed conformation, as designed. The C₉–C₁₀ bond length was estimated to be greater than 1.7 Å, which is much greater than the values previously reported for other Ar₄DHPs (Figure 4). The calculated HOMO level of 1 was raised to −5.25 eV, which is much higher than that of (4-CH₃OC₆H₄)₄DHP (−5.44 eV). Thus, we designed a stronger electron-donating hydrocarbon than Ar₄DHP with four CH₃O groups.
When dication 2 with the results of DFT calculations, 7 was isolated. The eight-membered-ring endoperoxide with BuLi and dibenzotropone [37]. Upon treatment of 2,2′-diiodobiphenyl as a starting material, diol 4 was prepared by successive reactions with HBF4 in trifluoroacetic anhydride (TFAA), the dication salt 2\(^{2+}\)(BF\(_4\)^-)\(_2\) was isolated as a deep purple powder with 85% yield. When dication 2\(^{2+}\) was reduced with Zn powder, the newly designed dispiro Ar\(_4\)DHP 1 with two spiro(dibenzocycloheptatriene) units was obtained quantitatively as a white solid (Scheme 1).

In the case of the spiro(dibenzocycloheptadiene) derivative, diol 5 was obtained in a similar manner and treated with TMSCIO\(_4\) [38] in 1,1,1,3,3,3-hexafluoro-2-propanol (HFIP). The resulting precursor dication was directly reduced with Zn powder to give the desired compound 3 in 33% yield as a white solid. The low yield of 3 can be partly explained by the formation of by-products/decomposition products. From the mixture, isomer 6 with a spiro(flourene) framework was isolated. The eight-membered-ring endoperoxide 7 was also generated as a decomposition pathway of 3.

X-ray analyses were performed at 150 K by using single crystals of 1 and 3 (Figure 6). Consistent with the results of DFT calculations, 1 adopts a nearly eclipsed geometry. For the DHP skeleton, the torsion angle \(\alpha\) over the C\(_9\)–C\(_{10}\) bond is 23.78(13)° and the dihedral angle \(\theta\) for the biphenyl unit is only 3.7°. Due to the lack of skewing deformation to reduce the “front” strain, the C\(_9\)–C\(_{10}\) bond is expanded. The length of 1.6665(17) Å is greater than any value ever reported for Ar\(_4\)DHPs [39].

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Figure 4. Optimized structures (left: front view, right: side view) of (a) 1 and (b) 3.

As shown in Figure 5, the HOMO of 1 has the character of \((\pi^+ - \sigma)\), as evidenced by the large orbital coefficients on the ethane bond. Thus, the effective TBI between \(\pi^+\) and \(\sigma\) causes the perturbation of both orbital levels, resulting in an increase in the energy of \((\pi^+ - \sigma)\) to become higher than that of \(\pi^-\) [next-highest OMO (NHOMO), \(-5.39\) eV].

Figure 5. (a) HOMO and (b) next-highest OMO (NHOMO) of 1 based on DFT calculations (B3LYP/6-31G*).

2.2. Preparation and X-ray Analysis

With 2,2’-diiodobiphenyl as a starting material, diol 4 was prepared by successive reactions with BuLi and dibenzotropone [37]. Upon treatment of 4 with HBF\(_4\) in trifluoroacetic anhydride (TFAA), the dication salt 2\(^{2+}\)(BF\(_4\)^-)\(_2\) was isolated as a deep purple powder with 85% yield. When dication 2\(^{2+}\) was reduced with Zn powder, the newly designed dispiro Ar\(_4\)DHP 1 with two spiro(dibenzocycloheptatriene) units was obtained quantitatively as a white solid (Scheme 1).

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As indicated by the DFT calculation shown in Figure 4, both of the crystallographically independent molecules of 3 adopt a twisted conformation. In this way, close contact between the two CH₂CH₂ units over the C₉–C₁₀ bond is avoided. However, the molecules still suffer from large steric hindrance between the two spiro(dibenzocycloheptadiene) units. Thus, the bond lengths for the C₉–C₁₀ bond [1.705(4) and 1.697(4) Å] of 3 are the highest values reported for the Ar₄DHP, including 1.

Scheme 1. Preparation of newly designed Ar₄HPEs 1 and 3.

![Scheme 1. Preparation of newly designed Ar₄HPEs 1 and 3.](image)

Figure 6. ORTEP drawings (left: front view, right: side view) of (a) 1; (b) 3 (mol-1) and (c) 3 (mol-2) at 150 K. Thermal ellipsoids are shown at the 50% probability level.

2.3. Redox Behavior

To investigate the electron-donating properties and reversibility of the redox behavior of 1, redox potentials were measured by cyclic voltammetry in CH₂Cl₂ (Figure 7). The one-wave two-electron oxidation peak was observed, which corresponds to the formation of the dication 2⁺ accompanied by C₉–C₁₀ bond cleavage. The corresponding reduction peak of 2⁺ appeared at +0.42 V. Such a separation...
of redox peaks is characteristic of 

of redox peaks is characteristic of dyrex systems. When scanning was repeated twice, no change was observed in the voltammogram, which indicates highly reversible interconversion between 1 and $2^{2+}$.

![Figure 7](image_url)

**Figure 7.** Cyclic voltammogram of 1 in CH$_2$Cl$_2$ containing 0.1 M Bu$_4$NBF$_4$ as a supporting electrolyte (scan rate 100 mV s$^{-1}$, Pt electrode).

The electron-donating properties of hydrocarbon 1 are striking. The oxidation potential (+0.95 V) is far less positive than that of (4-CH$_3$OCC$_6$H$_4$)$_4$DHP (+1.44 V) measured under similar conditions [28]. Such a change can be qualitatively accounted for by the different HOMO levels calculated by the DFT method, which demonstrates the validity of our concept for raising the HOMO level of Ar$_4$DHP through TBI without the aid of heteroatoms.

2.4. Electrochromic Behavior

Upon the electrochemical oxidation of 1 in CH$_2$Cl$_2$, from colorless to red, an electrochromic response was observed with an isosbestic point at 252 nm. The final UV/Vis spectrum is identical to that of the isolated dication $2^{2+}$ (BF$_4^{-}$)$_2$ (Figure 8). Regeneration of 1 with a drastic color change from red to colorless was attained by reverse electrolysis of the as-prepared dication solution (see Supplementary Materials Figure S1). This is a rare successful demonstration of electrochromic behavior based on pure hydrocarbon redox species [40].

![Figure 8](image_url)

**Figure 8.** Change in the UV/Vis spectrum upon electrochemical oxidation (20 µA) of 1 in CH$_2$Cl$_2$ containing 0.05 M Bu$_4$NBF$_4$ as a supporting electrolyte (every 4 min).
Quantitative interconversion between donor 1 and dication 2\(^{2+}\) can be conducted in a preparative manner by chemical oxidation and reduction. Thus, upon treatment of 1 with two equivalents of (4-BrC\(_6\)H\(_4\))\(_3\)N\(^{+}\)SbCl\(_6\)\(^{-}\) in CH\(_2\)Cl\(_2\), dication 2\(^{2+}\) was isolated quantitatively as a stable salt. The X-ray structural analysis of 2\(^{2+}\)(SbCl\(_6\)\(^{-}\))\(_2\)·2(CH\(_2\)Cl\(_2\)) crystal shows that there is a \(\pi-\pi\) stacking interaction between two dibenzotropylium units [dihedral angle: 12.1\(^{\circ}\); closest C–C contact: 3.31(2) Å] (Figure 9a).

While 2\(^{2+}\) was persistent in CH\(_2\)Cl\(_2\), it underwent faci transformation into spiro(fluorene)-type monocation 8\(^{+}\) in CH\(_3\)CN. When this conversion was followed by UV/Vis spectroscopy at 23 \(^{\circ}\)C, several isosbestic points were observed (Figure 10), from which a reaction rate of \(5.0 \times 10^{-4}\) s\(^{-1}\) was deduced. Friedel–Crafts-type cyclization also proceeded cleanly in a preparative scale, and 8\(^{+}\) was isolated as a stable salt with 94% yield. The spiro-structure was unambiguously determined by X-ray analysis of the 8\(^{+}\)(SbCl\(_6\)\(^{-}\))·(CH\(_3\)CN) crystal (Figure 9b).

Many dicationic dyes with a biphenyl-2,2\(^{\prime}\)-diyl skeleton adopt a twisted structure similar to that of 2\(^{2+}\) with a \(\pi-\pi\) stacking arrangement, for which solvent polarity affects the twisting angle: more acute in a non-polar/less polar solvent and nearly perpendicular in a polar solvent [41,42]. The observed solvent effects for the Friedel–Crafts-type cyclization of 2\(^{2+}\) might be related to this difference in dication conformation depending on the solvent. Facile degradation to the monocation 8\(^{+}\) in CH\(_3\)CN may be related to the largely twisted conformation to facilitate nucleophilic addition of a benzene ring.
to the cationic center. The lack of further spiro cyclization of $8^+$ might be related to the ring strain of the 4,8-dihydrocyclopenta[def]fluorene structure [43].

3. Experimental

3.1. General Methods

All reactions were carried out under an argon atmosphere. All commercially available compounds were used without further purification unless otherwise indicated. Dry CH$_2$CN was obtained by distillation from CaH$_2$ prior to use. Column chromatography was performed on silica gel I-6-40 (YMC) of particle size 40–63 µm. $^1$H and $^{13}$C NMR spectra were recorded on a Ascend$^\text{TM}$ 400 ($^1$H/400 MHz and $^{13}$C/100 MHz) spectrometer (BRUKER, Billerica, MA, USA). IR spectra were measured as a KBr pellet on a JIR-WINSPEC100 FT/IR spectrophotometer (JEOL, Tokyo, Japan). Mass spectra were recorded on a JMS-T100GCV spectrometer (JEOL, Tokyo, Japan) in FD mode (GC-MS&NMR were used without further purification unless otherwise indicated. Dry CH$_3$ (29,000), 246 (49,200).

Yamato MP-21 or Yanagimoto micro melting point apparatus and are uncorrected. UV/Vis spectra were recorded on a U-3500 spectrophotometer (Hitachi, Tokyo, Japan). DFT calculations were performed with the Gaussian 09W program package (Revision E.01, 2013, Gaussian Inc., Wallingford, CT USA).

The geometries of the compounds were optimized by using the B3LYP method in combination with the 6-31G* basis set. Redox potentials ($E^\text{ox}$ and $E^\text{red}$) were measured by cyclic voltammetry in dry CH$_2$Cl$_2$ containing 0.1 M Bu$_4$NBF$_4$ as a supporting electrolyte. All of the values shown in the text are in $E/V$ versus SCE measured at a scan rate of 100 mV s$^{-1}$. Pt disk electrodes were used as the working and counter electrodes. The working electrode was polished using a water suspension of aluminum oxide (0.05 µm) before use. The irreversible half-wave potentials were estimated from the anodic peak potentials ($E_{pa}$) as $E^\text{ox} = E_{pa} - 0.03$ or the cathodic peak potentials ($E_{pc}$) as $E^\text{red} = E_{pc} + 0.03$.

3.2. Synthetic Procedures

3.2.1. Dispiro(dibenzo[a,d]cycloheptatriene-5,9'-phenanthrene-10,5'-dibenzo[a,d]cycloheptatriene) 1

From 2$^{2+}$(SbCl$_6$)$^-_2$: To a solution of 2$^{2+}$(SbCl$_6$)$^-_2$ (46.3 mg, 38.5 µmol) in dry CH$_2$CN (1.8 mL) was added activated zinc powder (50.7 mg, 775 µmol) at $25^\circ$C. The mixture was stirred for 15 min, and then diluted with water. The whole mixture was extracted with CH$_2$Cl$_2$ three times. The combined organic layers were washed with water and brine, and dried over anhydrous Na$_2$SO$_4$. After filtration, the solvent was concentrated under reduced pressure. The crude product was purified by column chromatography on silica gel (CH$_2$Cl$_2$) to give 1 (20.4 mg) as a white solid with 100% yield.

From 2$^{2+}$(BF$_4$)$^-_2$: To a solution of 2$^{2+}$(BF$_4$)$^-_2$ (265 mg, 375 µmol) in dry CH$_2$CN (8.5 mL) was added activated zinc powder (496 mg, 7.59 mmol) at $24^\circ$C. The mixture was stirred for 1 h at $24^\circ$C, and then diluted with water. The whole mixture was extracted with CH$_2$Cl$_2$ three times. The combined organic layers were washed with water and brine, and dried over anhydrous Na$_2$SO$_4$. After filtration through silica gel, the solvent was concentrated under reduced pressure to give 1 (198 mg) as a white solid with 99% yield.

Mp: 259–260 $^\circ$C (decomp.); $^1$H NMR (CDCl$_3$): $\delta$/ppm 8.10 (2H, dd, $J = 1.2, 8.2$Hz), 7.33 (2H, ddd, $J = 1.3, 7.0, 8.2$Hz), 7.02 (2H, ddd, $J = 1.2, 7.0, 8.1$Hz), 6.90 (2H, dd, $J = 1.3, 8.1$Hz), 6.83 (4H, ddd, $J = 1.1, 6.8, 7.8$Hz), 6.74 (4H, dd, $J = 1.7, 7.8$Hz), 6.43 (4H, ddd, $J = 1.7, 6.8, 8.5$Hz), 6.32 (4H, dd, $J = 1.1, 8.5$Hz), 6.08 (4H, s); $^{13}$C NMR (CDCl$_3$): $\delta$/ppm 147.13, 143.59, 137.58, 137.15, 136.80, 133.25, 131.63, 131.26, 127.29, 126.56, 125.34, 121.29, 70.50; IR (KBr): $\nu$/cm$^{-1}$ 3049, 3030, 1952, 1591, 1491, 1457, 1441, 1426, 1306, 1285, 1172, 1159, 1071, 1053, 954, 880, 870, 797, 767, 746, 725; LR-MS (FD) $m/z$ (%): 534.26 (11), 533.26 (45), 532.25 (M$^+$, bp), 267.13 (2), 266.63 (10), 266.13 (22); HR-MS (FD) Calcd. for C$_{32}$H$_{28}$: 532.21910; Found: 532.22084; UV/Vis (CH$_2$Cl$_2$): $\lambda_{max}$/nm ($\epsilon$/M$^{-1}$cm$^{-1}$) 347 (9600), 290 (29,000), 246 (49,200).
3.2.2. 5,5'-[(1,1'-Biphenyl)-2,2'-diyl]bis(5H-dibenzo[a,d]cycloheptatrien-5-yl)im bis(tetrafluoroborate) 22+(BF4)−

To a solution of diol 4 (252 mg, 445 μmol) in TFAA (4.7 mL) was added 42% HBF4 aq (670 μL, 4.45 mmol) at 0 °C to give a deep red solution, and the mixture was stirred for 2 h at 24 °C. The addition of dry ether led to precipitation of the dication salt. The precipitates were filtered and washed with dry ether three times to give 22+(BF4)− 2 (267 mg) as a red powder with 85% yield.

Mp: 209 °C (decomp.); 1H NMR and 13C NMR spectra are identical to those of 22+(SbCl6)− 2; IR (KBr): ν/cm−1 3063, 1609, 1602, 1517, 1476, 1429, 1385, 1337, 1223, 1177, 1163, 1068, 897, 850, 836, 801, 781, 751, 735; LR-MS (FD) m/z (%): 548.22 (9), 534.23 (10), 533.23 (47), 532.22 (M+, bp), 531.22 (7), 266.61 (10), 266.11 (21); HR-MS (FD) Calcd. for C42H32SbCl6: 532.21910; Found: 532.21844; UV/Vis (CH3CN): λmax/nm (Σ/M−1cm−1) 567 (4550), 534 (4330), 432 (4360), 386 (6760), 313 (76900), 269 (26200).

3.2.3. 5,5'-[(1,1'-Biphenyl)-2,2'-diyl]bis(5H-dibenzo[a,d]cycloheptatrien-5-yl)bis(hexachloroantimonate) 22+(SbCl6)− 2

To a solution of 1 (225 mg, 422 μmol) in dry CH2Cl2 (10 mL) was added (4-BrC6H4)3N•SbCl6− (680 mg, 833 μmol) at 23 °C, and the mixture was stirred for 1 h. The addition of dry ether led to precipitation of the dication salt. The precipitates were filtered and washed with dry ether three times to give 22+(SbCl6)− 2 (2CH2Cl2) (500 mg) as a red powder in 86% yield.

Mp: 258 °C (decomp.); 1H NMR (CDCl3): δ/ppm 8.98 (4H, brs), 8.49 (4H, br, J = 7.0 Hz), 8.42 (4H, br, J = 7.0 Hz), 7.80 (4H, brs), 7.61–7.53 (4H, m), 7.50 (4H, brs), 7.46 (2H, br, J = 7.4 Hz), 7.06 (2H, brd, J = 7.4 Hz); 13C NMR (CDCl3): δ/ppm 180.00, 146.66, 145.95, 141.64, 140.49, 140.15, 139.40, 138.51, 136.97, 134.76, 134.14, 132.92, 130.99, 128.48; IR (KBr): ν/cm−1 3055, 2931, 1636, 1600, 1512, 1473, 1426, 1383, 1335, 1253, 1222, 1164, 829, 847, 829, 796, 750, 730; LR-MS (FD) m/z (%): 534.26 (11), 533.26 (45), 532.25 (M+, bp), 267.13 (2), 266.63 (10), 266.13 (22); HR-MS (FD) Calcd. for C42H32SbCl6: 266.10955 (M+); Found: 266.10683 (M+); UV/Vis (CH3CN): λmax/nm (Σ/M−1cm−1) 569 (5000), 535 (4780), 425 (61700), 387 (7740), 314 (85600), 270 (43000).

3.2.4. Dispiro(dibenzo[a,d]cycloheptadiene-5,9'-phenanthena-10',5''-dibenzo[a,d]cycloheptadiene) 3

To a solution of diol 5 in HFIP (3.0 mL) was added at 22 °C a solution of TMSCIO4 in toluene (0.79 M, 1.40 mL, 1.11 mmol), which was prepared from TMSCI (500 μL, 3.96 mmol) and AgClO4 (830 mg, 4.00 mmol) in dry toluene (5 mL). After stirring at 22 °C for 1 h, the solvent was evaporated. To the residue were added dry CH2CN (8 mL) and activated zinc powder (1.78 g, 27.2 mmol). The mixture was stirred at 22 °C for 1 h, and then diluted with water. The whole mixture was extracted with CH2Cl2 three times. The combined organic layers were washed with water and brine, and dried over anhydrous Na2SO4. After filtration, the solvent was concentrated under reduced pressure. The crude product was purified by column chromatography on silica gel (hexane/CH2Cl2 = 5) to give 3 (47.5 mg) as a white solid with 33% yield.

Mp: 197–200 °C (decomp.); 1H NMR (CDCl3): δ/ppm 7.96 (2H, dd, J = 1.4, 8.2 Hz), 7.28 (2H, ddd, J = 1.2, 7.0, 8.2 Hz), 6.95–6.87 (6H, m), 6.78 (4H, d, J = 7.0 Hz), 6.44 (4H, brs), 6.29 (2H, dd, J = 1.2, 8.1 Hz), 6.11 (4H, brs), 2.92 (4H, dd, J = 9.7, 14.4 Hz), 2.57 (4H, brs); 13C NMR (CDCl3): δ/ppm 147.57, 145.21, 143.04, 137.89, 135.52, 134.28, 129.31, 127.46, 126.55, 126.35, 123.55, 121.62, 71.24, 35.78; IR (KBr): ν/cm−1 3106, 3053, 3019, 2936, 2843, 2360, 2341, 1920, 1673, 1657, 1597, 1486, 1441, 1430, 1385, 1365, 1314, 1280, 1265, 1217, 1162, 1129, 1100, 1083, 1068, 956, 866, 790, 781, 759, 746, 708, 668, 646, 625, 621, 589, 561, 534; LR-MS (FD) m/z (%): 539.26 (2), 538.26 (11), 537.26 (46), 536.26 (M+, bp), 268.13 (2); HR-MS (FD) Calcd. for C42H32SbCl6: 536.25040, Found: 536.25269.

3.2.5. 2,2'-Bis(5-hydroxydibenzo[a,d]cycloheptatrien-5-yl) biphenyl 4 [37]

To a solution of 2,2'-diiodobiphenyl [44] (1.13 g, 2.78 mmol) in dry THF (10 mL) was added nBuLi (1.55 M in hexane, 4.3 mL, 6.67 mmol) dropwise over 2 min at −78 °C. After stirring at −78 °C for 1 h,
dibenzotropone (1.38 g, 6.67 mmol) was added to the suspension and the mixture was warmed to 23 °C. The resulting solution was stirred for 1 h at 23 °C, and then diluted with water. The whole mixture was extracted with EtOAc three times. The combined organic layers were washed with water and brine, and dried over anhydrous Na₂SO₄. After filtration, the solvent was concentrated under reduced pressure. The crude product was washed with methanol to give diol 4 (1.12 mg) as a white solid with 71% yield.

³¹H NMR (CDC1₃): δ/ppm 8.13 (2H, dd, J = 1.2, 7.9 Hz), 7.50 (2H, ddd, J = 1.8, 6.8, 7.9 Hz), 7.39 (2H, dd, J = 1.3, 8.0 Hz), 7.30 (2H, dt, J = 1.2, 6.8 Hz), 7.02 (2H, dd, J = 1.8, 6.8 Hz), 6.98 (2H, dd, J = 1.3, 7.3 Hz), 6.86 (2H, dt, J = 1.3, 7.3 Hz), 6.79 (2H, ddd, J = 1.3, 7.3, 8.0 Hz), 6.67 (4H, s), 6.61 (2H, ddd, J = 1.5, 7.5, 8.7 Hz), 6.46 (2H, dt, J = 1.2, 7.5 Hz), 6.39 (2H, dd, J = 1.2, 8.7 Hz), 6.05 (2H, dd, J = 1.5, 7.5 Hz), 5.04 (2H, s).

3.2.6. 2,2'-Bis(5-hydroxydibenzo[a,d]cycloheptadiene-5-yl)biphenyl 5

To a solution of 2,2'-diiodobiphenyl (409 mg, 1.01 mmol) in dry THF (5 mL) was added nBuLi (1.55 M in hexane, 1.55 mL, 2.40 mmol) dropwise over 2 min at –78 °C. After stirring at –78 °C for 1 h, dibenzoosuberone (432 µL, 2.40 mmol) was added to the suspension, and the mixture was warmed to 25 °C. The resulting solution was stirred for 15 h at 25 °C, and then diluted with water. The whole mixture was extracted with EtOAc three times. The combined organic layers were washed with water and brine, and dried over anhydrous Na₂SO₄. After filtration, the solvent was concentrated under reduced pressure. The crude product was washed with methanol to give 5 (467 mg) as a white solid with 81% yield.

³¹H NMR (CDC1₃): δ/ppm 8.13 (2H, dd, J = 1.5, 7.6 Hz), 7.25 (2H, dt, J = 1.5, 7.6 Hz), 7.20 (2H, dt, J = 1.5, 7.6 Hz), 7.05 (2H, dd, J = 1.5, 7.6 Hz), 7.02 (2H, dd, J = 1.5, 7.6 Hz), 6.96–6.84 (8H, m), 6.66 (2H, dt, J = 1.5, 7.6 Hz), 6.44 (2H, dt, J = 1.4, 7.5 Hz), 6.00 (2H, dd, J = 1.4, 7.5 Hz), 5.15 (2H, s), 3.07–2.86 (4H, m), 2.70–2.60 (2H, m), 2.48–2.39 (2H, m).

3.2.7. 5-(Spiro[dibenzo[a,d]cycloheptatriene-5,9'-fluoren]-4'-yl)-5H-dibenzo[a,d]cycloheptatriene-5-ylium hexachloroantimoniate 8*(SbCl₆⁻)

A solution of 2²⁺(SbCl₆⁻)₂ (83.9 mg, 69.8 µmol) in dry CH₃CN (3 mL) was stirred for 2 days at 22 °C. The addition of dry ether led to precipitation of the monocation salt. The precipitates were filtered and washed with dry ether three times to give 8*(SbCl₆⁻) (56.7 mg) as a red powder with 94% yield.

Mp: 218 °C (decomp.); ¹³C NMR (acetone-d₆): δ/ppm 181.17, 154.51, 154.39, 147.39, 145.88, 141.66, 141.23, 138.85, 138.12, 138.16, 136.78, 136.75, 136.66, 136.02, 135.89, 134.09, 133.20, 132.88, 129.89, 128.78, 128.73, 128.46, 128.34, 128.16, 127.96, 127.84, 127.32, 122.72, 66.00; IR (KBr): ν/cm⁻¹ 3061, 2935, 2859, 2360, 1600, 1513, 1479, 1428, 1385, 1337, 1258, 1177, 892, 848, 833, 801, 752, 734; LR-MS (FD) m/z (%): 588.24 (6), 566.19 (6), 533.23 (11), 532.23 (47), 531.22 (M⁺, bp); HR-MS (FD) Calcd. for C₃₂H₂₇S: 531.21128; Found: 531.21124; UV/Vis (CH₂Cl₂): λ_max/µm (Σ/M⁻¹·cm⁻¹) 567 (5200), 533 (4680), 406 (6310), 388 (7980), 315 (104,500), 275 (35,800), 242 (46,700).

3.3. Crystal Data

Data were collected with a Rigaku Mercury 70 diffractometer (Mo-Kα radiation, λ = 0.71075 Å). The structure was solved by the direct method (SIR2004) and refined by the full-matrix least-squares method on F² with anisotropic temperature factors for non-hydrogen atoms. All the hydrogen atoms were located at the calculated positions and refined with riding.
1: Crystals were obtained by recrystallization from CHCl3/hexane. MF: C42H28, FW: 532.68, colorless block, 0.20 × 0.20 × 0.20 mm3, monoclinic P21/c, a = 13.3661(12) Å, b = 12.6558(11) Å, c = 16.1662(2) Å, β = 95.8567(13)°, V = 27220.4(5) Å3, ρ (Z = 4) = 1.301 g cm−3. A total of 20359 reflections (2θ(max) = 55.0°) were measured at T = 150 K. Numerical absorption correction was applied (μ = 0.735 cm−1). The final R1 and wR2 values are 0.0439 (I > 2σI) and 0.1179 (all data) for 5332 reflections and 379 parameters. Estimated standard deviations are 0.0017–0.003 Å for bond lengths and 0.09–0.17° for bond angles. CCDC 1580196.

2+[SbCl6]2−·2(CH2Cl2): Crystals were obtained by recrystallization from CH2Cl2/ether. MF: C44H32Cl6Sb2, FW: 1371.48, red platelet, 0.60 × 0.10 × 0.10 mm3, monoclinic P21/c, a = 11.109(2) Å, b = 15.074(3) Å, c = 30.901(6) Å, β = 91.693(3)°, V = 5172(2) Å3, ρ (Z = 4) = 1.761 g cm−3. A total of 31500 reflections (2θ(max) = 55.0°) were measured at T = 150 K. Numerical absorption correction was applied (μ = 19.007 cm−1). The final R1 and wR2 values are 0.0983 (I > 2σI) and 0.2428 (all data) for 8942 reflections and 559 parameters. Estimated standard deviations are 0.004–0.03 Å for bond lengths and 0.13–0.8° for bond angles. CCDC 1580197.

3: Crystals were obtained by recrystallization from benzene/hexane. MF: C42H32, FW: 536.71, colorless prism, 0.40 × 0.30 × 0.20 mm3, monoclinic P21/n, a = 16.557(6) Å, b = 20.113(7) Å, c = 17.343(6) Å, β = 103.513(5)°, V = 5616(4) Å3, ρ (Z = 8) = 1.270 g cm−3. A total of 42292 reflections (2θ(max) = 55.0°) were measured at T = 150 K. Numerical absorption correction was applied (μ = 0.716 cm−1). The final R1 and wR2 values are 0.0679 (I > 2σI) and 0.1926 (all data) for 11017 reflections and 757 parameters. Estimated standard deviations are 0.003–0.005 Å for bond lengths and 0.15–0.3° for bond angles. CCDC 1580198.

8+[SbCl6]2−·(CH2CN): Crystals were obtained by recrystallization from CH2CN/ether. MF: C44H30Sb6Cl6, FW: 907.20, red platelet, 0.10 × 0.10 × 0.50 mm3, triclinic PÍ, a = 9.5187(8) Å, b = 13.5559(10) Å, c = 15.433(11) Å, α = 98.195(9)°, β = 92.202(13)°, γ = 102.889(11)°, V = 1916(2) Å3, ρ (Z = 2) = 1.572 g cm−3. A total of 14621 reflections (2θ(max) = 55.0°) were measured at T = 150 K. Numerical absorption correction was applied (μ = 11.707 cm−1). The final R1 and wR2 values are 0.0415 (I > 2σI) and 0.1049 (all data) for 7454 reflections and 454 parameters. Estimated standard deviations are 0.0013–0.007 Å for bond lengths and 0.09–1.2° for bond angles. CCDC 1580199.

4. Conclusions

Based on a molecular design that maximizes the effects of TBI, the novel Ar4DHP 1 with two spiro(dibenzocycloheptatriene) units was designed as an electron-donating hydrocarbon. Based on reversible interconversion with the bis(dibenzo[b]tritylum)-type dicationic dye 2+ accompanied by C–C bond formation/cleavage (dyrex behavior), the present redox pair exhibits electrochromism with a vivid change in color, and thus represents a class of less well-developed hydrocarbon-based systems. Compound 1 was predicted to have a highly strained structure during the molecular design by a DFT calculation, which was finally confirmed by X-ray analysis of the hydrocarbon prepared as a stable entity. A more highly strained structure was observed in the tetrahydro derivative 3, the C9–C10 bond length of which is the greatest among the values ever reported for Ar4DHPs. Thus, the spiro(dibenzocycloheptadiene) unit was proven to be a promising spiro skeleton for inducing extreme bond expansion when incorporated in the clamped/fused HPE skeleton.

**Supplementary Materials:** Supplementary materials are available online, Figure S1: Change in the UV/Vis spectrum upon electrochemical reduction (20 μA) of as-prepared 22+ in CH2Cl2 containing 0.05 M Bu4NBF4 as a supporting electrolyte (every 4 min).

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**Sample Availability:** Samples of the compounds 1–8 are available from the authors.

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