Transformation of Thia[7]helicene to Aza[7]helicenes and [7]Helicene-like Compounds via Aromatic Metamorphosis

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Abstract: [n]Helicenes with helically twisted structures have attracted increasing interest owing to their unique properties. Therefore, it has been an important issue to develop facile synthetic methodologies which allow access to a variety of [n]helicenes. Here we report the synthesis of [7]helicenes and [7]helicene-like compounds from the thia[7]helicene as a common starting material. Desulfurative dilithiation of the thia[7]helicene and the subsequent reaction with silicon and phosphorus electrophiles afforded the silole- and phosphole-fused [7]helicene-like compounds, respectively. The cyclopentadiene-fused [7]helicene-like compound and the pyrrole-fused aza[7]helicenes were also successfully synthesized via twofold SnAr reactions of the thia[7]helicene S,S-dioxide with the carbon and nitrogen nucleophiles, respectively. The thia[7]helicene S,S-dioxide showed a slightly red-shifted absorption spectrum than the parent thia[7]helicene, which was well demonstrated by the theoretical calculations. The substituents on the silicon atom of silole-fused [7]helicene-like compounds have little impact on the longest absorption maximum. Such little effect of the substituents on absorption properties was also observed for cyclopentadiene-fused [7]helicene-like compounds andaza[7]helicenes and was well demonstrated by the theoretical calculations. The thia[7]helicene S,S-dioxide and the silole-fused [7]helicene-like compound exhibited bright blue emission, and the cyclopentadiene-fused [7]helicene-like compound and the aza[7]helicenes showed strong violet emission. Each single enantiomer of the aza[7]helicenes showed circularly-polarized luminescence with the dissymmetry factors of 4.2 × 10⁻³.

Keywords: [n]helicenes; aromatic metamorphosis; desulfurative lithiation; nucleophilic aromatic substitution; thiophene; silole; phosphole; pyrrole; circular dichroism; circularly polarized luminescence

1. Introduction

[n]Helicenes are ortho-fused polycyclic aromatic compounds in which all rings are angularly arranged to form screw-shaped helical structures [1–3]. Those composed of six or more benzene rings exhibit stable helical chirality owing to an intramolecular steric repulsion. The π-extended, nonplanar, and chiral structures of [n]helicenes endow them with unique properties and offer promising applications in various fields such as nonlinear optics [4], organic semiconductors [5,6], organic light-emitting materials [7], catalysts [8,9], and circularly polarized luminescent materials [10]. The prototypical [n]helicene is carbo-helicenes whose aromatic rings consist of only carbon atoms. Another class of [n]helicenes is heterohelicenes with one or more heteroaromatic rings, such as pyrrole, furan, thiophene, and pyridine, in the helical skeletons. Besides these [n]helicenes composed of only aromatic rings, helicene-like compounds with non-aromatic rings, such as silole and phosphole, in the helical skeletons have also been developed. Heterohelicenes and helicene-like compounds have received growing interest because the introduction of heteroatom(s)
into the π-conjugated framework is effective for controlling the electronic structure of the π-conjugated molecules and providing unique functions.

A variety of synthetic methodologies for helicenes have been developed. Photocyclization of the stilbene skeleton has been a conventional and common one [11]. Diels–Alder reaction [12], Friedel–Crafts-type reaction [5,13], transition metal-catalyzed [2 + 2 + 2] cycloaddition [14–18], cross-coupling reaction [19], and olefin metathesis [20,21] have been found to be effective for helicene synthesis during the last decades. These synthetic methodologies have undoubtedly contributed to a great progress in helicene chemistry. However, they often require lengthy multistep synthesis, which would be disadvantageous for structural diversity of helicenes. Therefore, there is still a tremendous need for developing more diversity-oriented synthetic methodologies.

Recently, it has been reported that dibenzothiophenes can be converted into a variety of ring systems, such as dibenzosiloles, dibenzophospholes, fluorenes, and carbazoles [22–26]. Transformation to dibenzosiloles and dibenzophospholes involves (1) the generation of 1,1′-dianions via desulfurative dilithiation with lithium metal and (2) the subsequent reaction of the resulting 1,1′-dianions with silicon and phosphorus electrophiles, respectively. On the other hand, dibenzothiophenes are converted into fluorenes and carbazoles via (1) oxidation to S,S-dioxides and (2) the consecutive inter- and intramolecular S_N_Ar reactions with carbon and nitrogen nucleophiles. This methodology named as “aromatic metamorphosis” would be a powerful tool for synthesizing π-extended compounds since a wide range of compounds containing the dibenzothiophene unit have been reported. Indeed, [8]circulenes [27,28] and spiro π-conjugated compounds [29] have been prepared by using this methodology. Yorimitsu and co-workers have also reported the transformation of dithia[8]helicene to its carbazole and fluorene analogs [30].

In this context, we envisaged that a series of silole-, phosphole-, and cyclopentadiene-fused [7]helicene-like compounds 2–4 and pyrrole-fused aza[7]helicenes 5 can be synthesized from a single thia[7]helicene, diphenanthro[3,4-b:4′,3′-d]thiophene (1a) [31,32] as a common starting material (Figure 1a). We have previously reported the synthesis of 2a, 3a, and 5a (Figure 1b) [33–36]. Each heteroatom and the bridging carbon was introduced at the early stage in the synthetic sequence. In addition, all the synthetic routes require costly transition metal catalysts. In contrast, the synthetic routes in this work are transition metal-free and can provide a series of [7]helicene-like compounds and aza[7]helicenes via late-stage diversification of thia[7]helicene 1a, therefore achieving more diversity-oriented synthesis. Aromatic metamorphosis of the dibenzothiophene unit to carbazole and fluorene units has been applied to the helicene synthesis as described above [30], but its application to silole- and phosphole-fused helicene-like compounds has never been demonstrated. Herein we report the synthesis of a series of [7]helicene-like compounds and aza[7]helicenes through aromatic metamorphosis. Their photophysical properties are also discussed through comparison with the previously reported analogs.
2. Results and Discussion

Aromatic metamorphosis of thia[7]helicene 1a to its silole and phosphole oxide analogs 2 and 3, respectively, was first investigated (Scheme 1). The desulfurative dilithiation of 1a with lithium metal followed by the trapping with MeI has been examined previously [31]. In the literature, the desired methylated compound was not obtained, and the main product was found to be the reduced compound, 4,4′-biphenanthrene. On the other hand, we have recently reported that the dianion formed by the reaction of 3,3′-dibromo-4,4′-biphenanthrene with BuLi can react with SiCl4 to give the double [7]helicene-like compound with a silicon spiro center [37]. Encouraged by the recent progress in aromatic metamorphosis of dibenzothiophenes [26] and our recent result, we conducted the reaction of thia[7]helicene 1a with 5 equivalents of lithium metal in the presence of 2 equivalents of TMEDA in THF at room temperature, which is the optimized conditions in the recent report on aromatic metamorphosis (Scheme 1). To our delight, the subsequent reaction with commercially available Cl2SiMe2 successfully afforded the silole-fused [7]helicene-like compound 2a in 48% yield. In addition, the reaction with Cl2SiPh2 proceeded to give the new silole-fused [7]helicene-like compound 2b in 41% yield. Phosphorous electrophile was also found to be applicable, and the phosphole oxide-fused [7]helicene-like compound 3 was obtained through the reaction of the dianion with Cl2PPh and the following oxidation.
Transformation of thia[7]helicene S,S-dioxide 1b, which can be easily prepared from 1a by oxidation, was also examined (Scheme 2). According to the literature in which dibenzothiophene S,S-dioxides are converted into fluorenes [24], the reaction of 1b with xanthene, a commercially available carbon nucleophile, was conducted in the presence of KN(SiMe$_3$)$_2$ as a base. Although the yield is unexpectedly low, the desired spirocyclic carbo[7]helicene-like compound 4b was obtained (23% yield). Next, the transformation of 1b to aza[7]helicenes 5 was investigated. Arylamines with electron-donating groups, such as p-toluidine and p-anisidine, reacted with 1b smoothly, affording the corresponding aza[7]helicenes 5b and 5c, respectively, in high yields (80% for 5b and 88% for 5c). The reaction with electron-deficient arylamines, such as p-(trifluoromethyl)aniline and 4-aminobenzonitrile, resulted in no production of the aza[7]helicenes, which would be due to low nucleophilicity of these arylamines [22].

The UV–vis absorption and photoluminescence (PL) spectra of new silole- and cyclopentadiene-fused [7]helicene-like compounds 2b and 4b and pyrrole-fused aza[7]helicenes 5b and 5c were measured to evaluate their photophysical properties (Figure 2). Thiophene-fused thia[7]helicene 1a and its S,S-dioxide 1b are known compounds, but only the absorption property of 1a in cyclohexane has been reported. Therefore, their absorption and fluorescence spectra in CH$_2$Cl$_2$ were also measured. The photophysical data are summarized in Table 1, together with those of the previously reported 2a, 4a, and 5a for comparison. We also performed theoretical calculations by density functional theory (DFT) and time-dependent (TD) DFT methods at the B3LYP/6-31G(d) level of theory to understand the experimental photophysical properties.
Figure 2. UV−vis absorption (solid line) and PL (broken line) spectra of (a) 1a and 1b in CH₂Cl₂ (1.0 × 10⁻⁵ M) and (b) 2b, 4b, and 5b in CH₂Cl₂ (1.0 × 10⁻⁵ M). The spectra of 5c are omitted since they are almost the same as those of 5b.

Table 1. Photophysical properties of [7]helicene-like compounds and aza[7]helicenes.

|    | λ_{abs} (nm) | λ_{em} (nm) | φ  |
|----|--------------|-------------|----|
| 1a | 401          | 417, 435 (sh) (265) | <1 |
| 1b | 395, 415 (sh) | 458 (300)    | 4  |
| 2b | 389, 411 (sh) | 434 (sh), 454 (325) | 11 |
| 4b | 406          | 422, 441 (sh) (294) | 31 |
| 5b | 401          | 421, 444 (335) | 17 |
| 5c | 401          | 422, 445 (335) | 16 |
| 2a | 412          | 450 (320)    | 23 |
| 4a | 408          | 421, 441 (sh) (340) | 40 |
| 5a | 400          | 418, 442 (330) | 17 |

a The longest absorption maximum in CH₂Cl₂. sh: shoulder. b Emission maximum in CH₂Cl₂. Excitation wavelength in parenthesis. c Absolute quantum yield in CH₂Cl₂. d Reference [35]. e Reference [36]. f Reference [33].

Thia[7]helicene 1a showed the longest absorption maximum (λ_{abs}) at 401 nm and an emission maximum (λ_{em}) at 417 nm with a shoulder at 435 nm (Figure 2a). Its fluorescence intensity is weak, and the absolute quantum yield (φ) was estimated to be <1%. The S,S-dioxide compound 1b gave a slightly red-shifted absorption band with a shoulder at 415 nm. The emission maximum (458 nm) is significantly red-shifted with a higher absolute quantum yield (4%), compared to those of 1a, exhibiting a bright blue emission. The TD–DFT calculations demonstrated a slightly red-shifted absorption characteristic of 1b (Table S11 in the supplementary material). The longest absorption wavelengths of 1a and 1b were estimated to be 387 nm and 406 nm, respectively. The energy levels of the HOMO and HOMO−1 of 1a are almost identical (−5.42 eV and −5.45 eV), and the HOMO−1 → LUMO transition mainly contributes to the longest-wavelength absorption band of 1a. On the other hand, the HOMO → LUMO transition is the main contribution to the longest-wavelength absorption band of 1b. The HOMOs of both compounds are delocalized over the entire skeleton, but without any contribution from the sulfur atoms (Figure 3). The HOMO energy level of 1b is 0.40 eV lower than that of 1a owing to the electron-withdrawing effect of the S,S-dioxide moiety. The LUMOs are delocalized over the entire skeleton including the sulfur atoms. It is notable that the difference between LUMO energy levels (0.62 eV) is larger than that between the HOMO energy levels. It may be due to the effective σ*−π* conjugation between the exocyclic S–O σ* orbitals and the endocyclic butadiene π* orbital of the thiophene unit [38], effectively lowering the LUMO energy level of 1b.
Figure 3. Frontier molecular orbitals of 1a, 1b, 2b, 4b, and 5b calculated by DFT method at the B3LYP/6-31G(d) level of theory.

The silole-fused [7]helicene-like compound 2b exhibited a relatively broad absorption spectrum with a shoulder at 411 nm and an absorption edge at 434 nm, which are almost the same as those of the dimethyl analog 2a (λ_{abs}: 412 nm; absorption edge: 431 nm) [35]. The absorption wavelengths estimated by TD–DFT calculation (392 nm for 2a; 396 nm for 2b) agree with these experimental absorption characteristics (Table S11 in supplementary material). The features of a silole are well reflected in the LUMO of 2b, as observed for 2a. The LUMO is distributed over the silicon atom owing to the σ*–π* conjugation between Si–C(phenyl)σ* orbitals and the butadiene π* orbital of the silole unit (Figure 3). Except for this effect, the phenyl substituents of 2b have no obvious contribution in the frontier orbitals. Therefore, the molecular orbital distributions in the HOMOs and LUMOs of 2a and 2b are almost the same as each other, giving almost the same λ_{abs}. A bright blue emission was observed for 2b (λ_{em}: 454 nm), while the absolute quantum yield of 2b (11%) is much lower than that of 2a (23%).

Photophysical properties of the cyclopentadiene-fused [7]helicene-like compound 4b and the pyrrole-fused aza[7]helicenes 5b and 5c were also found to be similar to those of their analogs 4a and 5a, respectively. All compounds exhibit strong violet emission at 421–422 nm. As indicated by theoretical calculations (Table S11 in supplementary material), the bridging of the two phenyl groups in 4a by an oxygen atom and the introduction of the electron-donating groups at para-position of N-phenyl group in 5a have little impact on their electronic structure. The compound 4b contains a spirocyclic structure, but a through-space orbital interaction known as spiroconjugation [39–41] between the 4,4′-biphenanthrene and diphenyl ether units is not observed.

Fortunately, we found the optical resolution conditions for aza[7]helicenes 5b and 5c by using chiral HPLC. In addition, the absolute configuration of the obtained single enantiomer of 5c was unambiguously identified by single-crystal X-ray analysis (Figure 4). The chiroptical properties of 5b and 5c were evaluated by CD and CPL spectroscopies (Figure 5). Both 5b and 5c showed mirror-image CD spectra with a Cotton effect in the longest absorption region of their absorption spectra. The (P)-isomer of 5c showed a negative Cotton effect at its longest absorption band along with two positive Cotton effects at around 340 nm and 280 nm. Such a characteristic of the CD spectrum of (P)-5c is identical to that of (P)-5a we have previously reported [33]. Based on these facts, we assigned the absolute configuration of the single enantiomer of 5b, which showed a negative Cotton effect at around 340 nm, to be P. The dissymmetry factors in absorption (g_{abs}) of (P)-5b and (P)-5c were estimated to be $-3.8 \times 10^{-3}$ and $-4.0 \times 10^{-3}$, respectively, at the maximum of the first Cotton effects. Enantiomers of 5b and 5c exhibited clear CPL signals. The CPL sign from (P)-enantiomers of them is negative. The dissymmetry factors in emission (g_{lum}) of
(P)-5b and (P)-5c were $-4.2 \times 10^{-3}$ (429 nm) and $-4.4 \times 10^{-3}$ (432 nm), respectively, which are comparable to those of the previously reported chiral small organic molecules [42].

(Figure 4. ORTEP drawing of (M)-5c with thermal ellipsoids at 50% probability. All hydrogen atoms and the crystal solvent molecule (CHCl$_3$) are omitted for clarity.)

(Figure 5. CD (in CH$_2$Cl$_2$) and CPL (in 2-Me-THF) spectra of (a) 5b and (b) 5c. The $g_{\text{abs}}$ and $g_{\text{lum}}$ values were calculated at the longest CD maximum and the CPL maximum, respectively. Wavelength for the calculated values in parentheses.)

3. Materials and Methods

3.1. General Procedures

All manipulations involving air- and/or moisture-sensitive compounds were carried out with the standard Schlenk technique under argon. Analytical thin-layer chromatography was performed on a glass plate coated with 0.25-mm 230–400-mesh silica gel containing a fluorescent indicator. Column chromatography was performed by using silica gel (spherical neutral, particle size: 63–210 µm). The recycling preparative HPLC was performed with YMC-GPC T-2000 and T-4000 columns (chloroform as an eluent) (YMC Co., Ltd., Kyoto, Japan). Most of the reagents were purchased from commercial suppliers, such as Sigma-Aldrich Co. LLC (St. Louis, MO, USA), Tokyo Chemical Industry Co., Ltd. (Tokyo, Japan), and Kanto Chemical Co., Inc. (Tokyo, Japan), and used without further purification unless otherwise specified. Commercially available anhydrous solvents were used for air- and/or moisture sensitive reactions. Diphenanthro[3,4-b:4',3'-d]thiophene (1a) and diphenanthro[3,4-b:4',3'-d]thiophene S,S-dioxide (1b) were prepared according to the literature [31].

NMR spectra were recorded in CDCl$_3$ on a JEOL–ECX400 spectrometer ($^1$H 400 MHz; $^{13}$C 101 MHz) or JEOL–ECA500 spectrometer ($^1$H 500 MHz; $^{13}$C 126 MHz) (JEOL Ltd., Tokyo, Japan). Chemical shifts are reported in ppm relative to the internal standard signal (0 ppm for Me$_4$Si in CDCl$_3$) for $^1$H and the deuterated solvent signal (77.16 ppm for CDCl$_3$) for $^{13}$C. Data are presented as follows: chemical shift, multiplicity (s = singlet, d = doublet, t = triplet, m = multiplet and/or multiple resonances), coupling constant in hertz (Hz), and signal area integration in natural numbers. High resolution mass spectra are taken with a Bruker Daltonics microOTOF-QII mass spectrometer (Bremen, Germany) by
atmospheric pressure chemical ionization-time-of-flight (APCI–TOF) method. UV–Vis absorption spectra were recorded on a JASCO V–650 spectrophotometer (JASCO Corporation, Tokyo, Japan). Photoluminescence spectra were recorded on a JASCO FP–6500 spectrophotometer (JASCO Corporation, Tokyo, Japan). Absolute quantum yields were determined by absolute quantum yield measurement system with a JASCO ILF–533 integrating sphere (JASCO Corporation, Tokyo, Japan). HPLC analyses and optical resolution were carried out using a DAICEL CHIRALPAK® IA-3 column (Daicel Corporation, Tokyo, Japan) (4.6 mm × 250 mm) and a DAICEL CHIRALPAK® IA column (Daicel Corporation, Tokyo, Japan) (20 mm × 250 mm), respectively. Circular dichroism (CD) spectra were recorded on a JASCO J–725 spectrometer (JASCO Corporation, Tokyo, Japan). CPL spectra were measured by using a JASCO CPL–300 spectrometer (JASCO Corporation, Tokyo, Japan).

3.2. Synthesis

3.2.1. 9,9-Dimethyl-9H-diphenanthro[3,4-b:4’,3’-d]silole (2a)

A 20-mL Schlenk tube was charged with lithium (7 mg, 1.0 mmol), THF (1.0 mL), N,N,N’,N’-tetramethylethylenediamine (TMEDA) (60 µL, 0.40 mmol) under argon atmosphere. After the mixture was stirred at room temperature for 10 min, diphenanthro[3,4-b:4’,3’-d]thiophene (1a) (77 mg, 0.20 mmol) was added. The resulting mixture was stirred vigorously at room temperature for 4 h, and then cooled to −78 °C. After adding Cl3SiMe2 (48 µL, 0.40 mmol) under argon atmosphere, the resulting mixture was stirred at −78 °C for 15 min and at room temperature for an additional 1 h. The reaction was quenched with water at 0 °C, and the resulting mixture was extracted with AcOEt (10 mL × 3). The combined organic layers were washed with brine, dried over Na2SO4, filtered, and concentrated under reduced pressure. The crude residue was purified by silica-gel column chromatography (hexane as an eluent) and then by recycling preparative HPLC to give the title compound 2a as a yellow solid (39 mg, 48% yield). The 1H NMR data were identical to that reported in the literature [35].

3.2.2. 9,9-Diphenyl-9H-diphenanthro[3,4-b:4’,3’-d]silole (2b)

The crude product was obtained by using lithium (14 mg, 2.0 mmol), THF (4 mL), TMEDA (0.12 mL, 0.80 mmol), diphenanthro[3,4-b:4’,3’-d]thiophene (1a) (154 mg, 0.40 mmol), and Cl3SiPh2 (0.17 mL, 0.80 mmol) at room temperature for 19 h. Purification by silica-gel column chromatography (CH2Cl2/hexane = 7/3 as an eluent; Rf = 0.82) and then by recycling preparative HPLC gave the title compound 2b as a yellow solid (88 mg, 41% yield): 1H NMR (400 MHz, CDCl3) δ 7.95 (d, J = 7.3 Hz, 2H), 7.79 (d, J = 7.3, 2H), 7.77–7.75 (m, 5H), 7.65 (d, J = 8.7 Hz, 2H), 7.52 (d, J = 8.7 Hz, 2H), 7.43–7.35 (m, 8H), 7.32 (d, J = 7.8 Hz, 2H), 6.94 (t, J = 7.8 Hz, 2H), 6.37 (t, J = 8.2 Hz, 2H); 13C NMR (101 MHz, CDCl3) δ 149.0, 136.6, 135.8, 134.6, 132.8, 132.1, 130.4, 130.3, 130.0, 129.1, 128.4, 128.1, 127.8, 127.6, 126.5, 126.2, 125.1, 123.1; HRMS–APCI+ (m/z) calcd for C40H27Si+ ([M+H]+) 535.1877, found 535.1876.

3.2.3. 9-Phenylnaphtho[1,2-e]phenanthro[3,4-b]phosphindole 9-oxide (3)

A 20-mL Schlenk tube was charged with lithium (7 mg, 1.0 mmol), THF (2 mL), N,N,N’,N’-tetramethylethylenediamine (TMEDA) (60 µL, 0.40 mmol) under argon atmosphere. After the mixture was stirred at room temperature for 10 min, diphenanthro[3,4-b:4’,3’-d]thiophene (1a) (77 mg, 0.20 mmol) was added. The resulting mixture was stirred vigorously at room temperature for 4 h, and then cooled to −78 °C. After adding Cl2PPh (54 µL, 0.40 mmol) under argon atmosphere, the resulting mixture was stirred at −78 °C for 10 min and at room temperature for an additional 30 min. The reaction mixture was concentrated under reduced pressure. The resulting residue was dissolved with CH2Cl2 (10 mL) and passed through a short pad of neutral alumina under argon atmosphere. The alumina pad was washed with CH2Cl2 (5 mL), and the combined filtrates were concentrated under reduced pressure. The resulting residue was dissolved in CH2Cl2 (5 mL), and H2O2 (0.40 mL, 35% aqueous solution) was added in one portion. After the resulting mixture was stirred at room temperature for 30 min, saturated aqueous Na2S2O3 (5 mL) was added
slowly to the reaction mixture. The organic layer was separated, and the aqueous layer was extracted with AcOEt (10 mL × 2). The combined organic layers were dried over Na2SO4, filtered, and concentrated under reduced pressure. The resulting residue was purified by silica-gel column chromatography (CH2Cl2/AcOEt = 4/1 as an eluent, Rf = 0.25) and then by recycling preparative HPLC to give the title compound 3 as a yellow solid (32 mg, 34% yield). The 1H NMR data were identical to that reported in the literature [34].

3.2.4. Spiro[cyclopenta[1,2-c:4,3-c]diphenanthrene-9,9′-xanthene] (4b)

A 20-mL Schlenk tube was charged with diphenanthro[3,4-b:4′,3′-d]thiophene S,S-dioxide (1b) (50 mg, 0.12 mmol), xanthene (27 mg, 0.18 mmol), and 1,4-dioxane (2.5 mL) under argon atmosphere. A solution of KN(SiMe3)2 (0.5 M in toluene, 0.60 mL, 0.30 mmol) was added to the mixture at an ambient temperature. The resulting solution was stirred at 80 °C for 16 h. After the reaction was quenched with saturated aqueous NH4Cl solution (3 mL), the resulting mixture was extracted with CH2Cl2 (10 mL × 3). The combined organic layers were dried over Na2SO4, filtered, and concentrated under reduced pressure. The crude residue was dissolved in CH2Cl2 and passed through a short pad of silica gel. The filtrate was concentrated, and the resulting crude residue was purified by recycling preparative HPLC to give the title compound 4b as a pale-yellow solid (15 mg, 23% yield): 1H NMR (500 MHz, CDCl3) δ 7.81 (d, J = 7.5 Hz, 2H), 7.76–7.71 (m, 6H) 7.65 (d, J = 8.0 Hz, 2H), 7.47 (d, J = 8.0 Hz, 2H), 7.30 (dd, J = 8.0, 1.2 Hz, 2H), 7.23–7.20 (m, 2H), 7.14 (t, J = 7.5 Hz, 2H), 6.73–6.70 (m, 2H), 6.52 (dd, J = 7.7, 1.2 Hz, 2H), 6.36–6.33 (m, 2H); 13C NMR (101 MHz, CDCl3) 156.3, 152.0, 136.7, 132.6, 131.8, 129.6, 128.0, 128.5, 128.2, 127.7, 127.6, 127.0, 126.9, 126.8, 126.6, 123.7, 123.5, 123.4, 117.2, 54.49; HRMS–APCI+ (m/z) calcd for C49H35O+ ([M+H]+) 533.1900, found 533.1900.

3.2.5. 9-(4-Methylphenyl)-9H-dinaphtho[2,1-c′:1′,2′-g]carbazole (5b)

A 20-mL Schlenk tube was charged with diphenanthro[3,4-b:4′,3′-d]thiophene S,S-dioxide (1b) (63 mg, 0.15 mmol), p-toluidine (32 mg, 0.30 mmol), and 1,4-dioxane (2.5 mL) under argon atmosphere. A solution of KN(SiMe3)2 (0.5 M in toluene, 0.90 mL, 0.45 mmol) was added to the mixture at an ambient temperature. The resulting solution was stirred at 80 °C for 15 h. After the reaction was quenched with saturated aqueous NH4Cl solution (2 mL), the resulting mixture was extracted with ethyl acetate (10 mL × 3). The combined organic layers were washed with brine (10 mL), dried over Na2SO4, filtered, and concentrated under reduced pressure. The crude residue was purified by silica-gel column chromatography (hexane/ACOEt = 10/1 as an eluent; Rf = 0.40) to give the title compound 5b as a pale-yellow solid (55 mg, 80% yield). The compound 5b can be separated into enantiomerically-pure (P)-5b and (M)-5b by HPLC equipped with a DAICEL CHIRALPAK® IA-3 column (Daicel Corporation, Tokyo, Japan) (4.6 mm × 250 mm) [tR = 4.24 min for (P)-5b and 5.23 min for (M)-5b (flow rate: 1.0 mL; eluent: Hex/CHCl3 = 7/3)]; 1H NMR (500 MHz, CDCl3) δ 8.03 (d, J = 8.6 Hz, 2H), 8.01 (d, J = 8.6, 2H), 7.844 (d, J = 8.6 Hz, 2H), 7.835 (d, J = 7.5 Hz, 2H), 7.78 (d, J = 8.6 Hz, 2H), 7.63–7.61 (m, 2H), 7.53 (d, J = 8.0 Hz, 2H), 7.50 (d, J = 8.6 Hz, 2H), 7.22–7.18 (m, 2H), 6.28–6.25 (m, 2H); 13C NMR (126 MHz, CDCl3) δ 140.4, 138.7, 134.6, 131.5, 130.9, 130.5, 128.5, 128.3, 127.4, 127.1, 126.9, 126.8, 126.4, 126.1, 124.6, 122.8, 117.2, 110.9, 21.5; HRMS–APCI+ (m/z) calcd for C35H24N1+ ([M+H]+) 458.1903, found 458.1904.

3.2.6. 9-(4-Methoxyphenyl)-9H-dinaphtho[2,1-c′:1′,2′-g]carbazole (5c)

The crude product was obtained by using diphenanthro[3,4-b:4′,3′-d]thiophene S,S-dioxide (1b) (42 mg, 0.10 mmol), p-anisidine (25 mg, 0.20 mmol), 1,4-dioxane (1.5 mL), KN(SiMe3)2 (0.5 M in toluene, 0.60 mL, 0.30 mmol) at 80 °C for 19 h according to the procedure for 5b. Purification by silica-gel column chromatography with (hexane/ACOEt = 5/1 as an eluent; Rf = 0.48) gave the title compound 5b as a pale-yellow solid (42 mg, 88% yield). The compound 5c can be separated into enantiomerically-pure (P)-5c and (M)-5c by HPLC equipped with a DAICEL CHIRALPAK® IA-3 column (Daicel Corporation, Tokyo, Japan)
In summary, we have demonstrated that the thiophene-fused thia[7]helicene can be converted into a series of aza[7]helicenes and [7]helicene-like compounds via aromatic metamorphosis without a transition metal catalyst. Especially, the 3,3'-dianion of 4,4'-biphenanthrene formed by desulfurative dilithiation were found to react with heteroatom electrophiles to give the corresponding ring-closing product. The dianion can also be prepared from 3,3'-biphenanthrene formed by desulfurative dilithiation were found to react with heteroatom electrophiles to give the corresponding ring-closing product. The dianion can also be

### Supplementary Materials

The following supporting information can be downloaded. Figures S1–S8: 
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1H and 
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13C NMR spectra for 2b, 4b, 5b, and 5c; Table S1: Crystallographic data for (M)-5c; 
Table S2–S10: Coordinates and absolute energy of the optimized structures for (P)-1a–1b, (P)-2a–2b, (P)-4a–4b, and (P)-5a–5c; Figures S9–S13: Molecular orbitals for (P)-1a–1b, (P)-2a–2b, (P)-4a–4b, and (P)-5a–5c; Table S11: The selected absorption of 1, 2, 4, and 5 calculated by TD–DFT method; File S1: CIF file for (M)-5c.
