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Communication

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A Shape-Persistent Polyphenylene Spoked Wheel

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Supporting Information Placeholder

ABSTRACT: A shape-persistent polyphenylene with a “spoked wheel” structure was synthesized as a subunit of an unprecedented two-dimensional polyphenylene, which we name graphenylene. The synthesis was carried out through a six-fold intramolecular Yamamoto coupling of a dodecabromo-substituted dendritic polyphenylene precursor, which had a central hexaphenylbenzene unit as a template. Characterizations by nuclear magnetic resonance spectroscopy and matrix-assisted laser ionization time-of-flight mass spectrometry provided an unambiguous structural proof for the wheel-like molecule with molecular weight of 3815.4 g/mol. Remarkably, scanning tunneling microscopy (STM) visualization clearly revealed defined spoked-wheel structure of the molecule with six internal pores.

Two-dimensional (2D) carbon-based materials have attracted enormous interest in recent decades due to their unique structures, intriguing properties, and versatile applications,¹⁻⁴ especially after the landmark advent of graphene.⁵ With the combination of sp^2 -, sp^2 -, and sp^3 -hybridized carbon atoms, a number of 2D carbon materials with different architectures have also been synthesized and/or theoretically predicted.^{2,6-18} In particular, graphene has attracted major attention during the last decade, which also stimulated the studies of other 2D carbon materials such as γ -graphyne and graphdiyne (Figure 1).^{10,11,14,19,20} Atomically precise syntheses of γ -graphyne and graphdiyne remain elusive, but widely studied phenylene-ethynylene-based macrocycles can be regarded as their molecular subunits.²¹⁻²⁵

The electronic properties of 2D carbon materials are critically dependent on the composition and arrangement of the sp^2 -, and sp^2 -, and/or sp^3 -hybridized carbon atoms. For example, while graphene is a zero-bandgap semimetal, γ -graphyne and graphdiyne are both predicted to be semiconductors with open bandgaps.¹² Moreover, theoretical studies elucidate that graphdiyne has an excellent intrinsic electron mobility reaching 10^5 cm²/Vs.¹² Further exploration of the structural diversity of 2D carbon architecture is expected to furnish materials with unique and tunable properties. Nevertheless, the number of reported examples of 2D carbon materials is still limited.^{2,6-18}

Figure 1. Structural representations of 2D carbon materials: (a) graphene, (b) γ -graphyne, (c) graphdiyne, and two newly proposed, purely sp^2 -hybridized structures (d) graphenylene-1 and (e) graphenylene-2.

By replacing ethynylene groups of γ -graphyne and graphdiyne with p -phenylene units, two purely sp^2 -hybridized 2D carbon structures with hexagonal symmetry can be conceived, which we name graphenylene-1 and graphenylene-2, respectively (Figure 1). Notably, graphenylene-1 corresponds to a single-layer substructure of the hypothetical carbon allotrope “cubic graphite”,²⁶ the synthesis of which still remains elusive. Because the direct and precise fabrication of these 2D carbon architectures is also highly challenging, we start with the synthesis of their subunits with defined sizes and shapes. During the last decade, Höger *et al.* have developed elegant modular syntheses of large and shape-persistent “molecular spoked wheels”, consisting of phenylene-ethynylene units, with the diameter reaching ~ 12 nm.^{27,28} Adapting their modular protocol, we designed synthetic routes towards two polyphenylene spoked wheels **1** and **2**, consisting of 22 and 37 benzene rings, as nanosized subunits of graphenylene-1 and

graphenylene-2, respectively (Scheme 1). Although the preparation of **1** turned out to be elusive by the current method due to the high steric hindrance in the final ring-forming step, **2** could be obtained and unambiguously characterized by NMR, mass spectrometry, and scanning tunneling microscopy (STM). Compound **2** has a D_{6h} -symmetrical structure and is expected to be highly shape-persistent with its rigid polyphenylene backbone.

Towards the synthesis of **1** and **2**, dendritic polyphenylene precursors **8a** and **b**, respectively, are designed with twelve bromo groups at appropriate positions for forming the rims of the wheel-like molecules by six-fold aryl-aryl coupling reactions. The hexaphenylbenzene core is expected to serve as a shape-persistent template that facilitates the ring closure. Precursors **8a** and **b** can be synthesized through the cobalt-catalyzed cyclotrimerization of corresponding alkynes **7a** and **b**, respectively. On the one hand, **8a** with C_3 symmetry will be inevitably mixed with its undesired, constitutional isomer **8a'** (Scheme 1). We therefore introduce strongly polar benzoate groups at the peripheral phenyl rings of precursor **8a** to enable its isolation by column chromatography. On the other hand, **8b** with C_6 symmetry can be obtained as a single isomer and does not require polar groups for the isolation.

Dendritic polyphenylene precursor **8a** was prepared over four steps as shown in Scheme 1, and could be easily separated from its undesired regioisomer **8a'** through silica gel column chromatography. Matrix-assisted laser ionization time-of-flight (MALDI-TOF) mass spectrometry (MS) and ^1H NMR with H-H 2D COSY and nuclear Overhauser enhancement spectroscopy (NOESY) validated the chemical identity of **8a** (see the Supporting Information). The Yamamoto coupling of precursor **8a** was then attempted under various conditions, including the use of a microwave reactor (see Table S1), but failed to give the desired compound **1**, which is most probably due to steric hindrance of the *ortho*-bromo-biphenyl unit of **8a**.

In order to circumvent the steric problem in the last cyclization step, compound **2** was next targeted, replacing the phenylene bridges of **1** with biphenylene units. The synthesis of **2** was performed as shown in Scheme 1, following a similar strategy as the one towards **1**. Suzuki coupling of freshly prepared 9-dodecyl-9-borabicyclo(3.3.1)nonane with 4-bromobenzaldehyde gave 4-dodecylbenzaldehyde (**3b**), which was then subjected to Lewis acid-catalyzed condensation with 4-bromoacetophenone to yield triphenylpyrilium salt **4b**. Condensation of **4b** with sodium 4-iodophenacetate in refluxing acetic anhydride provided oligophenylene **5b**, and then Suzuki coupling between **5b** and 1,2-bis(4-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)phenyl)acetylene furnished alkyne **7b**. The dendritic precursor **8b** was obtained by the cobalt-catalyzed cyclotrimerization of **7b** in refluxing toluene (Scheme 1). MALDI-TOF MS and ^1H NMR analyses proved the chemical identity of precursor **8b** (Figure S28, and S29).

In contrast to the unsuccessful synthesis of **1**, the intramolecular ring closure of precursor **8b** to compound **2** could be efficiently achieved by nickel-catalyzed coupling under microwave treatment. This was most probably due to the reduced steric hindrance and the absence of the *o*-bromobiphenyl moiety in precursor **8b**, in comparison to the more congested precursor **8a**. Preparative gel permeation chromatography (GPC) allowed for the complete purification, providing **2** in 56% yield as a colorless solid. The MALDI-TOF MS spectrum of **2** indicated the presence of a single species with $m/z = 3815.5$, consistent with the desired molecular weight of 3815.4 g/mol, and the experimental isotopic distribution was in perfect agreement with the simulated pattern (Figure 2).

Scheme 1. Synthetic route towards compounds **1** and **2**.^a

^aReagents and conditions: (i) 4-bromoacetophenone, BF₃·Et₂O, 100 °C, 12 h; **4a**: 25%; **4b**: 46%; (ii) sodium 4-iodophenacetate Ac₂O, 150 °C, 12 h; **5a**: 47%; **5b**: 39%; (iii) **7a**: **6**, Pd(PPh₃)₂Cl₂, Et₃N, THF, 12 h, rt, 71%; **7b**: 1,2-bis(4-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)phenyl)acetylene, Pd(PPh₃)₄, K₂CO₃, THF, methanol/H₂O, 12 h, 60 °C, 58%; (iv) Co₂(CO)₈, toluene, refluxing; **8a**: 23%; **8b**: 34%; (v) Ni(COD)₂, COD, bipyridine, THF, 120 °C, microwave reactor; **1**: 0%; **2**: 58%. THF: tetrahydrofuran. COD: 1,5-cyclooctadiene.

Figure 2. MALDI-TOF MS spectrum of **2**. Inset: isotopic distribution (black) compared with mass spectrum simulated (red) for C₂₉₄H₂₈₂.

^1H NMR together with 2D COSY and NOESY spectra provided further structural proof for compound **2**, where every aromatic

proton signal could be unambiguously assigned (Figures 3, S9, and S10). The ^1H NMR spectrum displayed that only the signals from H_e , H_g , and H_h were slightly up-field shifted from 7.48, 7.23, and 7.20 ppm to 7.42, 7.15, and 7.14 ppm, respectively, when the temperature was raised from 298 to 366 K. This is presumably because these protons are pointing out of the plane at room temperature due to the steric repulsion, but have higher freedom of rotation at an elevated temperature, leading to a reduced deshielding effect from the adjacent phenylene ring. Furthermore, the diameter of **2** in tetrachloroethane could be estimated to be 3.89 nm based on the NMR diffusion-ordered spectroscopy (DOSY) measurement (Figure S11) with a calculated self-diffusion constant (D) of $7.06 \times 10^{-11} \text{ m}^2 \text{ s}^{-1}$, which was comparable with the theoretical molecular size (3.74 nm) of the polyphenylene core of **2** calculated by the density functional theory (DFT) at the level of B3LYP/6-31G(d) (Figure S4).

The theoretically calculated geometry of **2** (DFT, B3LYP/6-31G(d)) clearly showed that its polyphenylene backbone is confined into a two-dimensional plane, in contrast to three-dimensional polyphenylene dendrimers. Six ‘rim-like’ biphenylene units are supported by six ‘spoke-like’ biphenylene segments and they are all tilted out of the plane (Figures S3 and S4). The UV-Vis absorption spectrum of **2** displays a slight blue-shift of $\sim 10 \text{ nm}$ compared to that of precursor **7b**, which is presumably due to a decrease in the effective conjugation length of *p*-oligophenylene units by the tilting (Figure S2a). Indeed, the absorption spectrum of **2** is similar to those of cyclic nonaphenylene derivatives reported by Iyoda et al.,²⁹ suggesting the interruption of the π -conjugation. On the other hand, the fluorescence spectra display a red shift and a larger Stokes shift for **2** ($\lambda_{\text{fl}} = 422 \text{ nm}$, with a shift of 125 nm) compared to those of **8b** ($\lambda_{\text{fl}} = 393 \text{ nm}$, with a shift of 83 nm) (Figure S2c), which suggests that the excited state of **2** has a more planarized structure than that of **8b**, with extended π -conjugation in contrast to their ground-state structures.

To confirm the shape-persistent nature of the wheel-like molecule, a self-assembled monolayer of **2** was characterized at sub-molecular resolution using STM at the 1-phenyloctane/highly oriented pyrolytic graphite (HOPG) interface at room temperature (Figure 4). Wheel-like molecule **2** formed a hexagonal 2D pattern on the substrate with unit cell parameters of $a = b = 4.7 \pm 0.2 \text{ nm}$ and $\alpha = 60 \pm 3^\circ$. Figure 4a shows a high-resolution STM image of the monolayer where sub-molecular features within the aromatic part of the molecular spoked wheel are clearly visible. The conjugated polyphenylene cores appear as bright spoked wheels separated by darker regions wherein the peripheral dodecyl chains are adsorbed.

A molecular model (Figure 4b) built using the experimentally obtained lattice parameters reveals that the dodecyl chains possibly interact with each other via van der Waals interactions. The model also shows that the region in between the dodecyl chains is largely empty, and thus it is possibly filled by mobile solvent molecules. The dimension of the polyphenylene core obtained from calibrated STM images (as indicated by the dashed line in Figure 4d: $\sim 2.4 \text{ nm}$) matches closely with the one obtained from the theoretically calculated structure (2.2 nm). Furthermore, the STM contrast of the aromatic core matches closely with the anticipated structure of **2**. Each bright feature consists of a hexagonal frame (‘rim-like’ biphenylenes) filled with a star-shaped feature (‘spoke-like’ biphenylenes) with a central dark blob. The ‘spoke-like’ biphenylenes give rise to six adjacent pores as expected from the molecular structure (Figure 4c and 4d). The tilted biphenylenes have presumably enhanced the self-assembly by multiple “CH- π ” interactions with the graphite surface. The peripheral phenyl rings that form a part of the phenyl dodecyl substitution are also visualized (arrows in Figure 4a and 4b). Thus, the STM analysis not only proved the chemical struc-

ture of the targeted compound **2**, but also provided direct evidence for its shape-persistent conformation on a solid substrate, which was not accessible via conventional characterization methods.

Figure 3. ^1H NMR spectrum of **2** in $\text{C}_2\text{D}_2\text{Cl}_4$ at room temperature with the assignment of aromatic protons.

Figure 4. (a) High-resolution STM image of a self-assembled monolayer of **2**. The unit cell parameters are $a = b = 4.7 \pm 0.2 \text{ nm}$, and $\alpha = 60 \pm 3.0^\circ$ (one molecule per unit cell). Tunneling current, $I_{\text{set}} = 25 \text{ pA}$; Sample bias, $V_{\text{bias}} = -1.5 \text{ V}$. (b) Molecular model of the self-assembled network of **2**. (c) Digital zoom (4.1 nm $\times$ 4.1 nm) of the STM image provided in panel (a) showing the aromatic core of a single molecule of **2**. (d) Same image as in (c) but overlapped with the molecular structure of **2**.

In summary, we considered two 2D carbon architectures, graphenylene-1 and graphenylene-2, with hexagonal symmetry by replacing ethynyl groups of γ -graphyne and graphdiyne, respectively, with *p*-phenylene linkages, and synthesized a shape-persistent polyphenylene spoked wheel **2**, corresponding to a subunit of graphenylene-2. With the assistance of the hexaphenylbenzene core as the cyclization template, the macrocyclic backbone of **2** was constructed via six-fold intramolecular nickel-catalyzed coupling with high yield. Remarkably, a submolecular-resolution STM image could be obtained at the liquid/solid interface, clearly elucidating the shape-persistent “spoked wheel” structure of **2**. Such molecular structure with six internal pores (approximately 4 Å in diameter as measured both in models and STM images) can be highly interesting for studying supramolecular interactions with other guest molecules or ions, since they are expected to accommodate e.g. methyl groups or Br^- and I^- anions. These results open up a new avenue for designing and synthesizing a further variety of unprecedented 2D carbon structures and their subunits.

ASSOCIATED CONTENT

Supporting Information

Experimental details, cyclic voltammograms, NMR, UV-Vis absorption, fluorescence, and MALDI-TOF MS spectra, and additional STM images and molecular models. “This material is available free of charge via the Internet at <http://pubs.acs.org>.”

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Notes

The authors declare no competing financial interest.

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REFERENCES

- (1) Geim, A. K.; Novoselov, K. S. *Nature Mater.* **2007**, *6*, 183.
- (2) Segawa, Y.; Ito, H.; Itami, K. *Nat. Rev. Mater.* **2016**, *1*, 15002.
- (3) Pinzón, J. R.; Villalta-Cerdas, A.; Echegoyen, L. In *Unimolecular and Supramolecular Electronics I: Chemistry and Physics Meet at Metal-Molecule Interfaces*; Metzger, M. R., Ed.; Springer Berlin Heidelberg: Berlin, Heidelberg, 2012; p 127.
- (4) Jariwala, D.; Sangwan, V. K.; Lauhon, L. J.; Marks, T. J.; Hersam, M. C. *Chem. Soc. Rev.* **2013**, *42*, 2824.
- (5) Novoselov, K. S.; Geim, A. K.; Morozov, S. V.; Jiang, D.; Zhang, Y.; Dubonos, S. V.; Grigorieva, I. V.; Firsov, A. A. *Science* **2004**, *306*, 666.
- (6) Falcao, E. H. L.; Wudl, F. *J. Chem. Technol. Biotechnol.* **2007**, *82*, 524.
- (7) Georgakilas, V.; Perman, J. A.; Tucek, J.; Zboril, R. *Chem. Rev.* **2015**, *115*, 4744.
- (8) Zhang, S.; Zhou, J.; Wang, Q.; Chen, X.; Kawazoe, Y.; Jena, P. *Proceedings of the National Academy of Sciences* **2015**, *112*, 2372.
- (9) Sharma, B. R.; Manjanath, A.; Singh, A. K. *Sci. Rep.* **2014**, *4*, 7164.
- (10) Li, Y.; Xu, L.; Liu, H.; Li, Y. *Chem. Soc. Rev.* **2014**, *43*, 2572.
- (11) Hirsch, A. *Nature Mater.* **2010**, *9*, 868.
- (12) Diederich, F.; Rubin, Y. *Angew. Chem. Int. Ed. Engl.* **1992**, *31*, 1101.
- (13) Diederich, F. *Nature* **1994**, *369*, 199.
- (14) Baughman, R. H.; Eckhardt, H.; Kertesz, M. *J. Chem. Phys.* **1987**, *87*, 6687.
- (15) Bieri, M.; Treier, M.; Cai, J.; Ait-Mansour, K.; Ruffieux, P.; Groning, O.; Groning, P.; Kastler, M.; Rieger, R.; Feng, X.; Müllen, K.; Fasel, R. *Chem. Commun.* **2009**, 6919.
- (16) Eichhorn, J.; Nieckarz, D.; Ochs, O.; Samanta, D.; Schmitt, M.; Szabelski, P. J.; Lackinger, M. *ACS Nano* **2014**, *8*, 7880.
- (17) Klappenberger, F.; Zhang, Y.-Q.; Björk, J.; Klyatskaya, S.; Ruben, M.; Barth, J. V. *Acc. Chem. Res.* **2015**, *48*, 2140.
- (18) Li, Q.; Yang, B.; Lin, H.; Aghdassi, N.; Miao, K.; Zhang, J.; Zhang, H.; Li, Y.; Duhm, S.; Fan, J.; Chi, L. *J Am Chem Soc* **2016**, *138*, 2809.
- (19) Malko, D.; Neiss, C.; Viñes, F.; Görling, A. *Phys. Rev. Lett.* **2012**, *108*, 086804.
- (20) Zhou, J.; Gao, X.; Liu, R.; Xie, Z.; Yang, J.; Zhang, S.; Zhang, G.; Liu, H.; Li, Y.; Zhang, J.; Liu, Z. *J Am Chem Soc* **2015**, *137*, 7596.
- (21) Bhaskar, A.; Guda, R.; Haley, M. M.; Goodson *J Am Chem Soc* **2006**, *128*, 13972.
- (22) Haley, M. M.; Brand, S. C.; Pak, J. J. *Angew. Chem. Int. Ed. Engl.* **1997**, *36*, 836.
- (23) Iyoda, M.; Yamakawa, J.; Rahman, M. J. *Angew. Chem. Int. Ed.* **2011**, *50*, 10522.
- (24) H. F. Bunz, U.; Rubin, Y.; Tobe, Y. *Chem. Soc. Rev.* **1999**, *28*, 107.
- (25) Tahara, K.; Yoshimura, T.; Sonoda, M.; Tobe, Y.; Williams, R. V. *J. Org. Chem.* **2007**, *72*, 1437.
- (26) Baughman, R. H.; Cui, C. *Synth. Met.* **1993**, *55*, 315.
- (27) May, R.; Jester, S.-S.; Höger, S. *J Am Chem Soc* **2014**, *136*, 16732.
- (28) Mössinger, D.; Hornung, J.; Lei, S.; De Feyter, S.; Höger, S. *Angew. Chem. Int. Ed.* **2007**, *46*, 6802.
- (29) Rahman, M. J.; Yamakawa, J.; Matsumoto, A.; Enozawa, H.; Nishinaga, T.; Kamada, K.; Iyoda, M. *J. Org. Chem.* **2008**, *73*, 5542.

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Shape-persistent macrocycle of sp^2 -hybridized carbons