Porphyrinoids Hot Paper

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Fusing Porphyrins and Phospholes: Synthesis and Analysis of a Phosphorus-Containing Porphyrin

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Abstract: A phosphole-fused porphyrin dimer, as a representative of a new class of porphyrins with a phosphorus atom, was synthesized for the first time. The porphyrin dimer exhibits remarkably broadened absorption, indicating effective π -conjugation over the two porphyrins through the phosphole moiety. The porphyrin dimer possesses excellent electronaccepting character, which is comparable to that of a representative electron-accepting material, [60]PCBM. These results provide access to a new class of phosphorus-containing porphyrins with unique optoelectronic properties.

Porphyrins are macrocyclic 18π -conjugated molecules that have been actively studied in various fields owing to their potential for diverse applications, for example, in materials science and as catalysts, as well as owing to their vital roles in natural systems.^[1] As their attractive properties are attributed to the 18n electron conjugated system, their structural and electronic properties can be tuned by metalation, peripheral modification, and replacement of a pyrrole ring with another heterocycle.^[2,3] In particular, the chemistry of porphyrins with phosphorus atoms has been actively investigated in the light of the ability of the phosphorus atom to form coordination and covalent bonds (Figure 1). For instance, phosphoruscontaining porphyrin complexes can be used as a structural motif to create oligomeric architectures. The wheel-and-axle phosphorus(V) porphyrin arrays 1 were synthesized by Shimidzu and co-workers.^[4] Furthermore, phosphorus insertion has recently been shown to be an effective approach for creating new porphyrinoids.^[5] Latos-Grażyński and co-workers reported the phosphorus complex of N-fused porphyrin 2, a 20π porphyrin, as an isophlorin analogue.^[5a] As another example, peripherally phosphanylated porphyrins have also been synthesized, which can form peripherally metalated porphyrins.^[6] The PCP pincer complexes of bis(phosphanyl)porphyrin 3 were catalytically active depending on the central metal atom in the porphyrin core.^[6c]

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Figure 1. Examples of porphyrins with phosphorus atoms.

Phosphaporphyrins 4 have been presented by Matano and co-workers as core-modified porphyrins containing a phosphole ring.^[7] They corroborated that replacement of a pyrrole ring with a phosphole has an impact on the structure, aromaticity, and optical and electrochemical properties as well as on the coordination abilities. However, to the best of our knowledge, no porphyrins with a peripherally fused phosphole structure such as 5 have been reported in spite of extensive efforts to explore various phosphole-fused π-conjugated molecules.^[8] We envisioned that such phospholefused porphyrins would exhibit unique optical and electrochemical properties owing to the phosphole moiety. Towards this goal, we exploited a new synthetic method to create bis(pyrrolo)heteroles.^[9] On the basis of the reactivity of bis(pyrrolo)phosphole as a key building block, we expected that the two porphyrin rings would be integrated through condensation with the bis(pyrrolo)phosphole. Herein, we report for the first time the synthesis of phosphole-fused porphyrin dimers as a new class of porphyrins with a phosphorus atom.

The phosphole-fused porphyrin dimer 12 was synthesized by the route shown in Scheme 1. The treatment of 4,4'dibromo-3,3'-bipyrrole $6^{[9]}$ with *t*-BuLi and PhPCl₂ followed by addition of elemental sulfur afforded the triisopropylsilylprotected bis(pyrrolo)phosphole 7 in 58% yield. Whereas the P^{III} derivative is unstable under ambient atmosphere,^[9] the P= S compound 7 is sufficiently stable to be used as the precursor. Then, 7 was deprotected with tetrabutylammonium fluoride (TBAF) to give bis(pyrrolo)phosphole 8. The reaction of 5,10di(4-methylphenyl)tripyrrane 9 with Vilsmeier reagent provided diformyltripyrrane 10 in 52% yield. Tripyrrane dicarbinol 11 was obtained by the reduction of 10 with NaBH₄. Then, we attempted the acid-catalyzed condensation of 8 and 11. However, we detected only a trace amount of 12 under standard reaction conditions for porphyrin synthesis. After extensive screening, we finally succeeded in isolating the

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Scheme 1. Synthesis of phosphole-fused porphyrin dimer **12**. DCE = 1,2-dichloroethane, MSA = methanesulfonic acid, TBAF = tetrabutylammonium fluoride, TIPS = triisopropylsilyl.

desired porphyrin dimer **12** in 0.7% yield upon condensation of **8** and **11** with 0.4 equiv of methanesulfonic acid (MSA) at 60 °C in 1,2-dichloroethane for 1 h followed by oxidation with 6 equiv of 2,3-dichloro-5,6-dicyanobenzoquinone (DDQ) at room temperature for 3 h. While the yield was low, we performed the synthesis repeatedly and obtained a sufficient amount of **12** for all measurements (ca. 10 mg). The highresolution electrospray ionization time-of-flight (HR-ESI-TOF) mass spectrum of **12** exhibited a positive ion peak at m/z1117.3936 (calcd for C₇₄H₅₄N₈PS: 1117.3924 [M+H]⁺), which is consistent with the molecular structure of **12**. The ¹H NMR spectrum of **12** in CD₂Cl₂ is shown in Figure 2. The



Figure 2. ¹H NMR spectrum (aromatic region) of porphyrin dimer **12** in CD_2Cl_2 at 25 °C.

six resonances arising from the β -protons are observed in the range of $\delta = 8.90-10.03$ ppm, and the two resonances arising from the meso protons are observed at $\delta = 10.51$ and 12.07 ppm. It should be noted that the resonance of the meso-H^a protons is remarkably downfield-shifted because of the aromatic ring current of the neighboring porphyrin macrocycle. The ³¹P NMR spectrum displays a resonance from the P=S moiety at $\delta = 18.76$ ppm (see the Supporting Information, Figure S4).

To our delight, single crystals suitable for X-ray diffraction analysis were obtained by vapor diffusion of 2-propanol into a solution of **12** in 1,2-dichloroethane.^[10] The crystal structure of **12** revealed two almost co-planar porphyrin macrocycles, which suggests effective π -conjugation over the two porphyrin moieties (Figure 3). Whereas porphyrin ring B adopts



Figure 3. X-ray crystal structure of **12**: a) top view and b) side view. Thermal ellipsoids set at 50% probability. Solvent molecules and meso-aryl substituents omitted for clarity.

a highly planar conformation, a slight distortion was observed for porphyrin ring A. The mean-plane deviations (MPDs) of the porphyrin macrocycles A and B (24 core atoms) were calculated to be 0.234 and 0.035 Å, respectively. In the packing structure, **12** forms a face-to-face dimeric structure, in which the intermolecular distance is approximately 3.5 Å (Figure S5). Notably, this dimeric structure forms a slipped π - π -stacked structure (Figure S6). The distance between the two porphyrin substructures is also about 3.5 Å. This π - π interaction probably induces the distortion of porphyrin ring A in the packing structure. The dimeric structure and effective π - π interactions would be favorable for potential applications of **12** as a charge-transport material (see below).^[11]

The UV/Vis absorption spectra of 12 and 5,10-di(4methylphenyl)porphyrin 13 in CH₂Cl₂ are shown in Figure 4. In contrast to porphyrin monomer 13, porphyrin dimer 12 displays remarkably broadened absorption whereas the absorption coefficients of **12** at $\lambda \approx 400$ nm are smaller than those of 13. The split Soret bands are broadened in the range of $\lambda = 400-500$ nm. The lowest-energy Q band of 12 $(\lambda = 668 \text{ nm})$ is red-shifted by $\Delta \lambda = 43 \text{ nm}$ relative to that of 13 $(\lambda = 625 \text{ nm})$,^[12] and the absorption edge reaches to 700 nm. It is noteworthy that this red shift of the Q band for 12 is significantly larger than that for a usual β - β -linked porphyrin dimer ($\Delta \lambda = 8 \text{ nm}$).^[13] Thus these absorption features support an effective interaction between the two porphyrin macrocycles through the phosphole moiety. The steady-state fluorescence spectrum of 12 was recorded in CH₂Cl₂, and an emission peak was detected at $\lambda = 683$ nm (Figure 4). The Stokes shift of $12 (330 \text{ cm}^{-1})$ is rather comparable to that of 13

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Figure 4. UV/Vis absorption (solid lines) and emission (dashed lines) spectra of **12** (black) and **13** (gray) in CH_2Cl_2 . The samples were excited at $\lambda = 400$ nm.

 (230 cm^{-1}) , which is consistent with the highly rigid planar structure of **12**.

The electrochemical properties of 7, 12, and 13 were studied by cyclic voltammetry (CV) and differential pulse voltammetry (DPV) in CH2Cl2 vs. ferrocene/ferrocenium cation (Fc/Fc⁺) with tetrabutylammonium hexafluorophosphate (Bu_4NPF_6) as the electrolyte (Figure S7). Bis-(pyrrolo)phosphole 7 gave rise to an oxidation peak at 0.73 V whereas no reduction peak was visible at potentials up to -2.5 V. The electrochemical properties of 7 are similar to those of other bis(pyrrolo)heteroles.^[9] Porphyrin monomer 13 displays two reversible reduction peaks at -1.65 and -2.05 V and two quasi-reversible oxidation peaks at 0.50 and 0.91 V. On the other hand, 12 underwent three quasi-reversible reductions at -1.19, -1.44, and -1.90 V and two irreversible oxidations at 0.53 and 0.75 V. The first and second oxidation potentials of 12 stem mainly from the two porphyrin moieties and the phosphole conjugated to the porphyrins, respectively (see below). The effective π -expansion over the two porphyrins lowers the oxidation potentials while they are shifted in the positive direction by the electron-withdrawing effect of the phosphorus(V) center, offsetting the effects; therefore, the oxidation potentials of 12 are comparable to those of 13 and 7. In contrast, the three reduction peaks, including the unexpected new one, are significantly shifted in the positive direction, probably owing to the strongly electron-accepting character arising from the phosphole moiety (see below). The three one-electron reduction processes of 12 reflect distinct electronic communication over the two porphyrin moieties through the phosphole moiety. The large difference ($\Delta E_{\rm red}$ = 0.25 V) between the first and second reduction potentials also indicates effective electronic communication. It is worth noting that the first reduction potential of 12 is more positive than those of phosphole derivatives with high electron affinity.^[14] Given that the energy level of Fc/Fc⁺ is -4.8 eV under vacuum, the LUMO level of 12 was estimated to be $-3.61 \mbox{ eV}.^{[15]}$ This value is close to that of [60]PCBM (ca. -3.7 eV),^[16] a representative fullerene derivative that is used as an electron-accepting material for charge transport as well as an organic photovoltaic material.^[17] Therefore, this high electron-accepting ability, combined with effective π - π stacking in the crystal structure, suggests that **12** has potential as an n-type semiconductor.

To gain further insight into the electronic properties of 12, we performed DFT calculations at the B3LYP/6-31G(d,p) level of theory. First, we found that 12a is the most stable NH tautomer of 12 (Figure 5a). However, the difference in the



Figure 5. a) Three NH tautomers of 12 and their relative energies. b) Selected Kohn–Sham orbitals of 12a at the B3LYP/6-31G(d,p) level of theory.

relative energies of the tautomers 12a-12c is quite small (ca. 3 kJmol^{-1}), which agrees with the single NH peak in the ¹H NMR spectrum of **12** at room temperature (Figure S4). The Kohn-Sham frontier orbitals of 12a are illustrated in Figures 5b and S8. For the eight Kohn-Sham orbitals composed of four porphyrin orbitals, the orbital distributions are well delocalized over the whole π -conjugated system. HOMO and HOMO-1 of 12a originate from the HOMO of 13. As the phosphole ring is located on the node of HOMO/ HOMO-1, these orbitals mainly exhibit porphyrin character. However, the energy levels of HOMO and HOMO-1 of 12a are slightly shifted in the negative direction owing to the electron-withdrawing effect of the phosphorus(V) atom. On the other hand, the LUMO of 12a is well delocalized over the two porphyrin moieties as well as the phosphole ring (Figure 5b). Importantly, the LUMO of **12a** is derived from the LUMO +1 of 13 and remarkably stabilized compared to the LUMO + 1/LUMO + 2, which stem from the LUMO of 13 (Figure S8). The new reduction peak observed in the cyclic voltammogram was rationalized by the stabilized LUMO (Figure S7). As the LUMO displays a large orbital distribution on the phosphole ring, the LUMO possesses phosphole character to a large extent.^[8,14] This is a consequence of the stabilization solely by the fusion of the two porphyrins and the phosphole ring.

We also calculated nucleus-independent chemical shift (NICS) values for the optimized structure of **12a** (Figure S9). The NICS value at the center of the porphyrin moiety is -13.8 ppm, whereas the NICS value at the center of the phosphole ring is +8.33 ppm. This large positive NICS value results from the double diatropic ring current effect of the two porphyrin macrocycles. Although the phosphole ring can participate in π -conjugation with the two porphyrin moieties, it has little impact on the 18π aromatic character of the porphyrin moiety. Furthermore, we carried out time-dependent.

dent DFT (TD-DFT) calculations to evaluate the absorption (Figure S10 and Table S2). The oscillator strengths of the HOMO/LUMO and HOMO-1/LUMO transitions are small (f < 0.03) whereas the HOMO-2/LUMO transition has a large oscillator strength (f = 0.41). These results suggest that the absorption at $\lambda = 600-700$ nm is not caused by intramolecular charge transfer (ICT) from the porphyrin core to the phosphole ring, but mainly by $\pi - \pi^*$ transitions of the porphyrin dimer with phosphole moiety.

Finally, we assessed the charge-transport ability of 12 and 13 by flash photolysis time resolved microwave conductivity (FP-TRMC) measurements.^[18] The FP-TRMC profiles for drop-cast films of 12 and 13, prepared from toluene solution, upon irradiation at $\lambda = 355$ nm are displayed in Figure S11, with a charge recombination rate constant high enough compared to the time constant of the measurement system (ca. 100 ns), especially for 12. The maximum transient conductivity $(\phi \Sigma \mu)_{max}$ for 13 was determined to be 6.5 × $10^{-6} \text{ cm}^2 \text{V}^{-1} \text{s}^{-1}$, where ϕ and $\Sigma \mu$ are the charge carrier generation efficiency and the sum of the charge carrier mobilities. Importantly, the $(\phi \Sigma \mu)_{max}$ value for 12 $(5.5 \times$ $10^{-5} \text{ cm}^2 \text{V}^{-1} \text{s}^{-1}$) is one order of magnitude larger than that for 13. This remarkable difference is most likely due to the effective π - π interaction in the solid state, supporting the potential utility of the phosphole-fused structure as a chargetransport material.

In summary, we have reported the synthesis of phospholefused porphyrin dimer 12 as a representative of a new class of phosphorus-containing porphyrins, exemplifying the potential utility of our bis(pyrrolo)heteroles as building blocks for heterole-fused porphyrin dimers. Porphyrin dimer 12 exhibits effective π -conjugation over the two porphyrin macrocycles through the phosphole moiety. The low reduction potential of 12 is solely due to the integration of the two porphyrin moieties into one large π -system through the phosphole moiety. FP-TRMC measurements suggest the prospective use of 12 as a novel charge-transport material. The optical and electrochemical properties of the phosphole-fused porphyrin dimer could be tuned by further modifications of the phosphorus atom, metalation of the porphyrin core, or peripheral functionalizations. Furthermore, we believe that this report will enable the synthesis of new classes of porphyrins containing not only phosphorus but also other main-group elements, such as sulfur and silicon.

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Communications



Communications

Porphyrinoids

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T. Higashino,* T. Yamada, T. Sakurai, S. Seki, H. Imahori* _____ **IIII**-**IIII**

Fusing Porphyrins and Phospholes: Synthesis and Analysis of a Phosphorus-Containing Porphyrin



A phosphole-fused porphyrin dimer as a representative of a new class of phosphorus-containing porphyrins was synthesized. This structure exhibits remarkably broadened absorption as well as unique optoelectronic properties and is a good electron acceptor owing to the unique phosphole-fused structure.

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