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5,20-Bis(α -oligothienyl)-substituted [26]hexaphyrins possessing electronic circuits strongly perturbed by *meso*-oligothienyl substituents†

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A series of [26]hexaphyrins(1.1.1.1.1.1) bearing two α -oligothienyl substituents at 5,20-positions have been synthesised and are shown to have a dumbbell hexaphyrin conformation, to which the α -oligothienyl groups are linked with small dihedral angles to form an acyclic helix-like conjugated network. While their distinct diatropic ring currents and four reversible reduction waves characteristic of aromatic [26]hexaphyrins indicate that the [26]hexaphyrin aromatic circuits are viable, the absorption spectra and excited state dynamics are significantly perturbed, which becomes increasingly evident with elongation of the oligothenyl substituents. DFT calculations of these hexaphyrins indicated that the LUMO and LUMO + 1 are localised on the hexaphyrin circuit and the HOMO and HOMO – 1 are spread over the acyclic helix-like conjugation network, which can explain the perturbed absorption spectra.

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Introduction

In the last two decades, the chemistry of expanded porphyrins has gained increasing popularity, in light of their flexible structures, multi-metal coordination properties, anion sensing abilities, facile interconversions between multiple and neutral redox states, and unprecedented chemical reactivities such as drastic skeletal rearrangements.¹ Fairly flexible electronic systems of *meso*-aryl expanded porphyrins have been demonstrated by the exploration of versatile electronic states such as Hückel aromatic, Hückel antiaromatic,² Möbius aromatic,³ Möbius antiaromatic,⁴ stable monoradical,⁵ and singlet biradicaloid species.⁶ As an additional example, internally 5,20-aromatic-bridged [26]hexaphyrins have been recently developed as dual electronic systems consisting of [18]porphyrin and/or [26]hexaphyrin, which can be modulated by the internal bridging aromatic units (Fig. 1a).⁷ In this paper, we report the synthesis of a series of 5,20-bis(α -oligothienyl) [26]hexaphyrins(1.1.1.1.1.1) (T2–T4), which display characteristic optical and electronic properties due to the presence of 5,20-bis(α -oligothienyl) substituents (Fig. 1b).

Results and discussion

Synthesis of 5,20-bis(α -oligothienyl) [26]hexaphyrins(1.1.1.1.1.1)

[26]Hexaphyrins T2–T4 have been prepared by a modified method used for the synthesis of 5,20-(2-thienyl) [26]hexaphyrin(1.1.1.1.1.1) T1.⁸ Acid-catalysed condensation of 5,10-

(a) Previous Work

(b) This Work

Fig. 1 [26]Hexaphyrins possessing dual electronic systems; (a) internally aromatic-bridged [26]hexaphyrins (X = S, NH) and (b) bis(α -oligothienyl) [26]hexaphyrins. Effective conjugated networks are indicated by the coloured lines.

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bis(pentafluorophenyl)tripyrane **1** with corresponding α -oligothiophene-5-carbaldehydes followed by oxidation with 2,3-dichloro-5,6-dicyano-1,4-benzoquinone (DDQ) gave **T2–T4** in 22, 10, and 15% yields, respectively (Scheme 1). High-resolution electrospray ionisation time-of-flight (HR ESI-TOF) mass measurements showed the parent ion peaks at m/z 1457.0685 ($[M+H]^+$; calcd for $C_{70}H_{25}N_6F_{20}S_4$: 1457.0699) for **T2**, m/z 1621.0462 ($[M+H]^+$; calcd for $C_{78}H_{29}N_6F_{20}S_6$: 1621.0453) for **T3**, and m/z 1783.0059 ($[M-H]^-$; calcd for $C_{86}H_{31}N_6F_{20}S_8$: 1783.0062) for **T4** (see ESI†). 5,15-Bis(α -oligothienyl)-substituted porphyrins **P1–P4** were also prepared (ESI†) to examine the effects of α -oligothienyl substituents on the electronic system of porphyrins, which highlight the characteristically flexible electronic properties of [26]hexaphyrins.

The solid-state structures of **T2–T4** have been unambiguously determined by single crystal X-ray diffraction analysis.⁹ Hexaphyrins **T2–T4** all show a dumbbell hexaphyrin conformation, to which the two oligothiophenyl groups are appended with small dihedral angles, being favorable for conjugation with the hexaphyrin macrocycle (Fig. 2 and ESI†). The hexaphyrin frames of **T2–T4** are essentially the same as those of 5,20-unsubstituted hexaphyrin (**T0**)^{5a} and **T1**⁸ with regard to the planar dumbbell structure. The dumbbell conformation is intrinsically the most stable for [26]hexaphyrins, because of four possible energy-stabilising intramolecular hydrogen bonding interactions, but is only allowed for [26]hexaphyrins bearing small substituents at 5,20-positions.^{5a,8} Thus, **T0** is the most stable, having a strain-free fairly planar conformation with the largest diatropic ring current. The observed dumbbell structures of **T1–T4** have been similarly ascribed to small 2-thienyl and α -oligothienyl substituents.⁸ The two oligothiophenyl substituents are positioned above and below the macrocycle and are oriented toward the same side of the hexaphyrin to form a helix-like conjugated network involving the tripyrrodimethene segments of the hexaphyrin.

The 1H NMR spectra of **T0** and **T1** indicate sharp signals at room temperature, which are consistent with the dumbbell

Fig. 2 X-Ray crystal structures of (a and b) **T2**, (c and d) **T3**, and (e and f) **T4**. Solvent molecules are omitted for clarity. The thermal ellipsoids are scaled to a 30% probability level. One of the two molecules in the unit cell is shown for **T2** and **T4**.

structures.^{5a,8} The 1H NMR spectrum of **T1** exhibits only three signals at 7.38, 7.72, and 8.16 ppm due to the outer β -pyrrolic protons, indicating a very rapid rotation of the *meso*-thienyl substituents. In contrast, the 1H NMR spectra of **T2–T4** taken at room temperature are interesting, in that the protons of the hexaphyrin periphery appear as very broad signals but the protons of the oligothiophenyl substituents appear as sharp signals. At $-60^\circ C$, these 1H NMR spectra became sharp, featuring six signals due to the outer β -pyrrolic protons in the range $\delta = 8.21$ – 5.76 ppm and signals due to the *meso*-oligothienyl groups in the range $\delta = 7.25$ – 4.06 ppm. These data indicate substantial diatropic ring currents for the hexaphyrin in **T2–T4** in accordance with the solid-state dumbbell structures. Importantly, two singlets due to the two inner NH protons appeared differently around $\delta = 8.4$ – 8.2 and 5.4 – 5.1 ppm, implying that the right and left tripyrrodimethene segments of the hexaphyrin are electronically different, owing to the influence of the *meso*-oligothienyl substituents. Most probably the rotational dynamics of the *meso*-oligothienyl substituents are slow at room temperature as compared with the 1H NMR time scale.

Scheme 1 Synthesis of 5,20-(α -oligothienyl)-substituted [26]hexaphyrins **T1–T4** and structures of **P1–P4**.

Redox potentials and UV/Vis/NIR absorption spectra

The electrochemical properties of **T1–T4** were examined by cyclic voltammetry (CV) in CH_2Cl_2 containing 0.1 M Bu_4NPF_6 as a supporting electrolyte *versus* ferrocene/ferrocenium cation (see Fig. S20 and Table S1†). All the cyclic voltammograms of **T1–T4** showed four reversible reduction waves almost at the same potential, that correspond to stepwise reductions reaching their 30π states, hence indicating that the 26π hexaphyrin electronic circuits are all viable. On the other hand, the first oxidation potentials appeared at 0.38 V for **T1**, 0.27 V for **T2**, 0.22 V for **T3**, and 0.17 V for **T4**, indicating the rising of the HOMO levels upon elongation of the *meso*-oligothienyl chains.

The absorption spectrum of **T0** displays a sharp and intense Soret-like band at 549 nm and weak but distinct Q-bands in the range 700–1100 nm as characteristic features of typical aromatic porphyrinoids. The absorption spectrum of **T1** exhibits an ill-defined Soret-like band at 616 nm and broad Q-bands in the range 700–1150 nm, which is significantly different from that of **T0**, indicating that the 2-thienyl substituents cause significant perturbation on the electronic state of the [26]hexaphyrin. This trend is increasingly evident upon elongation of the oligothieryl chains. Namely, a Soret-like band, a characteristic attribute of aromatic [26]hexaphyrins, almost disappears in the absorption spectra of **T2–T4**, and instead a broad high-energy band is observed around 350 nm for **T1**, which is red-shifted and intensified with the elongation of the *meso*-oligothienyl chains.¹⁰ In addition, the spectra of **T2–T4** show broad Q-bands in the NIR region. Collectively, these absorption features of **T2–T4** are radically different from those of aromatic porphyrinoids, but are rather similar to those of reported acyclic oligopyrromethenes.¹¹ The absorption spectra of **T1–T4** became sharpened upon lowering the temperature (ESI†), indicating the importance of the dynamic motion of the oligothieryl substituents for broadening the absorption spectra. In sharp contrast, no such drastic changes were observed in the absorption spectra of the porphyrin counterparts **P1–P4** (Fig. 3b), underlining the unique and flexible electronic properties of the [26]hexaphyrins.

Theoretical calculations and excited state dynamics

To understand the anomalous absorption spectra, density functional theory (DFT) calculations of **T0–T4** were carried out using the Gaussian 09 program at the B3LYP/6-31G(d) level.¹² Geometry optimisations produced dumbbell structures that were respectively similar to those obtained *via* X-ray diffraction analyses. Since the calculated MO diagrams of **T2–T4** are quite similar (ESI†), we discuss here the MOs of **T3** in comparison with those of **T0** (Fig. 4). Similar to the rectangular shaped [26]hexaphyrin,¹³ the HOMO and LUMO of **T0** are, respectively, similar to the a_{1u} and a_{2u} HOMOs and two degenerate e_g^* LUMOs of porphyrins with a HOMO–LUMO gap of 1.88 eV. The LUMO and LUMO + 1 of **T3** are nearly localised on the hexaphyrin circuit, and their energy levels are only slightly destabilised from those of **T0**. In contrast, the HOMO – 1 is spread over the helix-like network and the HOMO is mainly spread over the same network, and both are largely destabilised, which causes a

Fig. 3 UV/Vis/NIR absorption spectra of (a) **T0–T4** and (b) **P1–P4** in CH_2Cl_2 .

small HOMO–LUMO gap of 1.340 eV. The MO features of **T2** and **T4** are nearly the same as those of **T3**, which are all consistent with the CV results. These MO features compelled us to consider a dual contribution of the 26π -hexaphyrin network and an acyclic conjugation along the helix-like structure (Fig. 1b), which plays an important role in the electronic excitation to the excited states.

The nucleus-independent chemical shift (NICS) values¹⁴ inside the [26]hexaphyrin macrocycle were calculated to be –15.54, –12.21, –8.50, –8.30, and –8.26 for **T0–T4**, respectively. The harmonic oscillator model of aromaticity (HOMA) values¹⁵ for the [26]hexaphyrin circuit were calculated from the real crystal structures and the optimised structures to be 0.783 and 0.737 for **T0**, 0.568 and 0.604 for **T1**, 0.464 (0.425) and 0.518 for **T2**, and 0.397 and 0.498 for **T3**. The HOMA value of **T4** was only calculated for the optimised structure to be 0.498 due to insufficient crystal data to discuss the bond length. These results indicate the increasing perturbation by the 5,20-oligothienyl substituents of **T1–T4**.

We have also investigated the excited state dynamics of **T1–T4** by using femtosecond transient absorption (fs-TA) spectroscopy (Fig. S46 and 47†) to confirm the electronic perturbation by the 5,20-oligothienyl substituents. S1-state lifetimes have been determined to be 35.4 ps for **T1**, 15.2 ps for **T2**, 10.0 ps for **T3**, and 8.5 ps for **T4**, respectively, which show a dramatic decrease compared to the value of 138 ps for **T0**.^{5a} These overall

Fig. 4 Kohn-Sham MO diagrams of T0 and T3 calculated with the Gaussian 09 package. All energy levels were calculated at the B3LYP/6-31G(d) level.

observations also confirmed the increasing electronic perturbation by the oligothieryl chains.

Collectively, these experimental and theoretical calculations indicate that the perturbation provided by the oligothieryl substituents is delicate, being not strong enough to spoil the aromatic [26]hexaphyrin network but large enough to alter the absorption properties and mitigate the aromatic characters of the [26]hexaphyrin segments as a rare case.

Conclusions

5,20-Bis(α -oligothienyl)-substituted [26]hexaphyrins(1.1.1.1.1.1) **T1-T4** were prepared and were shown to possess characteristic optical and electronic properties. The observed diatropic ring currents and reversible reduction waves indicate that the aromatic [26]hexaphyrin networks are viable in **T1-T4**. Nevertheless, the absorption spectra and the excited state dynamics of **T1-T4** are significantly perturbed by the appended oligothieryl substituents. DFT calculations indicated that the LUMO and LUMO + 1 are localised at the [26]hexaphyrin network but that the HOMO and HOMO – 1 are significantly spread over the helix-like network and are largely destabilised. Further explorations of expanded porphyrins possessing dual electronic networks are actively under way in our group.

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Notes and references

- (a) A. Jasat and D. Dolphin, *Chem. Rev.*, 1997, **97**, 2267; (b) H. Furuta, H. Maed and A. Osuka, *Chem. Commun.*, 2002, 1795; (c) J. L. Sessler and D. Seidel, *Angew. Chem., Int. Ed.*, 2003, **42**, 5134; (d) T. K. Chandrashekar and S. Venkatraman, *Acc. Chem. Res.*, 2003, **36**, 676; (e) M. Stępień, N. Sprutta and L. Latos-Grażyński, *Angew. Chem., Int. Ed.*, 2011, **50**, 4288; (f) S. Saito and A. Osuka, *Angew. Chem., Int. Ed.*, 2011, **50**, 4342; (g) A. Osuka and S. Saito, *Chem. Commun.*, 2011, **47**, 4330.
- (a) S. Mori and A. Osuka, *J. Am. Chem. Soc.*, 2005, **127**, 8030; (b) S. Mori, K. S. Kim, Z. S. Yoon, S. B. Noh, D. Kim and A. Osuka, *J. Am. Chem. Soc.*, 2007, **129**, 11344; (c) K. Naoda, H. Mori and A. Osuka, *Chem.-Asian J.*, 2013, **8**, 1395; (d) H. Mori, Y. M. Sung, B. S. Lee, D. Kim and A. Osuka, *Angew. Chem., Int. Ed.*, 2012, **51**, 12459.
- (a) M. Stępień, L. Latos-Grażyński, N. Sprutta, P. Chwalisz and L. Sztrenberg, *Angew. Chem., Int. Ed.*, 2007, **46**, 7869; (b) Y. Tanaka, S. Saito, S. Mori, N. Aratani, H. Shinokubo, N. Shibata, Y. Higuchi, Z. S. Yoon, K. S. Kim, S. B. Noh, J. K. Park, D. Kim and A. Osuka, *Angew. Chem., Int. Ed.*, 2008, **47**, 681; (c) J. Sankar, S. Mori, S. Saito, H. Rath, M. Suzuki, Y. Inokuma, H. Shinokubo, K. S. Kim, Z. S. Yoon, J.-Y. Shin, J. M. Lim, Y. Matsuzaki, O. Matsushita, A. Muranaka, N. Kobayashi, D. Kim and A. Osuka, *J. Am. Chem. Soc.*, 2008, **130**, 13568; (d) Z. S. Yoon, A. Osuka and D. Kim, *Nat. Chem.*, 2009, **1**, 113.
- (a) E. Pacholska-Dudziak, J. Skonieczny, M. Pawlicki, L. Sztrenberg, Z. Ciunik and L. Latos-Grażyński, *J. Am. Chem. Soc.*, 2008, **130**, 6182; (b) T. Higashino, J. M. Lim, T. Miura, S. Saito, J.-Y. Shin, D. Kim and A. Osuka, *Angew.*

- Chem., Int. Ed.*, 2010, **49**, 4950; (c) T. Higashino, B. S. Lee, J. M. Lim, D. Kim and A. Osuka, *Angew. Chem., Int. Ed.*, 2012, **51**, 13105.
- 5 (a) T. Koide, G. Kashiwazaki, M. Suzuki, K. Furukawa, M.-C. Yoon, S. Cho, D. Kim and A. Osuka, *Angew. Chem., Int. Ed.*, 2008, **47**, 9661; (b) H. Rath, S. Tokuji, N. Aratani, K. Furukawa, J. M. Lim, D. Kim, H. Shinokubo and A. Osuka, *Angew. Chem., Int. Ed.*, 2010, **49**, 1489.
- 6 T. Koide, K. Furukawa, H. Shinokubo, J.-Y. Shin, K. S. Kim, D. Kim and A. Osuka, *J. Am. Chem. Soc.*, 2010, **132**, 7246.
- 7 (a) H. Mori, J. M. Lim, D. Kim and A. Osuka, *Angew. Chem., Int. Ed.*, 2013, **52**, 12997; (b) G. Karthik, M. Sneha, V. P. Raja, J. M. Lim, D. Kim, A. Srinivasan and T. K. Chandrashekar, *Chem.-Eur. J.*, 2013, **19**, 1886.
- 8 M. Suzuki and A. Osuka, *Chem.-Eur. J.*, 2007, **13**, 196.
- 9 (a) Crystallographic data for **T2**: $2(\text{C}_{70}\text{H}_{24}\text{F}_{20}\text{N}_6\text{S}_4) \cdot 1.82(\text{C}_6\text{H}_{14}) \cdot 1.10(\text{CH}_2\text{Cl}_2)$, $M_r = 3165.36$, triclinic, space group $P\bar{1}$ (no. 2), $a = 18.9073(16)$, $b = 20.7452(6)$, $c = 21.091(2)$ Å, $\alpha = 67.38(2)$, $\beta = 77.21(2)$, $\gamma = 65.257(13)^\circ$, $V = 6917.2(9)$ Å³, $T = 93$ K, $\rho_{\text{calcd}} = 1.520$ g cm⁻³, $Z = 2$, 87 882 reflections measured, 23 494 unique ($R_{\text{int}} = 0.0351$), $R_1 = 0.0639$ ($I > 2\sigma(I)$), $R_w = 0.1805$ (all data), GOF = 1.042; (b) Crystallographic data for **T3**: $\text{C}_{78}\text{H}_{28}\text{F}_{20}\text{N}_6\text{S}_6 \cdot 0.51(\text{C}_7\text{H}_{16}) \cdot \text{C}_7 \cdot 1.89(\text{CH}_2\text{Cl}_2)$, $M_r = 1882.68$, monoclinic, space group $C2/c$ (no. 15), $a = 49.998(16)$, $b = 12.402(3)$, $c = 35.242(11)$ Å, $\beta = 131.613(5)^\circ$, $V = 16\,338(8)$ Å³, $T = 93$ K, $\rho_{\text{calcd}} = 1.513$ g cm⁻³, $Z = 8$, 51 126 reflections measured, 14 608 unique ($R_{\text{int}} = 0.0547$), $R_1 = 0.0850$ ($I > 2\sigma(I)$), $R_w = 0.2299$ (all data), GOF = 1.034; (c) Crystallographic data for **T4**: $2(\text{C}_{86}\text{H}_{32}\text{F}_{20}\text{N}_6\text{S}_8) \cdot 2(\text{C}_7\text{H}_{16}) \cdot 0.44(\text{C}_7) \cdot 1.53(\text{CHCl}_3) \cdot \text{O}$, $M_r = 4006.31$, triclinic, space group $P\bar{1}$ (no. 2), $a = 17.2734(8)$, $b = 18.9028(5)$, $c = 30.1289(17)$ Å, $\alpha = 105.000(14)$, $\beta = 92.92(2)$, $\gamma = 107.23(3)^\circ$, $V = 8988.7(7)$ Å³, $T = 93$ K, $\rho_{\text{calcd}} = 1.481$ g cm⁻³, $Z = 2$, 99 137 reflections measured, 27 782 unique ($R_{\text{int}} = 0.0733$), $R_1 = 0.1104$ ($I > 2\sigma(I)$), $R_w = 0.3750$ (all data), GOF = 1.073.
- 10 The intense bands of **T1–T4** in the UV region are red-shifted in a manner similar to those of the α -oligothiophenes as below; 353/354 nm for **T1** and terthiophene, 385/392 nm for **T2** and quaterthiophene, 414/417 nm for **T3** and quinquethiophene, and 432/436 nm for **T4** and sexithiophene. R. S. Becker, J. Seixas de Melo, L. Macanita and F. Elisei, *J. Phys. Chem.*, 1996, **100**, 18683. One of the reviewers suggested charge transfer interactions between the [26]hexaphyrin networks and the oligothieryl substituents, but the solvent polarity effects were only marginal for **T1–T4** (ESI†).
- 11 (a) P. Morosini, M. Scherer, S. Meyer, V. Lynch and J. L. Sessler, *J. Org. Chem.*, 1997, **62**, 8848; (b) J.-Y. Shin, S. S. Hepperle, B. O. Patrick and D. Dolphin, *Chem. Commun.*, 2009, 2323; (c) S. Saito, K. Furukawa and A. Osuka, *J. Am. Chem. Soc.*, 2010, **132**, 2128.
- 12 For the full Gaussian citation, see ESI†.
- 13 T. K. Ahn, J. H. Kwon, D. Y. Kim, D. W. Cho, D. H. Jeong, S. K. Kim, M. Suzuki, S. Shimizu, A. Osuka and D. Kim, *J. Am. Chem. Soc.*, 2005, **127**, 12856.
- 14 Z. Chen, C. S. Wannere, C. Corminboeuf, R. Puchta and P. v. R. Schleyer, *Chem. Rev.*, 2005, **105**, 3842.
- 15 T. M. Krygowski and K. M. Cryański, *Chem. Rev.*, 2001, **101**, 1385.