

Iron and Copper Complexes of Tetraphenyl-*m*-benzporphyrin: Reactivity of the Internal C–H Bond

Chen-Hsiung Hung,^{*†} Fei-Chien Chang,[†] Cheng-Yu Lin,[†] Krystyna Rachlewicz,[‡] Marcin Stępień,[‡] Lechosław Latos-Grażyński,^{*‡} Gene-Hsiang Lee,[§] and Shie-Ming Peng^{||}

Department of Chemistry, National Changhua University of Education, Changhua 50058, Taiwan, Department of Chemistry, University of Wrocław, 14 F. Joliot-Curie St., Wrocław 50 383, Poland, Instrumentation Center, National Taiwan University, Taipei, Taiwan, and Department of Chemistry, National Taiwan University, Taipei, Taiwan

Received February 12, 2004

Iron and copper complexes of tetraphenyl-*m*-benzporphyrin (TP-*m*BPH)H have been prepared and structurally characterized. The iron system, (TP*m*BPH)Fe^{II}Br, contains a high-spin Fe(II) center. In the solid state the complex forms dimeric units linked by weak CH...Br hydrogen bonds. The Cu complex contains a tetrameric copper cluster with a Cu₂Cl₄²⁻ unit bridging two [(TP*m*BPC)Cu^{II}]⁺ fragments. The formation of (TP*m*BPC)H represents an example of copper-catalyzed chlorination on the internal carbon atom of (TP*m*BPH)H.

The idea of replacing a pyrrole ring in the porphyrin macrocycle with a carbocycle (heterocarbocycle) to exploit the coordinating properties of porphyrin cores has initiated studies on carbaporphyrins, a new class of porphyrin analogues.^{1–4} Recently, it was shown that N-confused tetraphenylporphyrin (CTPPH)₂ can stabilize unusual metal oxidation states⁵ and provides a unique environment for studying the formation and reactivity of organometallic metal–carbon bonds.^{6–8} It has also been found that the internal C–H bond of the (CTPPH)₂ system is susceptible to substitution reactions. The products of alkylation,⁹ nitra-

tion,¹⁰ internal fusion,¹¹ and cyanation¹² have been reported to date. More recently, tetraaryl-*m*-benzporphyrin and its para isomer, which have a pyrrole ring replaced with a benzene ring, have been prepared and structurally characterized.^{13,14} Both benzporphyrins have already been shown to coordinate selected metal(II) ions.^{13–15} In the reactions of *m*-tetraphenylbenzporphyrin with silver salts, the pyridination¹⁶ and acetoxylation products¹⁴ have been isolated. Reactions of 22-acetoxy-*m*-benzporphyrin with zinc(II) or cadmium(II) give metal complexes containing an unmodified acetoxy substituent while the reactions with nickel(II) or palladium(II) cleave the Ac–O bond and form 22-hydroxy-*m*-benzporphyrin species.¹⁷ The unusual reactivity of the internal carbon has inspired us to further exploit the chemistry of carbaporphyrins. Herein we report the syntheses and characterization of iron and copper complexes of tetraphenyl-*m*-benzporphyrin (Scheme 1). Interestingly, the reaction of copper(II) chloride with *m*-benzporphyrin yields a 22-chloro-*m*-benzporphyrin complex from a chlorination reaction of the internal carbon on the *m*-benzporphyrin core.

When refluxing a THF solution of free base (TP*m*BPH)H and FeBr₂ for 4 h in the presence of a few drops of 2,6-lutidine under anaerobic conditions, (TP*m*BPH)Fe^{II}Br (**2**) can be worked up in 71% yield.¹⁸ The final product gives a Soret band at 431 nm and *Q* bands at 770 and 860 nm (Figure 1). Compared with the free base of (TP*m*BPH)H the Soret band is red-shifted only slightly but the bathochromic shift of the *Q* bands is significant. The compound is relatively stable toward dioxygen in contrast to the instantaneous reaction with O₂ in the case of (HCTPPH)Fe^{II}Br.¹⁹

* Authors to whom correspondence should be addressed. E-mail: chhung@cc.ncue.edu.tw (C.-H.H.); llg@wchuwr.chem.uni.wroc.pl (L.L.-G.).

† National Changhua University of Education.

‡ University of Wrocław.

§ Instrumentation Center, National Taiwan University.

|| Department of Chemistry, National Taiwan University.

- (1) Furuta, H.; Asano, T.; Ogawa, T. *J. Am. Chem. Soc.* **1994**, *116*, 767.
- (2) Chmielewski, P. J.; Latos-Grażyński, L.; Rachlewicz, K.; Głowiak, T. *Angew. Chem., Int. Ed. Engl.* **1994**, *33*, 779.
- (3) Berlin, K.; Breitmaier, E. *Angew. Chem., Int. Ed. Engl.* **1994**, *33*, 1246.
- (4) Lash, T. D.; Chaney, S. T. *Angew. Chem., Int. Ed. Engl.* **1997**, *36*, 839.
- (5) Furuta, H.; Ogawa, T.; Uwatoko, Y.; Araki, K. *Inorg. Chem.* **1999**, *38*, 2676.
- (6) Chen, W.-C.; Hung, C.-H. *Inorg. Chem.* **2001**, *40*, 5070.
- (7) Bohle, D. S.; Chen, W.-C.; Hung, C.-H. *Inorg. Chem.* **2002**, *41*, 3334.
- (8) Hung, C.-H.; Chen, W.-C.; Lee, G.-H.; Peng, S.-M. *Chem. Commun.* **2002**, 1516.
- (9) Chmielewski, P. J.; Latos-Grażyński, L.; Głowiak, T. *J. Am. Chem. Soc.* **1996**, *118*, 5690.

- (10) Ishikawa, Y.; Yoshida, I.; Akaiwa, K.; Koguchi, E.; Sasaki, T.; Furuta, H. *Chem. Lett.* **1997**, 453.
- (11) Furuta, H.; Ishizuka, T.; Osuka, A.; Ogawa, T. *J. Am. Chem. Soc.* **1999**, *121*, 2945.
- (12) Xiao, Z.; Patrick, B. O.; Dolphin, D. *Chem. Commun.* **2003**, 1062.
- (13) Stępień, M.; Latos-Grażyński, L. *J. Am. Chem. Soc.* **2002**, *124*, 3838.
- (14) Stępień, M.; Latos-Grażyński, L. *Chem. Eur. J.* **2001**, *7*, 5113.
- (15) Stępień, M.; Latos-Grażyński, L.; Sztrenberg, L.; Panek, J.; Latajka, Z. *J. Am. Chem. Soc.* **2004**, *126*, 4566.
- (16) Stępień, M.; Latos-Grażyński, L. *Org. Lett.* **2003**, *5*, 3379.
- (17) Stępień, M.; Latos-Grażyński, L. *Inorg. Chem.* **2003**, *42*, 6183.

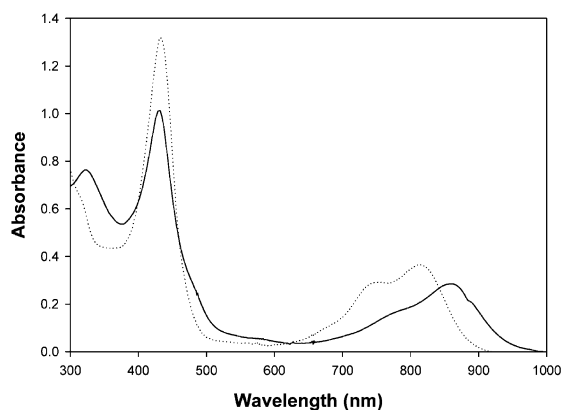
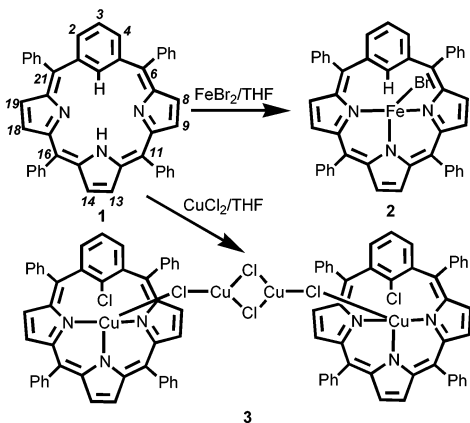


Figure 1. The absorption spectra of **2** (—) and **3** (···).

Scheme 1



A magnetic susceptibility measurement performed in solution for **2** using the Evans method gives an effective magnetic moment of $4.65 \mu_B$. The value is in the range for a high-spin Fe(II) center.

As shown in Figure 2, the β -pyrrolic protons of **2** are located at 49.4, 21.1, and 6.5 ppm at 293 K as identified by ^2H NMR for (TPmBPH- d_6)Fe^{II}Br. These values are close to those recently reported for high-spin nickel(II) complexes of 22-acetoxyporphyrin¹⁷ and tetraphenyl-*m*-benzporphyrin.¹⁵ The hydrogens on the *m*-phenylene ring have been identified by default at 22-H, 440; 2,4-H, 65.8; 3-H, -33.2 ppm (298 K). Interestingly, the 440 ppm shift of the inner C–H proton is much smaller than the corresponding shift observed for (HCTPPH)Fe^{II}Br (812 ppm, 298 K)²⁰ but similar to the value for (TPmBPH)Ni^{II}Cl (386 ppm, 298 K).¹⁵ An *agostic mechanism* of spin density transfer, described

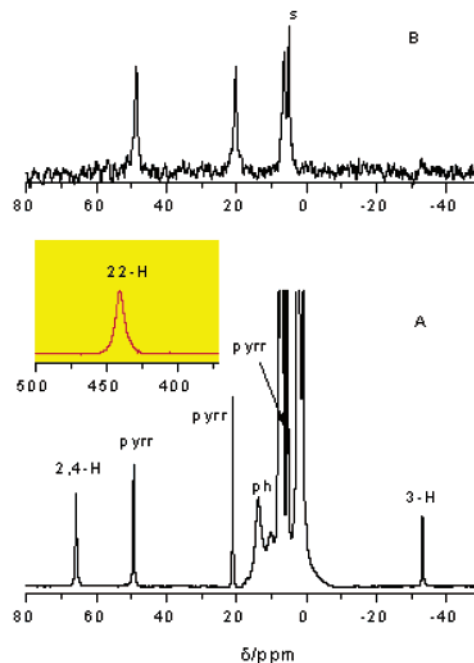
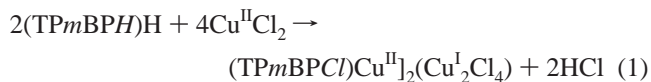


Figure 2. NMR spectra of iron(II) tetraphenyl-*m*-benzporphyrin: (A) (TPmBPH)Fe^{II}Br (^1H NMR, dichloromethane- d_2 , 298 K); (B) (TPmBPH- d_6)Fe^{II}Br (^2H NMR, dichloromethane, 298 K). Inset in trace A presents the downfield region with the 22-H resonance. Peak labels follow the systematic numbering of the porphyrin ring or denote proton groups: pyrr, pyrrole ring; ph, *meso*-phenyl; s, solvent.

previously for (TPmBPH)Ni^{II}Cl, explains the shifts of (TPmBPH)Ni^{II}Cl as resulting from electron donation from the CH bond to the metal.¹⁵

The reaction between Cu(II) and (TPmBPH)H has been carried out under much milder conditions. Using CuCl₂ as the metal carrier under anaerobic conditions at room temperature in the absence of base, [(TPmBPH)Cu^{II}]₂(Cu^I₂Cl₄) (**3**) can be readily obtained in 75% yield.²¹ The UV–vis spectrum of **3** (Figure 1) resembles that of **2** with the Q band slightly blue-shifted. The elemental analysis suggested that the compound is a tetranuclear complex with a [Cu₂Cl₄]²⁻ unit bridging two copper(II) benzporphyrins. Importantly, the mass peak at m/z 721.2 suggests that a chlorination reaction proceeds cleanly according to eq 1. This reactivity resembles the silver-mediated 22-substitutions observed earlier for the reactions of (TPmBPH)H with acetate ions¹⁴ and pyridine.¹⁶



The structures of both **2** and **3** are confirmed by X-ray structural analysis. The iron complex crystallized in the $P\bar{1}$

(18) Absorption spectrum (CH₂Cl₂) [λ_{max} , nm (log ϵ , M⁻¹ cm⁻¹): 339 (4.06), 431 (4.34), 770 (3.53, sh), 860 (3.81)]. MS (ESI): m/z = 760.9 (calcd m/z = 760.1 for M + H⁺), 680.4 (calcd m/z = 680.2 for M⁺ – Br), 626.5 (calcd m/z = 625.3 for M + H⁺ – Br – Fe). ^1H NMR (CD₂Cl₂): 400 (1 H, 22-H), 65.8 (2 H, 2,4-H), 49.4 (2 H, pyrr), 21.1 (2 H, pyrr), 6.9 (2 H, pyrr), -33.2 (2 H, 3-H). Crystallographic data for **2**: C₅₃H₃₈BrFeN₃, T = 150(2) K, MW = 852.62, triclinic, space group $P\bar{1}$, a = 11.0498(13) Å, b = 11.5449(13) Å, c = 16.5738(19) Å, α = 78.544(2)°, β = 79.527(2)°, γ = 77.699(2)° V = 2003.0(4) Å³, Z = 2, D_c = 1.414 Mg/m⁻³, λ = 0.71073 Å, μ = 1.416 mm⁻¹, $F(000)$ = 876. The structure was solved by direct methods and refined by least squares against F^2 to R_1 = 0.0414 (wR_2 = 0.0788) and S_{GOF} = 0.957.

(19) Rachlewicz, K.; Wang, S.-L.; Ko, C.-L.; Hung, C.-H.; Latos-Grażyński, L. *J. Am. Chem. Soc.* **2004**, *126*, 4420.

(20) Rachlewicz, K.; Wang, S.-L.; Peng, C.-H.; Hung, C.-H.; Latos-Grażyński, L. *Inorg. Chem.* **2003**, *42*, 7348.

(21) Absorption spectrum (CH₂Cl₂) [λ_{max} , nm (log ϵ , M⁻¹ cm⁻¹): 433 (5.12), 757 (4.46, sh), 814 (4.56)]. MS (ESI): m/z = 721.2 (calcd m/z = 721.1 for [Cu(TPmBPH)]⁺). Anal. Calcd for Cu₄Cl₆N₆C₉₂H₅₈·CH₂Cl₂·C₆H₁₄: N, 4.45; C, 63.06; H, 3.95. Found: N, 4.78; C, 64.92; H, 3.73. Crystallographic data for **3**: C₄₈H₃₃Cl₇Cu₂N₃, T = 150(1) K, MW = 1027.00, monoclinic, space group $P2_1/c$, a = 14.5784(5) Å, b = 25.0959(8) Å, c = 12.6346(5) Å, β = 108.462(1)°, V = 4384.6(3) Å³, Z = 4, D_c = 1.556 Mg/m⁻³, λ = 0.71073 Å, μ = 1.556 mm⁻¹, $F(000)$ = 2074. The structure was solved by direct methods and refined by least squares against F^2 to R_1 = 0.0588 (wR_2 = 0.1329) and S_{GOF} = 1.091.

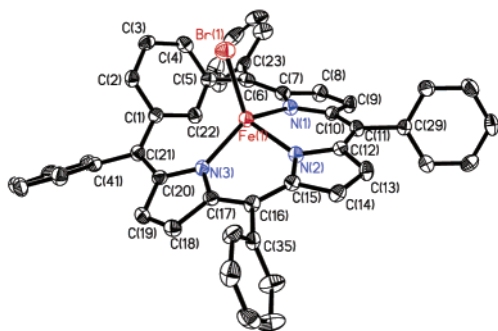


Figure 3. The crystal structure of **2** with 50% thermal ellipsoids.

space group with one solvate toluene per asymmetric unit (Figure 3). The iron center binds to three pyrrolic nitrogens and a bromide in a distorted tetrahedral structure. The bromide bends toward the *m*-phenylene ring and expands the angle of N2–Fe1–Br1 to 116.15(9)°. The distance of 2.073(3) Å for Fe1–N2 is slightly shorter than 2.131(3) and 2.141(3) Å for Fe1–N1 and Fe1–N3. The distances between iron and pyrrolic nitrogens are in the range for high-spin Fe(II) porphyrin complexes.²² The tripyrrolic fragment arranges in a dome-shaped geometry. The average deviation of three pyrrole rings and four *meso*-carbons from the three nitrogen plane is 0.35 Å. The iron sits 0.694(2) Å above the three nitrogen plane. The large deviation of iron from the porphyrin plane has been observed in five-coordinated iron(II) porphyrin complexes.²² As expected, the *m*-phenylene ring is tilted away from the mean porphyrin plane. The distances from iron to the inner carbon (C22) and inner hydrogen (H22) are 2.579(4) and 2.439(36) Å, respectively. These distances are longer than the corresponding distances of 2.361 and 1.971 Å in (HCTPPH)Fe^{II}Br wherein we confirmed the existence of an agostic interaction between Fe and the inner C–H bond.⁸ Nevertheless, these values are small compared to the van der Waals contact²³ and similar to those seen in chloronickel(II) *m*-benzporphyrin (Ni···C22, 2.549(3) Å; Ni···H22 2.40(3) Å),¹⁵ and they evidence the formation of a weak agostic bond. Importantly, the angle between Fe1–C22 and the *m*-phenylene ring is 110.5°, which is much less than 126.2° for (HCTPPH)Fe^{II}Br.⁶ In the crystal lattice, **2** forms a cyclic dimeric structure with mutual hydrogen bonding interactions between the axial bromides and H3 atoms on the *m*-phenylene rings (see Supporting Information).

The crystal of **3** belongs to the $P2_1/c$ space group with the molecule located at a symmetry center. The tetranuclear complex (Figure 4) is a neutral compound, and four solvate dichloromethanes per tetranuclear complex are located in the unit cell.

(22) Scheidt, W. R.; Reed, C. A. *Chem. Rev.* **1981**, *81*, 543.

(23) Bondi, A. J. *J. Phys. Chem.* **1964**, *68*, 441.

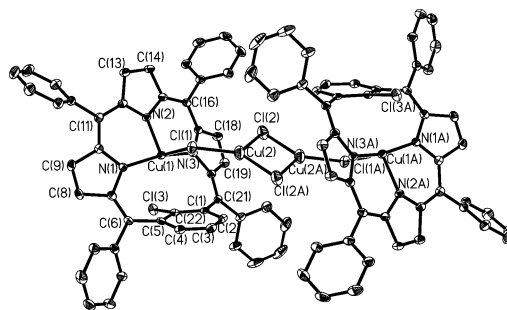


Figure 4. The crystal structure of **3** with 50% thermal ellipsoids which shows the tetranuclear complexes and TPmBPCl cores.

In agreement with the elemental analysis described above, the copper complex is a tetranuclear complex with a bridging [Cu₂Cl₄]²⁻ unit. The bridging ligand of [Cu₂Cl₄]²⁻ is rare but has been reported in several structures including the recent one by Busch et al.²⁴ The structure parameters of [Cu₂Cl₄]²⁻ in **3** are comparable to the literature values. A chlorine atom on the *m*-phenylene moiety can be clearly identified with a distance of 1.751(3) Å for C22–Cl3. The TPmBPCl moiety has a tilted *m*-phenylene ring and a saddle arrangement of the tripyrrolic unit. The average distance between copper and pyrrolic nitrogens of 1.976(3) Å agrees with the presence of a copper(II) ion in the carbaporphyrin central cavity.²⁵ The distance between copper(II) and the inner carbon C22 is 2.698(5) Å, which is significantly longer than the distance in **2**.

In conclusion, iron and copper complexes of *m*-benzporphyrin have been prepared. The studies on the NMR spectroscopy and crystal structure of **2** suggest a weaker interaction between the iron center and internal C–H for **2** compared with (HCTPPH)Fe^{II}Br. The formation of a tetranuclear copper complex and the copper-assisted chlorination is unexpected but further demonstrates the versatile chemistry that can be carried out in carbaporphyrin systems.

Acknowledgment. This work was supported by the National Science Council of Taiwan under Grant NSC 92-2113-M-018-003 (C.-H.H.) and by the State Committee for Scientific Research of Poland (KBN) under Grant 4 T09A 147 22 (L.L.-G. and M.S.). M.S. thanks the Foundation for Polish Science for financial support.

Supporting Information Available: Experimental procedures, spectroscopic data, VT plots for **2** in toluene-*d*₈, and X-ray crystallographic data for **2** and **3**. This material is available free of charge via the Internet at <http://pubs.acs.org>.

IC049821L

(24) Rybak-Akimova, E. V.; Busch, D. H.; Kahol, P. K.; Pinto, N.; Alcock, N. W.; Clase, H. J. *Inorg. Chem.* **1997**, *36*, 510.

(25) Scheidt, W. R.; Turowska-Tyrk, I. *Inorg. Chem.* **1994**, *33*, 1314.