

## Synthesis and chemical reactivity of new zinc porphyrin diazoacetates catalyzed by ruthenium and iron porphyrins

Daniel Carrie, Hassan Srour, Paul Le Maux and Gerard Simonneaux\*

*Ingénierie Chimique et Molécules pour le Vivant, UMR 6226, Campus de Beaulieu, 35042 Rennes Cedex, France*

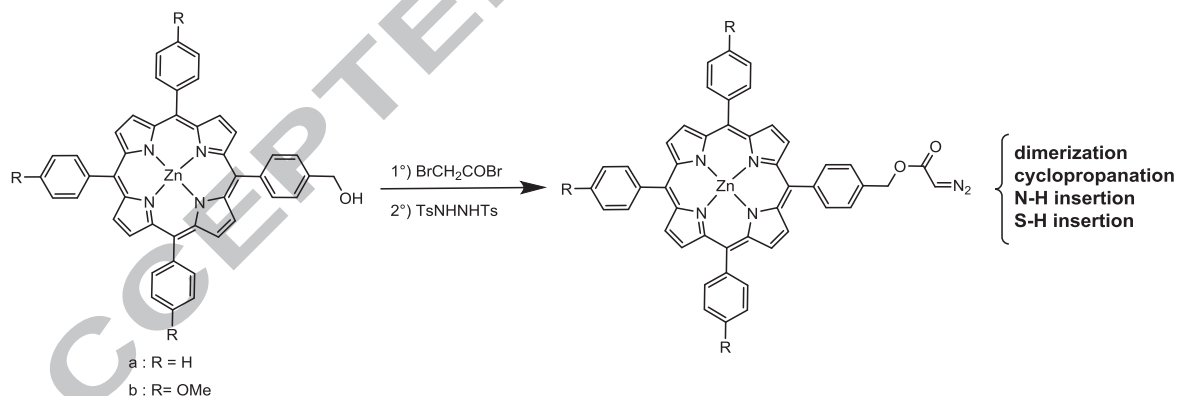
e-mail address : [gerard.simonneaux@univ-rennes1.fr](mailto:gerard.simonneaux@univ-rennes1.fr)

fax number: 33(0)2 23 23 56 37

### Abstract

Zinc porphyrin diazoacetates were synthesized from the corresponding bromoacetates by treatment with *N,N'*-ditosylhydrazine in ~70% yield. The transfer of carbenoids derived from the porphyrin diazo compounds catalyzed by metalloporphyrins (Ru,Fe) undergoes dimerization, cyclopropanation, N-H and S-H insertion reactions, respectively.

### Graphical abstract

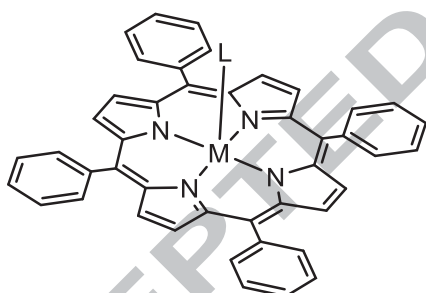


### Keywords:

Meso-phenyl-diazo-porphyrin ester, ruthenium porphyrin, iron porphyrin, catalytic carbene transfer

The applications of diazocarbonyl compounds to organic synthesis still continue to attract the attention of many chemists.<sup>1</sup> Various methodologies have been developed for

preparing different types of diazocarbonyl compounds and there are now well-established procedures.<sup>1,2</sup> Important examples of these processes include the functionalization of porphyrin macrocycles using diazo compounds which has been recently reviewed.<sup>3</sup> Thus porphyrin derivatives such as chlorins with a diazo function in the macrocycle core were first reported by Cavaleiro and coworkers in 1997.<sup>4</sup> These chlorin diazoketones can be used as versatile intermediates for the synthesis of new compounds.<sup>5</sup> In 2006, Zaleski's group reported an alternative strategy to obtain 2-diazo-3-oxochlorins.<sup>6</sup> In all these cases, the diazo group is directly inserted in the core yielding chlorin derivatives and to our knowledge, there is no diazo group directly appended to the periphery of the porphyrin ring. We previously reported the use of metalloporphyrins for catalytic insertion of diazo compounds in N-H bonds,<sup>7,8</sup> including insulin protein.<sup>9</sup> Asymmetric cyclopropanation was also developed using chiral iron and ruthenium porphyrins as catalysts.<sup>10,11</sup> In this paper, we report the first synthesis of zinc porphyrin diazoacetates and some of their reactivity using metalloporphyrins as catalysts (Figure 1).

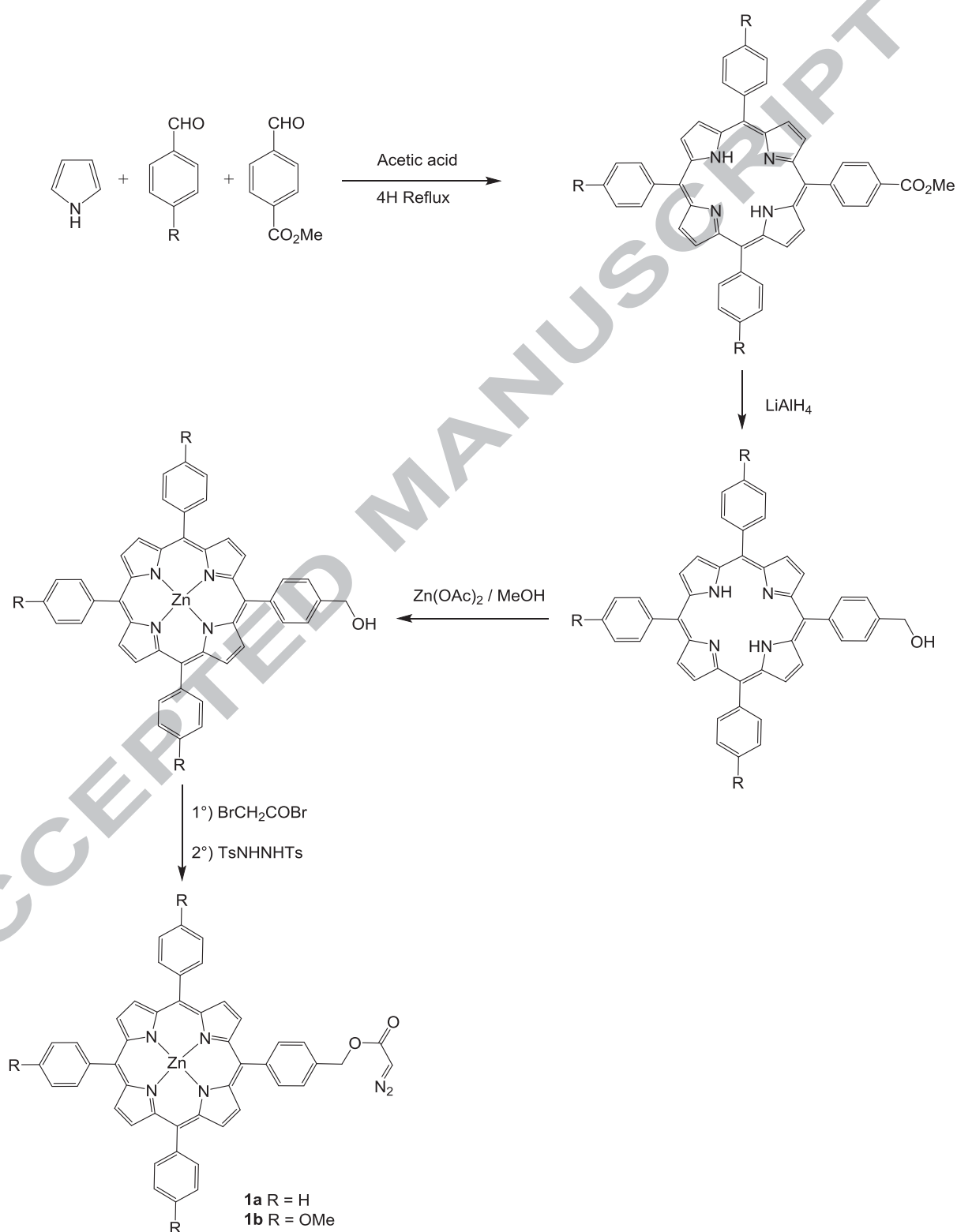


M-L RuCO: TPPRuCO  
M-L Fe-Cl: TPPFeCl

Figure 1. Structure of the catalysts.

A convenient synthetic method for the preparation of a variety of diazoacetates from the corresponding bromoacetates by treatment with  $N,N'$ -ditosylhydrazine has been recently reported.<sup>12</sup> Herein, we successfully used this recent method with hydroxymethyl-porphyrins that were prepared as previously reported by Tamiaki.<sup>13</sup> After Zn metalation, reaction with bromoacetate and then with  $N,N'$ -ditosylhydrazine yielded the expected diazo derivatives. From 5-[4-(hydroxymethyl)phenyl]-10,15,20-triphenylporphyrin, the corresponding zinc diazoporphyrin ester **1a** was prepared in 72% yield (Scheme 1). Similar results were obtained with 5-[4-(hydroxymethyl)phenyl]-10,15,20-5-[4-(hydroxymethyl)phenyl]-10,15,20-tris(4-methoxyphenyl)porphyrin giving **1b** with 67% yield. These diazo esters were first characterized by  $^1\text{H}$  and  $^{13}\text{C}$  NMR and mass spectrometry. A characteristic diazo proton at

~5.0 ppm as a broad peak was observed in the  $^1\text{H}$  NMR spectrum. It should be noted that a preliminary zinc insertion is essential for the synthesis since free base porphyrins gave only traces of diazo derivatives. The general synthetic pathway is summarized in Scheme 1.

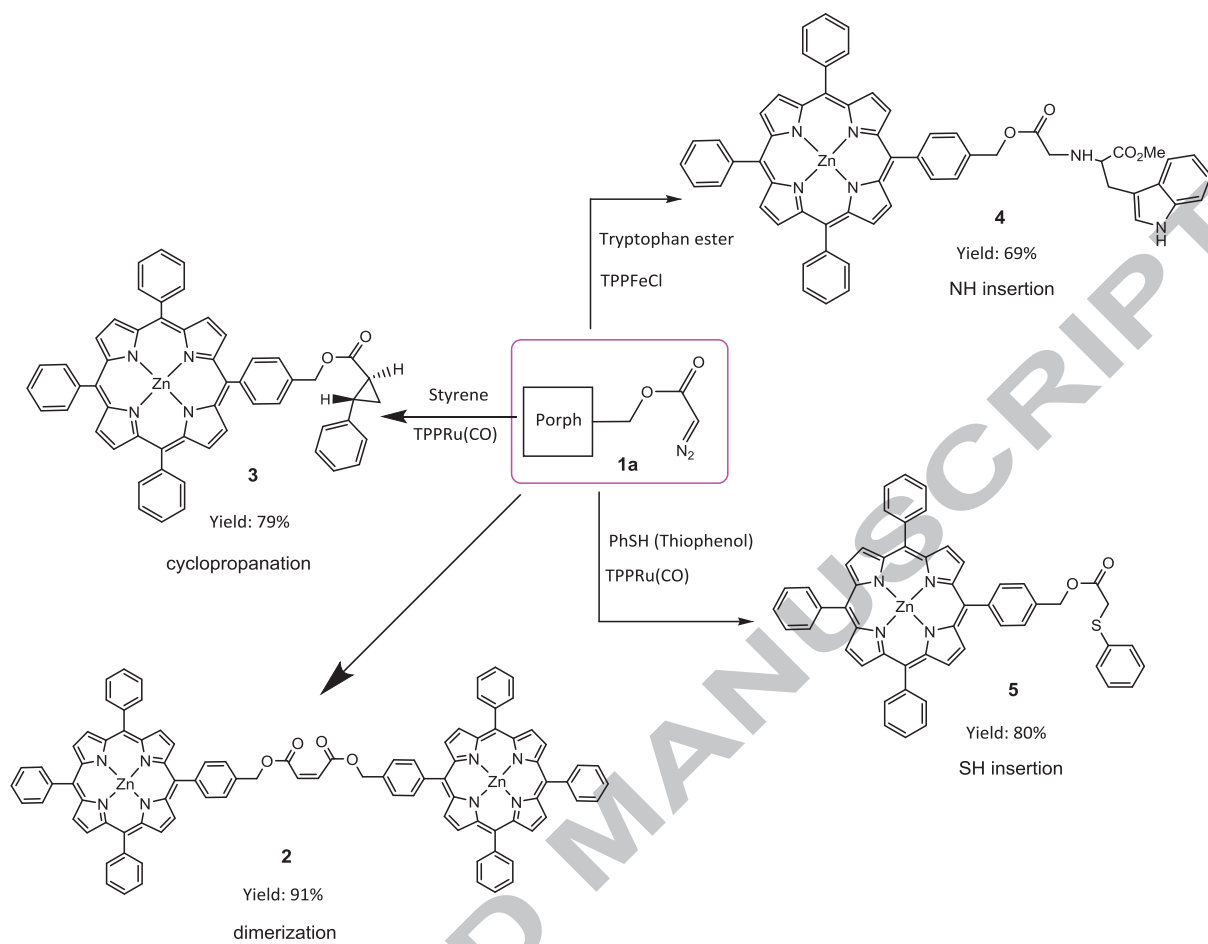


Scheme 1. General pathway for the synthesis of Zn diazoporphyrin ester derivatives.

To demonstrate the presence of possible metalcarbene intermediate, first its decomposition to form olefinic products at room temperature in the presence of tetraphenylporphyrin ruthenium carbon monoxide (Figure 1) as catalyst was examined. Starting from **1a**, this coupling reaction catalyzed by ruthenium porphyrin to form olefins, proceeds slowly, possibly through a metalcarbene intermediate, yielding the *cis* isomer **2** in overall 91% yield (Scheme 2). In our hands, the *trans* isomer was not detected.<sup>14</sup> However the reaction needs 24 hours to be completed. In comparison, a similar ruthenium(II) porphyrin species has been reported to catalyze rapidly the decomposition of ethyl diazoacetate to form diethyl maleate and diethyl fumarate with a large *cis/trans* ratio of 15.<sup>15</sup> The stereochemical constraints that result from the steric interaction of the axial porphyrin carbene atoms with atoms of the porphyrinato core<sup>16</sup> may explain the decrease of the rate. The generally accepted reaction pathway for transition metal catalyzed carbene dimerization of diazo derivatives involves initial attack of the diazo compound at the metal center to generate a metal carbene.<sup>17,18</sup> Nucleophilic attack on the metal carbene by diazo compound, followed by dissociation of the olefin from the metal complex, completes the catalytic cycle. The origin of the *cis* selectivity during the coupling of  $\alpha$ -diazo compounds has been previously interpreted on the basis of steric and electronic effects.<sup>19-21</sup>

To evaluate the reactivity of diazo porphyrin ester, its catalytic decomposition was also examined in the presence of styrene in toluene at room temperature using tetraphenylporphyrin ruthenium carbon monoxide (Figure 1) as catalyst. The cyclopropane **3** was formed with 79% yield and complete *trans* diastereoselectivity with a concomitant formation of the dimer (15%), resulting from coupling of two carbene precursors. The formation of the *trans* isomer, which was confirmed by NOESY spectroscopy, is similar to that observed when ethyl diazoacetate is used as reactant.<sup>22</sup>

The insertion of electrophilic carbenes in the N-H bonds of protected  $\alpha$ -amino esters or amides is a powerful method for N-alkylating this class of compounds.<sup>23</sup> Since we<sup>7,8</sup> and others<sup>24-27</sup> previously reported N-H insertion of diazoesters catalyzed by metalloporphyrins (Fe, Ru), N-H insertion with a diazo porphyrin ester was also investigated (Scheme 2). The results presented in Scheme 2 show that the iron complex is a good catalyst for the transformation of tryptophan methyl ester into the expected N-H inserted products with 69% yield. It should be noted that only the dimerization was observed with the ruthenium complex.



Scheme 2. Reactivity of diazoporphyrin esters.

Since peptidyl diazomethyl ketones appeared initially to be specific inactivators of cysteine proteinases,<sup>28</sup> insertion of diazoporphyrin ester into S-H bonds, catalyzed by metalloporphyrins, was assayed (Scheme 2). Treatment of thiophenol with diazoporphyrin ester catalyzed by complex **1a** gave insertion of the diazo derivative into the S-H bond with 85 % yield. The mechanisms of the cyclopropanation reaction, N-H insertion and S-H insertion, catalyzed by metalloporphyrins, have been previously discussed by us<sup>8</sup> and others<sup>25-27</sup> several times, and will not be presented herein. All the new derivatives were identified by NMR and mass spectral analysis.<sup>29</sup>

In summary, the synthesis of new zinc diazoporphyrin esters occurs with good yields offering for the first time a general access to original porphyrins. Studies on the application of the insertion reaction into N-H and S-H bonds for bioconjugation of these potentially fluorescent probes to proteins are in progress.

## References and notes

1. Ford, A.; Miel, H.; Ring, A.; Slattery, C. N.; Maguire, A. R.; McKervey, M. A. *Chem. Rev.* **2015**, *115*, 9981-10080.
2. Zhang, Z.; Wang, J. *Tetrahedron* **2008**, *64*, 6577-6605.
3. Gomes, A. T. P. C.; Neves, M. G. P. M. S.; Cavaleiro, J. A. S. *J. Porphyrins Phthalocyanines* **2011**, *15*, 835-847.
4. Cavaleiro, J. A. S.; Gerdan, V. M.; Hombrecher, H. K.; Neves, M. G. P. M. S.; Silva, A. M. S. *Heterocycl. Commun.* **1997**, *3*, 253-261.
5. Hombrecher, H. K.; Gerdan, V. M.; Cavaleiro, J. A. S.; Neves, M. G. P. M. S. *Heterocycl. Commun.* **1997**, *3*, 453-461.
6. Köpke, T.; Pink, M.; Zaleski, J. M. *Synlett* **2006**, *14*, 2183-2186.
7. Galardon, E.; Le Maux, P.; Simonneaux, G. *J. Chem. Soc., Perkin Trans. 1* **1997**, 2455-2456.
8. Galardon, E.; Le Maux, P.; Simonneaux, G. *Tetrahedron* **2000**, *56*, 615-621.
9. Srour, H. F.; Le Maux, P.; Chevance, S.; Carrie, D.; Le Yondre, N.; Simonneaux, G. *J. Mol. Cat. A.: Chem.* **2015**, *407*, 194-203.
10. Nicolas, I.; Roisnel, T.; Le Maux, P.; Simonneaux, G. *Tetrahedron Lett.* **2009**, *50*, 5149-5151.
11. Simonneaux, G.; Tagliatesta, P. *J. Porphyrins Phthalocyanines* **2004**, *8*, 1166-1171.
12. Toma, T.; Shimokawa, J.; Fukuyama, T. *Org. Lett.* **2007**, *9*, 3195-3197.
13. Tamiaki, H.; Kumon, K.; Shibata, R. *J. Porphyrins Phthalocyanines* **2007**, *11*, 434-441.
14. Le Maux, P.; Nicolas, I.; Chevance, S.; Simonneaux, G. *Tetrahedron* **2010**, *66*, 4462-4468.
15. Collman, J. P.; Rose, E.; Venburg, G. D. *J. Chem. Soc., Chem. Commun.* **1993**, 934-935.
16. Le Maux, P.; Roisnel, T.; Nicolas, I.; Simonneaux, G. *Organometallics* **2008**, *27*, 3037-3042.
17. Woo, L. K.; Smith, D. A. *Organometallics* **1992**, *11*, 2344-2346.
18. Graban, E.; Lemke, F. R. *Organometallics* **2002**, *21*, 3823-3826.
19. Oshima, T.; Nagai, T. *Tetrahedron Lett.* **1980**, *21*, 1251-1254.
20. Shankar, B. K. R.; Shechter, H. *Tetrahedron Lett.* **1982**, *23*, 2277-2280.
21. Hodgson, D. M.; Angrish, D. *Chem. Eur. J.* **2007**, *13*, 3470-3479.
22. Galardon, E.; Le Maux, P.; Simonneaux, G. *J. Chem. Soc., Chem. Commun.* **1997**, 927-928.
23. Zaragoza, F. *Tetrahedron* **1997**, *53*, 3425-3439.
24. Aviv, I.; Gross, Z. *Chem. Commun.* **2006**, 4477-4479.
25. Baumann, L. K.; Mbuvi, H. M.; Du, G.; Woo, L. K. *Organometallics* **2007**, *26*, 3995-4002.
26. Aviv, I.; Gross, Z. *Chem. Eur. J.* **2008**, *14*, 3995-4005.
27. Ho, C. M.; Zhang, J. L.; Zhou, C. Y.; Chan, O. Y.; Yan, J. J.; Zhang, F. Y.; Huang, J. S.; Che, C. M. *J. Am. Chem. Soc.* **2010**, *132*, 1886-1894.
28. Shaw, E. *Methods Enzymol.* **1994**, *244*, 649-656.

29. All new compounds reported here gave spectral data consistent with the assigned structures. Reaction conditions and selected data: For **1**. To a distilled CH<sub>2</sub>Cl<sub>2</sub> solution (5 mL) of Zinc 5-[4-(hydroxymethyl)phenyl]-10,15,20-triphenylporphyrin<sup>13</sup> (100 mg, 0.14 mmol), 3 eq of DBU (0.42 mmol) were first added under argon at 0°C and then 1.5 eq of bromo acetyl bromide (0.21 mmol). The reaction mixture was stirred for 10 min at room temperature. After cooling the solution again to 0°C, a THF solution of ditosylhydrazine (2 eq) and DBU (5 eq) was added and the mixture was stirred for 30 min at room temperature. The solution was then evaporated, dissolved in CH<sub>2</sub>Cl<sub>2</sub> and purified through a silica gel column (CH<sub>2</sub>Cl<sub>2</sub>). Yield = 72%. <sup>1</sup>H NMR (CD<sub>2</sub>Cl<sub>2</sub>, ppm, 400 MHz): δ 8.99 (m, 8H, β pyrrole), 8.33-8.20 (m, 8H, phe), 7.90-7.72 (m, 11H, phe), 5.55 (s, 2H, CH<sub>2</sub>O), 5.00 (s (broad), 1H, CHN<sub>2</sub>). <sup>13</sup>C NMR (CD<sub>2</sub>Cl<sub>2</sub>, ppm, 125 MHz): δ 166.8(CO), 150.1, 142.5, 135.5, 134.5, 132.0, 127.5, 126.5, 126.2, 121.1, 66.4 (CH<sub>2</sub>O), 30.6 (CHN<sub>2</sub>). HRMS (ESI): calcd for C<sub>47</sub>H<sub>30</sub>N<sub>6</sub>O<sub>2</sub>Zn (M+H)<sup>+</sup>: 774.17162, found: 774.1719. UV-vis (CH<sub>2</sub>Cl<sub>2</sub>): λ<sub>max</sub>, nm (ε cm<sup>-1</sup> mM<sup>-1</sup>): 421 (365.90), 549 (14.39), 592 (3.47). For **2**. To a distilled CH<sub>2</sub>Cl<sub>2</sub> solution (1 mL) containing 0.9 mg of TPPRuCO, 8.6 mg (0.01 mmol) of **1** dissolved in 2 ml of CH<sub>2</sub>Cl<sub>2</sub> was added under argon. The mixture was stirred for 24h at room temperature. The solution was then evaporated and purified by chromatography on silica gel column (CH<sub>2</sub>Cl<sub>2</sub>). Yield = 91 %. <sup>1</sup>H NMR (CD<sub>2</sub>Cl<sub>2</sub>, ppm, 400MHz): δ 9–8.85 (m, 16H, β pyrrole), 8.3-8.1 (m, 18H, Phe), 7.90-7.72 (m, 20H, Phe), 6.62 (s, 2H, CH<sub>2</sub>), 5.67 (s, 4H, CH<sub>2</sub>O). <sup>13</sup>C NMR (CD<sub>2</sub>Cl<sub>2</sub>, ppm, 100 MHz): δ 165.2 (CO), 151.0, 143.0, 135.1, 134.9, 132.5, 130.1 (CH<sub>cis</sub>), 127.5, 126.4, 126.2, 67.2(CH<sub>2</sub>O). HRMS (ESI): calcd for C<sub>94</sub>H<sub>60</sub>N<sub>8</sub>O<sub>4</sub>Zn<sub>2</sub> (M)<sup>+</sup>: 1492.33149, found: 1492.3311. UV-vis (CH<sub>2</sub>Cl<sub>2</sub>): λ<sub>max</sub>, nm (ε cm<sup>-1</sup> mM<sup>-1</sup>): 421 (485.59), 550 (20.58), 594 (5.15). For **3**. To a distilled CH<sub>2</sub>Cl<sub>2</sub> solution (1mL) containing 25 μl of styrene and 0.9 mg of TPPRuCO under argon, 15.8 mg of **1** dissolved in 2 ml of CH<sub>2</sub>Cl<sub>2</sub> was added progressively (1ml/h). The mixture was stirred overnight. The solution was then evaporated and purified by chromatography on silica gel column. Yield = 79 %. <sup>1</sup>H NMR (CD<sub>2</sub>Cl<sub>2</sub>, ppm, 400 MHz): δ 8.99 (m, 8H, pyrrole), 8.31-8.21 (m, 8H, phe), 7.90-7.72 (m, 11H, phe), 7.38-7.30 (m, 2H, phe), 7.28-7.23 (m, 1H, phe), 7.22-7.16 (m, 2H, phe), 5.53 (s, 2H, CH<sub>2</sub>), 2.64-2.57 (m, 1H, cyclo), 2.12-2.05 (m, cyclo), 1.72-1.64 (m,1H, cyclo), 1.49-1.41 (m,1H, cyclo). <sup>13</sup>C NMR (CD<sub>2</sub>Cl<sub>2</sub>, ppm, 100 MHz): δ 173.6 (CO), 150.1, 142.8, 142.7, 140.2, 135.6, 134.5, 132.0, 128.4, 127.5, 126.5, 126.2, 121.6, 121.2, 66.4, 26.2, 24.1, 17.1. HRMS

(ESI): calcd for  $C_{55}H_{38}N_4O_2Zn$  ( $M+H$ )<sup>+</sup>: 850.22807, found: 850.2276. UV-vis ( $CH_2Cl_2$ ):  $\lambda_{max}$ , nm ( $\epsilon$  cm<sup>-1</sup> mM<sup>-1</sup>): 421 (357.50), 550 (10.43), 592 (1.08). For **4**. To a distilled  $CH_2Cl_2$  solution (1mL) containing 2.3  $\mu$ l of triethylamine (1.5 eq), 0.6 mg of TPPFeCl and 2.8 mg (0.01 mmol) of L- tryptophan methyl ester hydrochloride, 8.6 mg (0.01 mmol) of **1** dissolved in 2 ml of  $CH_2Cl_2$  was added under argon. The mixture was stirred for 15 min. The solution was then evaporated and purified by chromatography on silica gel column ( $CH_2Cl_2/CH_3OH$ : 98/2). Yield = 69%. <sup>1</sup>H NMR ( $CD_2Cl_2$ , ppm, 400 MHz):  $\delta$  8.97 (m, 8H,  $\beta$  pyrrole), 8.36-8.24 (m, 8H, Phe), 7.93 (s, 1H, NH), 7.81-7.60 (m, 11H, Phe), 7.27 (d, 1H, trypt), 7.11 (t, 1H, trypt), 6.94 (t, 1H, trypt), 6.84 (d, 1H, trypt), 5.87 (s, 1H, trypt), 4.93 (s, 2H,  $CH_2O$ ), 3.05 (s, 3H,  $CH_3$ ), 1.4 and 0.5 (m, 2H,  $CH_2$ ), 0.4 (m, 2H,  $CH_2N$ ), 0.39 (m, 1H,  $CH^*$ ). <sup>13</sup>C NMR ( $CD_2Cl_2$ , ppm, 100 MHz):  $\delta$  169.5 (CO), 168.0 (CO), 149.1, 141.5, 135.0, 134.5, 132.0, 127.0, 126.5, 126.0, 121.1, 110.0, 67 ( $CH_2O$ ), 57.5 (C\*), 43.0 ( $CH_2N$ ), 52.0 ( $CH_3$ ), 24.8 ( $CH_2$ ). HRMS (ESI): calcd for  $C_{59}H_{45}N_6O_4Zn$  ( $M+H$ )<sup>+</sup>: 965.27882, found: 965.2790. UV-vis ( $CH_2Cl_2$ ):  $\lambda_{max}$ , nm ( $\epsilon$  cm<sup>-1</sup> mM<sup>-1</sup>): 421 (272.90), 551 (11.47), 592 (1.90). For **5**. To a distilled toluene solution (1mL) o2.86 mg (0.026 mmol) of thiophenol and 0.6 mg of TPPRuCO under argon, 2.9 mg (0.02 mmol) of **1** dissolved in 0.5 ml of toluene was added over a period of 30 min. The reaction was then stirred for 4 hours at room temperature. The mixture was concentrated by vacuum evaporation and purified by silica gel chromatography. Yield = 85%). <sup>1</sup>H NMR ( $CD_2Cl_2$ , ppm, 400 MHz):  $\delta$  9.10-8.90 (m, 8H, pyrrole), 8.35-8.15 (m, 8H, phe), 7.95-7.75 (m, 9H, phe), 7.75-7.68 (m, 2H, phe), 7.56-7.50 (m, 2H, phe), 7.43-7.35 (m, 1H, phe), 7.33-7.25 (m, 2H, phe) 5.52 (s, 2H,  $CH_2O$ ), 3.88 (s, 2H,  $CH_2S$ ). <sup>13</sup>C NMR ( $CD_2Cl_2$ , ppm, 100 MHz):  $\delta$  169.7 (CO), 150.3, 142.9, 127.0, 134.5, 132.0, 130, 129.1, 127.5, 127.1, 126.6, 126.3, 121.2, 120.4, 67.36 ( $CH_2O$ ), 36.6 ( $CH_2S$ ). HRMS (ESI): calcd for  $C_{53}H_{36}N_4O_2SZn$  ( $M+H$ )<sup>+</sup>: 856.18449, found: 856.1841. UV-vis ( $CH_2Cl_2$ ):  $\lambda_{max}$ , nm ( $\epsilon$  cm<sup>-1</sup> mM<sup>-1</sup>): 422 (251.76), 550 (8.93), 592 (1.32).



Highlights :

First synthesis of Zinc porphyrin diazoacetates

NH of tryptophan ester by porphyrin diazo esters

Cyclopropanation of styrene by porphyrin diazo esters

Preparation of porphyrin dimers by diazo coupling reaction

SH insertion of thiophenol by porphyrin diazo esters

ACCEPTED MANUSCRIPT