

Supplementary Information

Enantioselective recognition by highly ordered porphyrin-assembly on chiral molecular gel

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1. Materials and Generals

5-(4-methoxycarbonylphenyl)-10,15,20-triphenyl-21H,23H-porphine and amino acid methyl esters were purchased from Wako Pure Chemical Industries and Tokyo Chemical Industry, respectively. *N*^l,*N*⁵-didodecyl-L-glutamide (**g**) and zinc porphyrin lipid (**g-TPP/Zn**) were synthesized by the previously reported procedure with slight modification. UV-visible, CD and fluorescence spectra were measured with V-560 (JASCO), J725 (JASCO) and FP-6500 (JASCO), respectively. The Stern-Volmer constants were determined in the concentrations below 0.30 equimolar of His-OMe for **g-TPP/Zn**. (50 μM).

2. Sample Preparation for Spectroscopic Measurement¹

- (1) The free amino acid methyl ester solution was prepared from its hydrochloride salt. 10 mg of amino acid methyl ester was dissolved in 1.0 mL of methanol with Na₂CO₃ (100 mg).
- (2) Solvent (methanol) was removed under a stream of nitrogen and then placement under high vacuum (0.1 Torr) for an hour.
- (3) Toluene was added to yield the free amino acid methyl ester solution (4.0 mM).
- (4) An aliquot of 5 μL of the amino acid methyl ester solution (1 equiv) was added to the prepared 0.05 mM of zinc porphyrin lipid (**g-TPP/Zn**) solution to obtained **g-TPP/Zn**/ amino acid methyl ester complex.

1. X. Huang, B. H. Rickman, B. Borhan, N. Berova, and K. Nakanishi, *J. Am. Chem. Soc.*, 1998, **120**, 6185–6186.

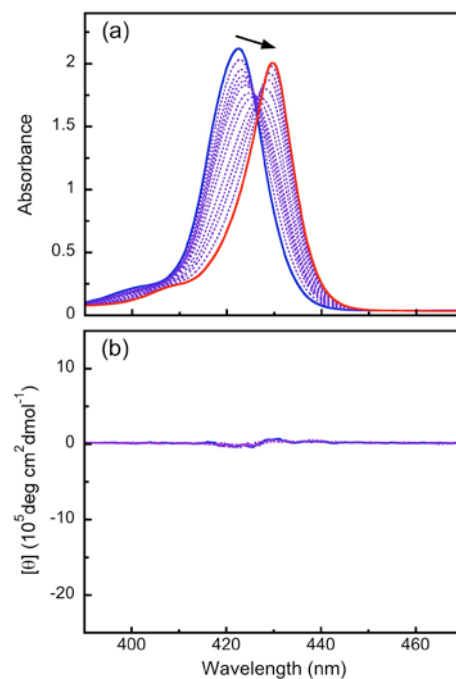


Figure S1. (a) UV-vis and (b) CD spectral changes of *g*-TPP/Zn (50 μM) upon addition of L-His-OMe (0 - 500 μM) in chloroform at 20 °C.

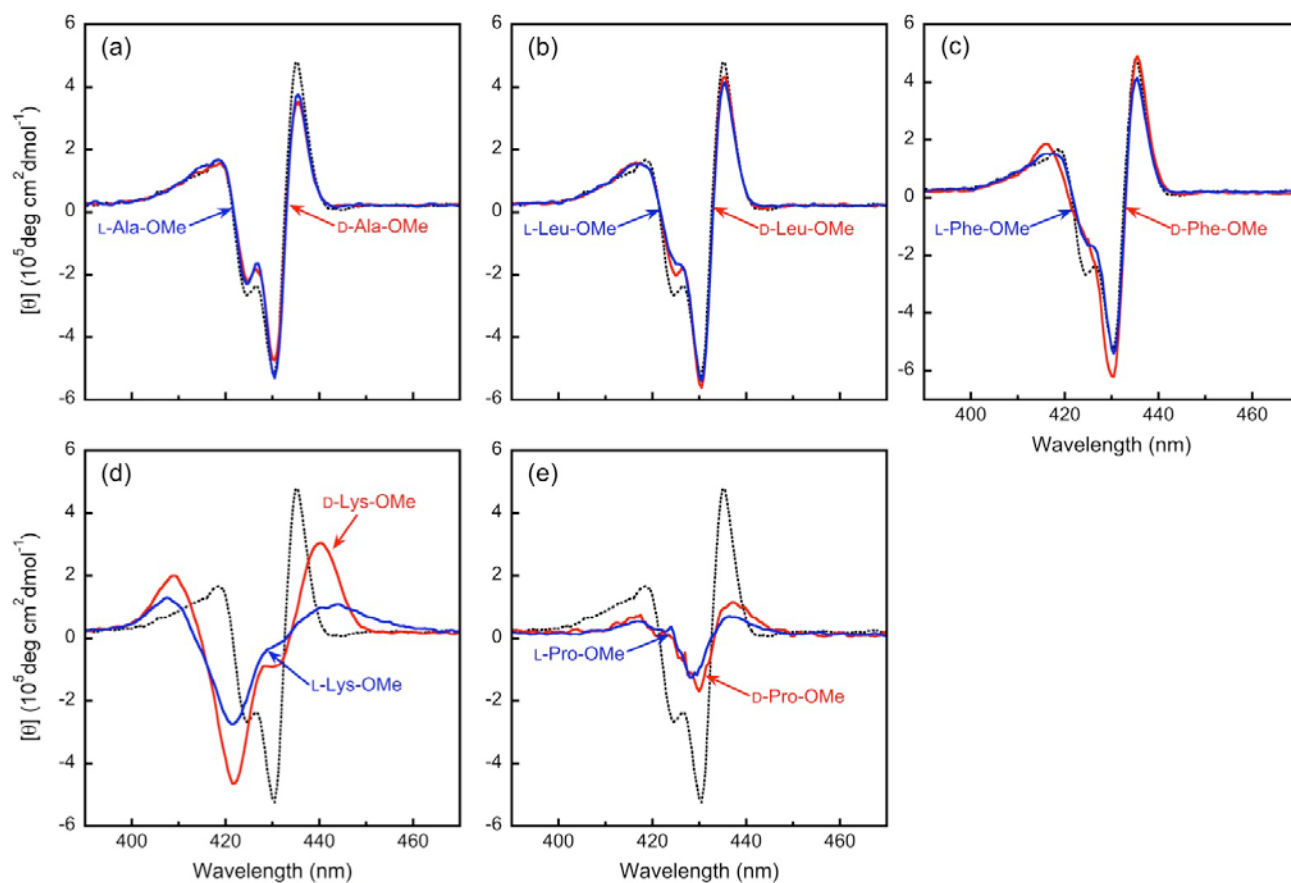


Figure S2. CD spectra of *g*-TPP/Zn (50 μM) with and without an equimolar of (a) Ala-OMe, (b) Leu-OMe, (c) Phe-OMe (d) Lys-OMe, and (e) Pro-OMe in cyclohexane at 20 °C.