Ruthenium(IV) N-confused porphyrin μ-oxo-bridged dimers: acid-responsive molecular rotors†

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Ruthenium(IV) N-confused porphyrin μ-oxo-bridged complexes were synthesized via oxidative dimerization of a ruthenium(II) N-confused porphyrin complex using 2,2,6,6-tetramethylpiperidine 1-oxyl. The deformed core planes in the dimers conferred a relatively high ring rotational barrier of ca. 16 kcal mol⁻¹. Rotation of the complexes was controlled by protonating the peripheral nitrogen.

The design/creation/operation of molecular rotors, gears, and switches is an interesting topic in machinery nanotechnology, allowing the development of adaptive materials, switchable chiral sensors, stereo-controlled catalysts, etc.¹ The molecular approach to control nanoscale motions (e.g., rotation) is advantageous for realizing well-defined molecular devices or sets of practical supramolecular assembled systems.² As active components of molecular rotors, metal double-decker bisporphyrins and related macrocycles (e.g., phthalocyanines) are representative candidates that afford specific features, such as long-term stability, ease of fabrication, and visual responses that are detectable by microscopy (Fig. 1a).³ Bis-metal μ-oxo (or nitride)-bridged porphyrin complexes are also considered alternatives for modulating complexes’ rotational mode. However, there are few investigations concerning molecular rotation because of the limitation in synthesizing molecular rotamers comprising unsymmetrical core skeletons with diamagnetic and oxophilic metal ions (for which the rotational dynamics can be probed by NMR spectroscopy) (Fig. 1b).⁴

To develop new rotors based on metal μ-oxo-bridged porphyrin dimers, we envisioned that a C₄ᵥ-symmetric N-confused porphyrin (NCP) isomer possessing an inverted pyrrole ring in the macrocyclic scaffold could act as an alternative ring motif for the complexes (Fig. 1c).⁵ The NCPs bind various metal ions at the NNNC core, forming unique organometallic complexes.⁶ For example, redox-active metal species such as iron facilitate the generation of metal–oxo linkages to form μ-oxo-bridged dimer complexes.⁷ In the case of iron-coordination of NCP, a unique bis-iron μ-hydroxo-bridged NCP dimer was developed, in which the structure was restricted by the coordination of peripheral sodium, as evident from X-ray crystallography (Fig. 1d).⁸

Fig. 1 Chemical structures of (a) double-decker metal porphyrin complexes and (b) bis-metal μ-oxo-bridged porphyrin complex as representative motifs of molecular rotors (R¹ = R³ = p-methoxyphenyl, R² = R⁴ = 4-pyridyl). (c) N-confused porphyrin as a key rotor ligand and its (d) bis-iron μ-hydroxo-bridged porphyrin complex. (e) Schematic of the rotational mode of unsymmetrical N-confused porphyrin μ-oxo dimer complex. Sky-blue moiety in the planes is a confused pyrrole ring.
We herein report the synthesis of novel ruthenium(IV) NCP µ-oxo-bridged dimers \( \text{Ru}_2-2 \) and \( \text{Ru}_2-3 \) as a porphyrin-based molecular rotor. The ruthenium(n) NCP complex \( \text{Ru}-1 \) with an axial carbonyl and pyridine (−py) exhibits catalytic reactivity for alkene epoxidation (Scheme 1). The simultaneous formation of isomers can be anticipated owing to the two stereogenic centers in the NCP planes of \( \text{Ru}_2-2 \) and \( \text{Ru}_2-3 \), which differ in the relative position of the nitrogen atoms of the confused pyrrole rings in the scaffold. The \( \mu \)-oxo dimers showed distinct ring rotation in solution when probed via NMR spectroscopy (Fig. 1e). Moreover, symmetry-dependent restriction of the rotation in solution was achieved through protonation at the peripheral amino-nitrogen moieties in \( \text{Ru}_2-2 \) and \( \text{Ru}_2-3 \) using trifluoroacetic acid (TFA).

Treatment of monomer \( \text{Ru}-1 \) with 2,2,6,6-tetramethylpiperidine 1-oxyl (TEMPO) in refluxing o-dichlorobenzene solution under argon atmosphere afforded two products with approximately the same yields (40% each) (Scheme 1). The fast atom bombardment mass spectra of the two products revealed m/z 1430.2 (M+) with estimated mean plane deviation values (de 0.276 Å and 0.294 Å for \( \text{Ru}_2-2 \) and \( \text{Ru}_2-3 \) and 0.286 Å and 0.292 Å for \( \text{Ru}_2-3 \).

As anticipated above, the formal oxidation state of both ruthenium centers is expected to be +4 (d^4) for the complexes. The X-ray photoelectron spectra of \( \text{Ru}_2-2 \) and \( \text{Ru}_2-3 \) revealed higher Ru (3p_3/2) binding energies of 462.2 and 462.1 eV, respectively, than that of the divalent ruthenium (d6) complex, \( \text{Ru}_2 \) (460.9 eV, Fig. S3 in ESI†).

The structures of \( \text{Ru}_2-2 \) and \( \text{Ru}_2-3 \) were further characterized by \( ^1 \)H NMR spectroscopy in 1,1,2,2-tetrachloroethane-d2 (TCE-d2) (Fig. 3a and b). At 298 K, complicatedly overlapped signals were observed in the aromatic region (7-10 ppm). To distinguish the signals from the individual groups (β-pyrrole and meso-phenyl) in the complexes, we prepared the deuterated-phenyl derivatives (i.e., \( \text{Ru}_2-2D \) and \( \text{Ru}_2-3D \)) as shown in Scheme 2 (see, ESI†). The \( ^1 \)H NMR spectra of \( \text{Ru}_2-2D \) and \( \text{Ru}_2-3D \) (Fig. S4†) show two singlets at 9.38 and 9.89 ppm for \( \text{Ru}_2-2 \) and 9.32 and 9.05 for \( \text{Ru}_2-3 \). The remaining six β-CHs are expected to have signals around 8 ppm for both complexes. This spectral feature suggests the existence of two possible rotational isomers in solution with an almost 1:1 molecular ratio. The 1D exchange nuclear Overhauser effect spectra revealed chemical exchange of the rotamers at 298 K (Fig. S5 and S6†), which

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**Scheme 1** Synthesis of ruthenium(IV) NCP µ-oxo dimers, \( \text{Ru}_2-2 \) and \( \text{Ru}_2-3 \).

**Fig. 2** Top (left) and side (right) views of crystal structures of (a) \( \text{Ru}_2-2 \) and (b) \( \text{Ru}_2-3 \) with thermal ellipsoids set at 50% probability level. Hydrogen atoms and solvent molecules excepting the H-atoms hydrogen-bonded to the peripheral nitrogen atoms were omitted for clarity.
indicated that the NCP rings of Ru2-2 and Ru2-3 rotate slowly at 298 K. The rotational mode of the NCP rings was seemingly restricted at 243 K (Fig. S7 and S8†). With increasing temperature, the signal of the set of \(1H\)-pyrrole protons gradually disappeared and coalesced into a single peak at 9.15 ppm for Ru2-2 and 9.27 ppm for Ru2-3 at 408 K (Fig. 3c and d). Signals of the meso-phenyl protons also appeared with an averaged spectral feature in the region of 7.5–8.0 ppm (Fig. S9 and S10†). These results suggest that the rotation rate of the NCP rings is faster than the NMR timescale above 353 K. Furthermore, for the ruthenium(IV) tetrakis(phenyl)-substituted porphyrin \(\mu\)-oxo dimer (e.g., Ru(TPP)\(_2\)O) used as the reference complex, the inward and outward \(1H\) signals (\(i.e., \alpha\) - and \(m\)-CHs) of the meso-phenyl rings did not coalesce even at 408 K in the variable temperature \((VT\) NMR spectra (Fig. S11†). In this regard, the rotation of the bulky phenyl ring could be restricted by the spatially locked dimeric structure. The rotation mode of the NCP planes could be thus mutually un-synchronized with the rotational motion of the individual meso-phenyl rings.

To gain further insight into the rotational barrier of the complexes, a line-shape analysis of the VT \(1H\) NMR spectra was conducted to estimate the rate constant for interconversion \((k)\) in the specific temperature range. The free Gibbs’s energy \((\Delta G^\ddagger = \Delta H^\ddagger - T\Delta S^\ddagger)\) of Ru2-2 and Ru2-3 was determined to be \((16.0 \pm 0.8)\) kcal mol\(^{-1}\) and \((16.1 \pm 0.4)\) kcal mol\(^{-1}\), respectively, at 298 K, from the Eyring plots (Fig. S12†). The rate constant \((k < 200\) s\(^{-1}\)) at 333 K indicates that the energy barrier of the ring rotations of Ru2-2 and Ru2-3 is more significant than that of the double-decker metal complexes \((k = 760\) s\(^{-1}\) at 333 K).\(^{14}\)

Considering the structure of the complexes, the bent confused pyrrole moieties in the planes may play an essential role in the rotational motion beyond the barrier.

Upon lowering the temperature, the ratio of the integrated \(1H\)-CH signals for the rotamer mixture of Ru2-2 changed to 0.85 (at 9.33 ppm) and 0.15 (at 8.93 ppm) (Fig. S13†). The ratio of the two-rotamers of Ru2-3 also changed when the temperature was further lowered \((-223 K\), affording a thermodynamically stable conformer (Fig. S14†). To understand the structure-dependent features of Ru2-2 and Ru2-3, we analyzed the thermodynamic energy profiles of the possible rotational isomers \((e.g., Ru2-2a-f and Ru2-3a-c)\) by theoretical calculation with the B3LYP method (Tables S3 and S4†). The most stable isomers \((i.e., Ru2-2c and Ru2-3c)\) obtained in the optimization process are highly consistent with the crystal structures. Notably, the torsion angles of C17–Ru1–Ru2–C61(C62) in the structures of Ru2-2c and Ru2-3b are \(-102.8°\) and 98.5°, respectively, indicating stability \((\Delta E = +1.19 + 1.45\) kcal mol\(^{-1}\), respectively relative to Ru2-2c/-3c. Therefore, two energetically close conformers are present in the solution under the experimental conditions.

The effect of protonation on the rotation of the rotamers was evaluated.\(^{11}\) Upon addition of trifluoroacetic acid (TFA), the absorption spectra of Ru2-2 and Ru2-3 gradually changed with isosbestic points (Fig. S15 and S16†), reflecting protonation at the peripheral imino-nitrogen of the confused pyrrole moieties (Fig. 4a and b).\(^{14}\) In both complexes, a relatively large amount of TFA (at least 8 equiv) was necessary until spectral saturation (Fig. S15a and S16a). Strikingly, in the case of Ru2-3, the relatively crowded imino-nitrogen site surrounded by meso-phenyl rings interfered with double protonation, as seen in the acid-spectral titration change after the addition of 8 equiv. of TFA (Fig. 4d, e, S15b and S16b†). In the \(1H\) NMR spectrum, the resulting dication species of Ru2-2 \((i.e., Ru2-2H_2^{2+})\) was well-resolved, demonstrating the interlocked single isomer at 298 K, as evident from the NMR spectral features (Fig. S17a†). The

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VT NMR spectra revealed that ring rotation becomes feasible with increasing temperature due to the dissociation of the TFA molecule from the complex at equilibrium (Fig. S18f). In sharp contrast, the $^1$H NMR spectrum of the protonated species of the anti-symmetrical complex (i.e., Ru$_2$-3H$^+$) exhibited a significantly broadened spectral feature at 298 K in TCE-d$_2$. The unsymmetrical spectral feature was observed at 243 K (Fig. S17b†), which may be due to the above steric issue in the conformation of Ru$_2$-3. Therefore, the predominant formation of mono-protonated species was anticipated in Ru$_2$-3. Eventually, upon the addition of triethylamine as a base, the original spectral signal of the neutral species was reproduced for both complexes (Fig. S15c and S16c†).

In summary, novel cofacial ruthenium(IV) NCP μ-oxo dimers, Ru$_2$-2 and Ru$_2$-3, were synthesized as motifs for molecular rotors. The resulting complexes, Ru$_2$-2 and Ru$_2$-3, are syn/anti-stereo-isomers having the nitrogen at different positions of the confused pyrrole rings. The symmetrical difference between the complexes leads to distinct rotational behavior of the Ru–NCP planes through the oxo-axis of the complex, as proven by VT $^1$H NMR analysis. Additionally, considering the intrinsic basicity of the peripheral nitrogen sites, the rotational profile can be controlled by protonation. Further study on the cofacial metal–μ-oxo NCP dimers can enable applications in molecular machinery.

Conflicts of interest
There are no conflicts to declare.

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Notes and references


11 There are structural disorders (occupancies = 75% and 65% for each plane in Ru$_2$-2; occupancies = 75% and 56% in Ru$_2$-3) originating from the positions of carbon and nitrogen atoms of the confused pyrrole rings in the crystals. The CHCl$_3$ solvent molecules co-crystallized in the structure point to the external nitrogen site with possible hydrogen-bonding interaction.


14 We have obtained the preliminary crystal structure of the protonated species of Ru$_2$-2 from the CHCl$_3$/hexane solution in the presence of TFA. The discrete syn conformer possessing the C17–Ru1–Ru2–C62 torsion angle of $-150.9(4)^\circ$ was speculated in the crystal structure. See Fig. S19 in ESL†