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Relativistic effects in metallocorroles: Comparison of molybdenum and tungsten biscorroles

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The homoleptic sandwich compounds – Mo and W biscorroles – have afforded a novel platform for experimental studies of relativistic effects. A 200-mV difference in reduction potential and a remarkable 130-nm shift of a near-IR spectral feature have been identified as manifestations of relativistic effects on the properties of these complexes.

15 Detailed comparative studies of analogous 4d and 5d transition metal complexes can potentially provide insights into the importance of relativistic effects.^{1,2,3,4,5,6} Yet, there have been relatively few such studies on typical inorganic and organometallic complexes^{7,8,9} and no more than a couple on metalloporphyrins and related complexes.¹⁰ Important opportunities in this regard are provided by recent syntheses of several 5d metallocorroles,^{11,12,13,14,15,16,17,18} which have garnered attention as unique size-mismatched complexes combining a large 5d transition metal and a sterically constrained corrole ligand. Thus, in a recent study, we were able to attribute a ~28-nm redshift in the Soret maxima of rhenium(V)-oxo corroles, relative to analogous technetium(V)-oxo corroles, to relativistic effects.¹⁹ Molybdenum biscorroles, reported herein, and their tungsten analogues, reported previously,²⁰ now provide an opportunity for experimental studies of relativistic effects in these unique Group 6 sandwich compounds.

The synthetic coordination chemistry of Mo and W corroles already points to significant chemical differences between the two metals. Whereas the high-temperature interaction of Mo(CO)₆ and a free-base *meso*-tris(*p*-X-phenyl)corrole, H₃[TpXPC], readily affords Mo^V[TpXPC]O,²¹ the corresponding W^VO complexes are unknown.²² Instead, upon refluxing in decalin in the presence of K₂CO₃, W(CO)₆ and H₃[TpXPC] were found to yield W[TpXPC]₂ (X = CF₃, H, Me), the only examples of homoleptic corrole sandwich compounds reported to date.¹⁴ While the reason underlying the elusiveness of W^VO corroles remains unclear, we conjectured that the interaction of H₃[TpXPC] and Mo(CO)₆ under strict exclusion of oxygen might afford Mo biscorroles. Indeed, upon prolonged heating (186 °C, 16 h) in decalin under anoxic conditions, the interaction of H₃[TpXPC], Mo(CO)₆, and K₂CO₃ afforded, after work-up, 25–30% yields of Mo[TpXPC]₂ along with 15–20% yields of Mo^V[TpXPC]O, where X = H, Me, OMe. As delineated below,

Table 1. Soret and Q maxima (nm), redox potentials (V), and electrochemical HOMO-LUMO gaps (V) of M[TpXPC]₂ (M = Mo, W) as a function of X.

X	UV-vis		$E_{1/2ox2}$	$E_{1/2ox1}$	$E_{1/2red1}$	$E_{1/2red2}$	DE
<i>Molybdenum</i>							
H	356	908	0.72	0.23	-0.83	-1.78	1.06
Me	362	909	0.67	0.18	-0.88	-1.84	1.06
OMe	350	909	0.66	0.19	-0.89	-1.85	1.08
<i>Tungsten</i>							
CF ₃	356	778	0.88	0.45	-0.88	-1.71	1.33
H	357	781	0.75	0.25	-1.07	-1.79	1.32
Me	359	783	0.72	0.21	-1.11	-1.82	1.32

electrochemical and UV-vis measurements indeed reveal multiple differences between Mo and W biscorroles that may be ascribed to relativistic effects (Table 1).

As with the previously reported W biscorroles,¹⁴ the ¹H NMR spectra (including variable temperature experiments) of Mo[TpXPC]₂ exhibited strongly overlapping peaks in the aromatic region, which defied assignment. For X = Me and OMe, however, the aliphatic regions exhibited three equally intense methyl signals, consistent with C₁ local symmetry of the individual corroles and overall C₂ symmetry of the complexes. Fortunately, one of the complexes, Mo[TpMePC]₂, proved amenable to single-crystal X-ray diffraction analysis (Figure 1), allowing a detailed structural comparison of the Mo and W complexes.²³ Thus, both metals exhibit square antiprism coordination geometries and essentially identical M-N distances of ~2.2 Å and M-N₄ displacements of 1.18 Å.

The Mo[TpXPC]₂ derivatives exhibit two reversible oxidations and one reversible reduction in CH₂Cl₂ (Figure 2). As shown in Table 1, for a given *para* substituent X, the first oxidation potentials of Mo and W biscorroles are essentially identical. The first reduction potentials of Mo biscorroles on the other hand are some 200 mV higher than those of their W counterparts. The more negative reduction potentials of the W biscorroles relative to Mo raise the question whether the difference arises, to a

Figure 1. Thermal ellipsoid (50%) plots for Mo[TpMePC]₂: (a) side and (b) top views; (c) Mo-N distances (Å).

Figure 2. Cyclic voltammograms of Mo[TpXPC]₂ (X = H, Me and OMe).

The above differences between Mo and W corroles were investigated with DFT and TDDFT calculations,^{24,25} with relativistic effects taken into account with the zeroth order regular approximation (ZORA) to the two-component Dirac equation applied as a scalar correction. Use of a nonrelativistic Hamiltonian in all-electron B3LYP/STO-TZP calculations resulted in nearly identical HOMO and LUMO orbital energies, ionization potentials and electron affinities for Mo[TPC]₂ and W[TPC]₂. Deploying the ZORA Hamiltonian upshifted the LUMO orbital energy of W[TPC]₂ by ~200 meV, relative to a nonrelativistic Hamiltonian, but had a much smaller impact on Mo[TPC]₂. The same effect was also reproduced with DSCF calculations of electron affinities, where the ZORA Hamiltonian decreased the electron affinity of W[TPC]₂, relative to a nonrelativistic Hamiltonian, by ~170 meV.^{26,27} For both Mo[TPC]₂ and W[TPC]₂, the LUMO has substantial metal(d_{z2}) character (Figure 4), which explains why it should be relativistically destabilized in W case. Thus, the ~200 mV difference in reduction potentials between the Mo[TpXPC]₂ and W[TpXPC]₂ series is largely attributable to relativistic effects in the W series.

Relativistic ZORA TDDFT (CAMY²⁸-B3LYP/STO-TZP/COSMO; solvent = CH₂Cl₂) calculations readily assigned the lowest-energy near-IR feature of both Mo[TPC]₂ and W[TPC]₂ to a (HOMO – 1)[]LUMO transition (Figure 4). Comparison with TDDFT calculations, which employed a nonrelativistic Hamiltonian but an otherwise identical set of computational parameters, clearly implicated relativistic destabilization of the 5d_{z2} orbital of W as the major contributor to the higher energy of this transition in the W case or, equivalently,

Figure 3. UV vis spectra of (a) Mo[TpXPC]₂ (X = H, Me and OMe) and of (b) Mo[TPC]₂ (blue) and W[TPC]₂ (orange).

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Figure 4. Selected frontier MOs and Mo[TPC]₂ along with their orbital energies (eV) and % d character.

10 to the dramatic redshift observed in the Mo case. The calculations did not, however, provide an assignment for the second, higher-energy near-IR feature of the Mo biscalloles, which suggests that it may be a combination/vibronic band. No significant relativistic effect was found for the Soret band position of Mo[TPC]₂ and W[TPC]₂, in apparent agreement with the near-identical Soret maxima observed experimentally for both compounds (Figure 3).

Conclusions

20 As the sole representatives of homoleptic corrole sandwich compounds, Mo and W biscalloles provide a novel platform for experimental studies of relativistic effects. X-ray crystallographic analyses have shown that the two metals exhibit near-identical M-N distances (~2.2 Å) and M-N₄ displacements (~1.18 Å). The Mo complexes exhibit distinctly higher reduction potentials (i.e., are easier to reduce), by a margin of ~200 mV, than the analogous W complexes. The UV-vis spectra of Mo biscalloles also exhibit a near-IR feature at 908-909 nm, which is redshifted by some 130 nm relative to an analogous feature of W biscalloles. TDDFT calculations have identified this feature as a (HOMO - 1) → LUMO transition. For both the Mo and W biscalloles, the LUMO has substantial metal d₂₂ character. Relativistic destabilization of the W(5d) orbitals thus explains both the higher reduction potentials and the lower energy of the near-IR transition for the Mo biscalloles.

Notes and references

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- 45 [†] Electronic Supplementary Information (ESI) available: Crystallographic information file (cif), which has also been deposited to the Cambridge Crystallographic Data Center and assigned the deposition no. [CCDC 1534096](#).
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