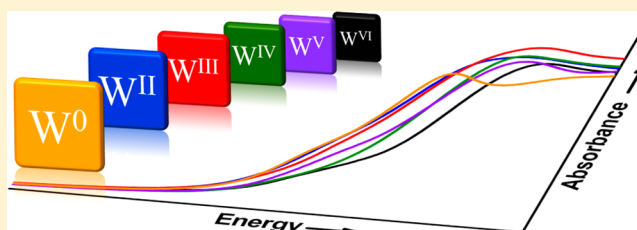


X-ray Absorption Spectroscopy Systematics at the Tungsten L-Edge

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Supporting Information

ABSTRACT: A series of mononuclear six-coordinate tungsten compounds spanning formal oxidation states from 0 to +VI, largely in a ligand environment of inert chloride and/or phosphine, was interrogated by tungsten L-edge X-ray absorption spectroscopy. The L-edge spectra of this compound set, comprised of $[W^0(PMe_3)_6]$, $[W^{II}Cl_2(PMePh_2)_4]$, $[W^{III}Cl_2(dppe)_2][PF_6]$ ($dppe = 1,2$ -bis(diphenylphosphino)ethane), $[W^{IV}Cl_4(PMePh_2)_2]$, $[W^V(NPh)Cl_3(PMe_3)_2]$, and $[W^{VI}Cl_6]$, correlate with formal oxidation state and have usefulness as references for the interpretation of the L-edge spectra of tungsten compounds with redox-active ligands and ambiguous electronic structure descriptions. The utility of these spectra arises from the combined correlation of the estimated branching ratio of the $L_{3,2}$ -edges and the L_1 rising-edge energy with metal Z_{eff} , thereby permitting an assessment of effective metal oxidation state. An application of these reference spectra is illustrated by their use as backdrop for the L-edge X-ray absorption spectra of $[W^{IV}(mdt)_2(CO)_2]$ and $[W^{IV}(mdt)_2(CN)_2]^{2-}$ ($mdt^{2-} = 1,2$ -dimethylethene-1,2-dithiolate), which shows that both compounds are effectively W^{IV} species even though the mdt ligands exist at different redox levels in the two compounds. Use of metal L-edge XAS to assess a compound of uncertain formulation requires: (1) Placement of that data within the context of spectra offered by unambiguous calibrant compounds, preferably with the same coordination number and similar metal ligand distances. Such spectra assist in defining upper and/or lower limits for metal Z_{eff} in the species of interest. (2) Evaluation of that data in conjunction with information from other physical methods, especially ligand K-edge XAS. (3) Increased care in interpretation if strong π -acceptor ligands, particularly CO, or π -donor ligands are present. The electron-withdrawing/donating nature of these ligand types, combined with relatively short metal–ligand distances, exaggerate the difference between formal oxidation state and metal Z_{eff} or, as in the case of $[W^{IV}(mdt)_2(CO)_2]$, exert the subtle effect of modulating the redox level of other ligands in the coordination sphere.



INTRODUCTION

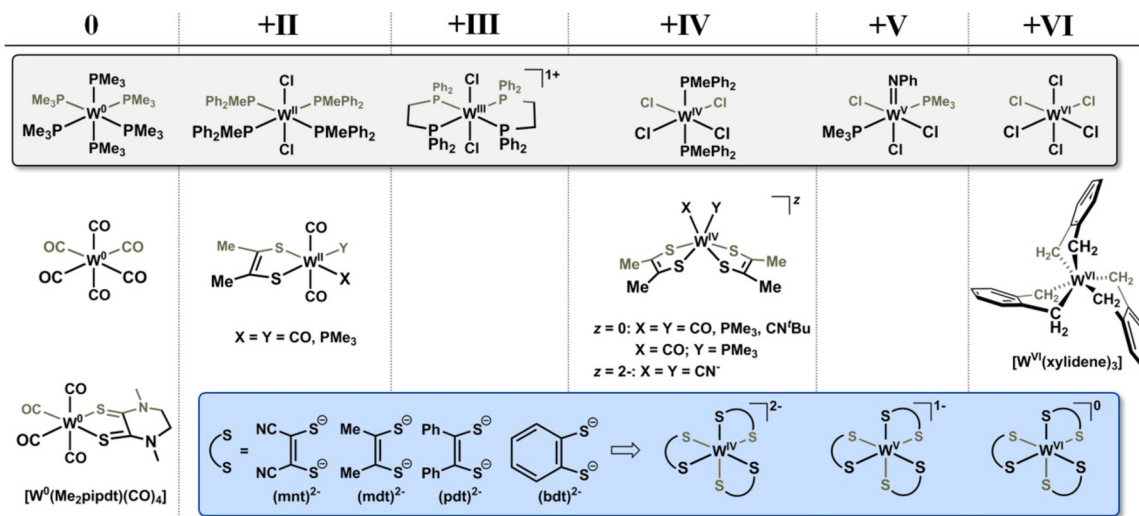
The distinction that may be made between the *formal* and *physical* oxidation state of a metal atom in a transition metal complex was first articulated by Jørgensen.¹ A formal oxidation state is a nonmeasurable integer commonly defined as the charge remaining on the metal after the ligands have been removed in their closed-shell form.² In principle, a physical oxidation state is a value derived from a measurable quantity (e.g., from spectroscopy) that diagnoses a d^n configuration.¹ Where these two descriptions can appreciably diverge is in situations where the bonding between metal and ligand(s) is substantially covalent such that the charge associated with the metal atom, evaluated by accounting methods that assume integer charges or complete neutrality to specific ligand types,³ differs from a metal effective nuclear charge (Z_{eff}) gauged by experimental methods. Redox-active ligands such as NO and heterodiene-type ligands, such as catecholates, *o*-diimines, and

dithiolenes, are widely recognized as “noninnocent” ligands that can readily assume radical character and lead to significant departures from the electron-counting schemes that partition charge in a heterolytic fashion. Recent work reported from several research groups, in which computational insights are typically used in conjunction with information from a battery of physical methods, has demonstrated that ligand redox activity extends beyond NO and heterodiene ligands to include such other ligand types as phenolates,⁴ corroles,^{5,6} bis(imino)pyridines,⁷ and tris(amido)-⁸ and amido bis(phenolate)⁹ pincer-type ligands.

From the perspective of understanding the intrinsic nature of a metal complex and anticipating its reactivity and properties, the spectroscopic oxidation state may constitute a preferred

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Chart 1. Pictorial Representation of the Complexes Organized According to Tungsten Formal Oxidation State^a

^aFor organizational purposes, all dithiolene-type ligands other than Me_2pipdt are treated as fully reduced.

basis for thought. Although no experimental method produces a metal “oxidation state” as immediate output, metal K-edge X-ray absorption spectroscopy (XAS), especially when taken in conjunction with ligand K-edge XAS data,^{10,11} is a direct experimental method for evaluating a metal Z_{eff} . The rising-edge is dominated by electric dipole-allowed $1s \rightarrow np$ transitions and is effective for experimentally gauging Z_{eff} because the core $1s$ orbital shifts to deeper binding energy upon oxidation of the metal. For first row metals, the core-hole lifetime is sufficiently long that the spectrum may be resolved into the pre-edge and near-edge constituents.¹² The pre-edge arises from electric quadrupole-allowed excitations to vacancies in the $3d$ manifold, which gain intensity through $d-p$ mixing via a departure from centrosymmetry and are diagnostic of the geometry of the complex.¹³ At this high resolution (1–1.5 eV), the pre-edge transition is the preferred metric for Z_{eff} .^{14–17} The corresponding K-edge for third row metals occurs at prohibitively high energies (>60 keV). Consequently, attaining detail about the local electronic structure is confined to L-edge XAS. These edges comprise three components: L_{3-} and L_{2-} edges are dipole-allowed excitations of core $2p$ electrons to partially filled or empty $5d$ -based orbitals, while the L_{1-} edge is dominated by $2s \rightarrow 6p$ dipole-allowed transitions. The latter represents a viable alternative to K-edge XAS; however, tangible electronic structure details are washed out by lifetime broadening of the $2s$ core state.¹² As such, data is almost exclusively restricted to oxides of $5d$ metals where well-resolved pre-edge peaks inherent to tetrahedral moieties are distinguished from the featureless rising-edge for octahedral centers.^{18,19}

The application of L_{1-} edge XAS to molecular systems with third row metals has yet to be undertaken. Its use hinges upon the availability of data for a well-defined set of reference compounds whose oxidation states are both relatively unambiguous and span a complete range of chemically accessible redox levels (Chart 1). Recent work^{20–22} in our laboratory with organometallic tungsten complexes bearing redox-active 1,2-dimethylethene-1,2-dithiolate, $(\text{mdt})^{2-}$, ligands has presented some challenge to finding a satisfactory description of their electronic structure and has motivated an effort to develop a small library of XAS reference compounds

that would have some general utility for the assessment of Z_{eff} and thereby assist a description of electronic structure in ambiguous situations (Chart 1). Accompanying these reference compounds are L-edge XAS data we have collected over a period of years on a broad set of mono-, bis-, and tris(dithiolene) complexes of tungsten. Here, we derive the estimated branching ratio (EBR), defined as the relative intensity of the L_{3-} edge over the sum of the L_{3-} and L_{2-} edges, $[I(L_{3-})/I(L_{3-} + L_{2-})]$,^{23–26} and the L_{1-} rising-edge energy for each reference compound. These two metrics are used to gauge whether the level of detail provided by L-edge XAS is sufficient to make an assessment of Z_{eff} across a range of tungsten dithiolene compounds. As X-ray absorption spectroscopy continues to grow as a method to complement the insights obtained by other physical methods, we shall illustrate that appropriately calibrated L-edge XAS for third row metals is a useful and informative alternative to the inaccessible K-edge for these heavy elements.

EXPERIMENTAL SECTION

Literature procedures were employed for the syntheses of $[\text{W}^{\text{II}}\text{Cl}_2(\text{PMePh}_2)_4]$,²⁷ $[\text{W}^{\text{II}}\text{Cl}_2(\text{dppe})_2]$ ($\text{dppe} = 1,2$ -bis-(diphenylphosphino)ethane),²⁸ $[\text{W}^{\text{IV}}\text{Cl}_4(\text{PMePh}_2)_2]$,²⁹ $[\text{W}^{\text{V}}(\text{NPh})\text{Cl}_3(\text{PMe}_3)_2]$,³⁰ $[\text{W}^{\text{VI}}(\text{xylylene})_3]$,³¹ $[\text{W}^0(\text{Me}_2\text{pipdt})(\text{CO})_4]$ ($\text{Me}_2\text{pipdt} = 1,4$ -dimethylpiperazine-2,3-dithione),²¹ $[\text{W}^{\text{II}}(\text{mdt})(\text{CO})_4]$,^{20,21} $[\text{W}^{\text{II}}(\text{mdt})(\text{CO})_2(\text{PMe}_3)_2]$,²⁰ $[\text{W}^{\text{IV}}(\text{mdt})_2(\text{CO})_2]$,^{20,32} $[\text{W}^{\text{IV}}(\text{mdt})_2(\text{CO})(\text{PMe}_3)]$,²⁰ $[\text{W}^{\text{IV}}(\text{mdt})_2(\text{PMe}_3)_2]$,²⁰ $[\text{W}^{\text{IV}}(\text{mdt})_2(\text{CN}^t\text{Bu})_2]$,²² $[\text{NEt}_4]_2[\text{W}^{\text{IV}}(\text{mdt})_2(\text{CN})_2]$,²² $[\text{Ph}_4\text{P}]_2[\text{W}^{\text{IV}}(\text{mnt})_3]$ ($\text{mnt}^{2-} = \text{maleonitriledithiolate}$),³³ $[\text{PPh}_4]_2[\text{W}^{\text{IV}}(\text{bdt})_3]$ ($\text{bdt}^{2-} = \text{benzene-1,2-dithiolate}$),³⁴ $[\text{NEt}_4]_2[\text{W}^{\text{IV}}(\text{bdt})_3]$,³⁴ $[\text{Et}_4\text{N}]_2[\text{W}^{\text{IV}}(\text{mdt})_3]$,³⁵ $[\text{PPh}_4][\text{W}^{\text{V}}(\text{bdt})_3]$,³⁴ $[\text{Et}_4\text{N}][\text{W}^{\text{V}}(\text{mdt})_3]$,³⁵ $[\text{W}^{\text{VI}}(\text{bdt})_3]$,³⁶ $[\text{W}^{\text{VI}}(\text{mdt})_3]$,²⁰ $[\text{W}^{\text{VI}}(\text{pdt})_3]$ ($\text{pdt}^{2-} = 1,2$ -diphenylethene-1,2-dithiolate).³⁶ Commercial sources of $[\text{W}^0(\text{CO})_6]$ and $[\text{W}^{\text{VI}}\text{Cl}_6]$ were purified by vacuum sublimation before use.

Syntheses. $[\text{W}^{\text{II}}\text{Cl}_2(\text{dppe})_2][\text{PF}_6]$. A 100 mL Schlenk flask with a stir bar was charged with a sample of $[\text{W}^{\text{II}}\text{Cl}_2(\text{dppe})_2]$ (0.397 g, 0.378 mmol) and 25 mL of dry dichloromethane. In a separate 50 mL Schlenk flask, a crystalline sample of $[\text{FeCp}_2][\text{PF}_6]$ (0.125 g, 0.378 mmol) was dissolved in 15 mL of dichloromethane. This solution was added to the yellow-orange solution of $[\text{W}^{\text{II}}\text{Cl}_2(\text{dppe})_2]$. Over the first 15 min, a light green color developed in the solution; stirring was continued for another 2 h. The solvent was removed under reduced

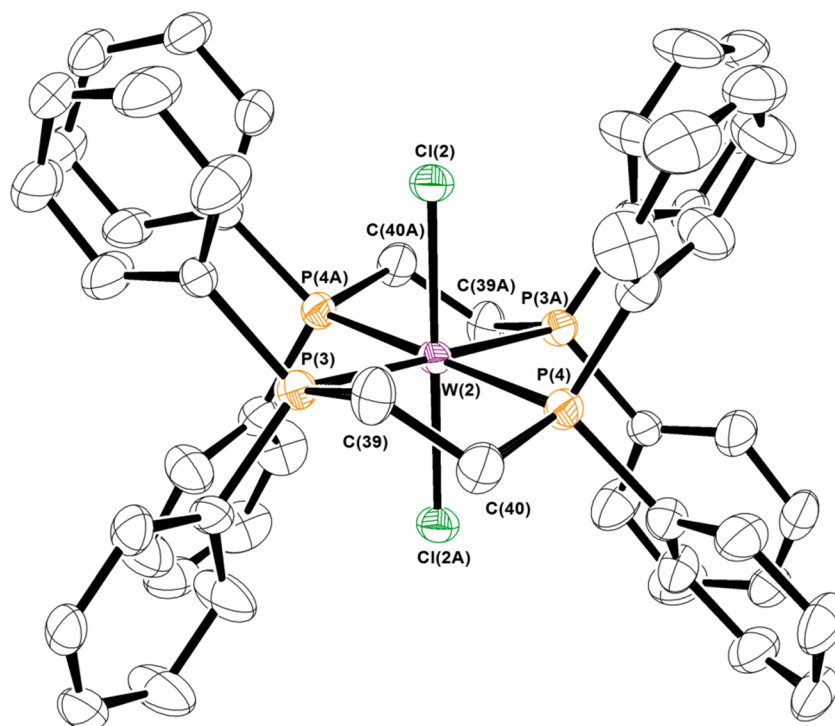


Figure 1. Thermal ellipsoid plot at the 50% probability level of one of the cations in the unit cell of crystals of $[W^{III}Cl_2(dppe)_2][PF_6]$. Hydrogen atoms are omitted for clarity.

pressure, affording a light greenish solid, which then was washed with Et_2O until the washings were colorless. After drying of the crude product overnight, orange crystals of $[W^{III}Cl_2(dppe)_2][PF_6]$ were obtained by the diffusion of Et_2O or pentane vapor into a concentrated 1,2-dichloroethane solution. Yield: 0.322 g (71%). Anal. Calcd for $C_{52}Cl_2F_6H_{48}P_5W$: C, 52.20; H, 4.04; P, 12.94. Found: C, 51.95; H, 4.28; P, 12.72%.

X-ray Crystallographic Data Collection and Structure Refinement. Diffraction-quality, yellow-orange block-shaped crystals of $[W^{III}Cl_2(dppe)_2][PF_6]$ were obtained by slow diffusion of Et_2O vapor into a 1,2-dichloroethane solution, while orange block-shaped crystals of $[W^{VI}(xylylene)_3]$ deposited from a hexanes solution upon standing. Crystals were coated with paratone oil and mounted on the end of a nylon loop attached to the end of a goniometer. Data were collected with a Bruker Smart APEX CCD diffractometer equipped with a Kryoflex attachment supplying a nitrogen stream at 100 K. Full spheres of data were obtained by the collection of three sets of 400 frames in ω ($0.5^\circ/\text{scan}$), collected at $\varphi = 0.00, 90.00,$ and 180.00° followed by two sets of 800 frames in φ ($0.45^\circ/\text{scan}$) collected with ω constant at -30.00 and 210.00° . Frame times of 15 and 10 s were used for $[W^{III}Cl_2(dppe)_2][PF_6]$ and $[W^{VI}(xylylene)_3]$, respectively. Data were collected under control of the APEX2 software package.³⁷ Raw data were reduced to F^2 values using the SAINT³⁸ software, and a global refinement of the unit cell parameters was performed using ~ 9900 selected reflections from the full data sets. Data were corrected for absorption on the basis of multiple measurements of symmetry-equivalent reflections with the use of SADABS.³⁹ Structure solutions were obtained by Patterson methods using SHELXS,⁴⁰ while refinements were accomplished by full-matrix least-squares procedures using SHELXL.⁴¹ Both the SHELXS and SHELXL programs are incorporated into the SHELXTL software suite.⁴²

The asymmetric unit of the unit cell for $[W^{III}Cl_2(dppe)_2][PF_6]$ was found to be composed of two independent $[PF_6]^{1-}$ anions, one of which was disordered, and four independent half $[W^{III}Cl_2(dppe)_2]^{1+}$ cations residing on inversion centers. Because of the imposed centrosymmetry, three of the four independent half-cations showed disorder to varying extents in the diphosphine ligand. This disorder was modeled by treating the affected phenyl groups as rigid entities and restraining equivalent metrical parameters to be approximately

equal. A similar treatment was accorded the disordered $[PF_6]^{1-}$ anion. In the final stages of refinement, hydrogen atoms were added in calculated positions and included as riding contributions with isotropic displacement parameters 1.2–1.5 times those of the carbon atoms to which they were attached. The thermal ellipsoid plot in Figure 1 and in the supplementary data were created with the use of XP, which also is part of the SHELXTL package.⁴² Final unit cell data and refinement statistics are collected in Table 1.

X-ray Absorption Spectroscopy. XAS data were measured at the Stanford Synchrotron Radiation Lightsource (SSRL) with the SPEAR storage ring containing 200–300 mA at 3.0 GeV. Tungsten L-edge spectra were collected on beamline 7–3 operating with a wiggler field of 2 T. A Si(220) double-crystal monochromator was used. Beamline 7–3 is equipped with a rhodium-coated vertical collimating mirror upstream of the monochromator and a downstream bent-cylindrical focusing mirror (also rhodium-coated). Incident and transmitted X-ray intensities were monitored using nitrogen-filled ionization chambers. Data were measured in transmittance mode, and samples were maintained at 10 K using an Oxford Instruments CF1208 continuous flow liquid helium cryostat. Internal energy calibrations were performed by simultaneous measurement of the W reference foil placed between the second and third ionization chamber with inflection points assigned as 12 100, 11 544, and 10 207 eV for the L_1 , L_2 , and L_3 -edges, respectively. Data were processed by fitting a second-order polynomial to the pre-edge region and subtracting this background from the entire spectrum.⁴³ A three-region cubic spline was used to model the smooth background above the edge. The data were normalized by subtracting the spline and normalizing the postedge to 1.0. Fits to the L_2 - and L_3 -edges modeled by pseudo-Voigt lines were carried out using the program EDG_FIT⁴³ with a fixed 1:1 ratio of Lorentzian to Gaussian contributions.

Other Physical Methods. Variable-temperature (2–290 K) magnetic susceptibility data were recorded in 1 T external field using an MPMS Quantum Design SQUID magnetometer. The experimental data were corrected for underlying diamagnetism using tabulated Pascal's constants. S- and X-band fluid solution spectra were collected using a Bruker EMX Micro spectrometer, and frozen solution spectra were obtained using a Bruker E580 spectrometer. Simulations

Table 1. Crystallographic Data for [W^{III}Cl₂(dppe)₂][PF₆] and [W^{VI}(xylylene)₃]

| | | |
|--|---|-----------------------------------|
| chem formula | C ₅₂ H ₄₈ Cl ₂ F ₆ P ₅ W | C ₂₄ H ₂₄ W |
| fw | 1196.50 | 496.28 |
| crystal system | triclinic | triclinic |
| space group | P $\bar{1}$ | P $\bar{1}$ |
| color/shape | orange/slab | orange/block |
| <i>a</i> , Å | 13.2421(8) | 6.8375(4) |
| <i>b</i> , Å | 19.247(1) | 11.1829(6) |
| <i>c</i> , Å | 19.638(1) | 11.8409(6) |
| α , deg | 100.3790(1) | 105.100(1) |
| β , deg | 98.9950(1) | 92.924(1) |
| γ , deg | 90.9070(1) | 90.392(1) |
| <i>V</i> , Å ³ | 4857.7(5) | 872.80(8) |
| <i>Z</i> | 4 | 2 |
| <i>T</i> , K | 100(2) | 100(2) |
| ρ calcd, g cm ⁻³ | 1.636 | 1.888 |
| reflns collected/2 Θ _{max} | 85 478/56.00 | 15 457/56.50 |
| unique reflns/ <i>I</i> > 2 σ (<i>I</i>) | 23 262/16 625 | 4267/4102 |
| parameters/restraints | 1130/220 | 274/0 |
| λ , Å/ μ (K α), mm ⁻¹ | 0.710 73/2.713 | 0.710 73/6.620 |
| GoF ^a | 1.055 | 1.058 |
| <i>R</i> ¹ / <i>b</i> , ^c / <i>wR</i> ² / <i>c</i> , ^d | 0.0453/0.0986 | 0.147/0.0332 |
| residual density, e Å ⁻³ | 3.58/−3.10 | 0.773/−0.579 |

^aGoF = $\{\sum[w(F_0^2 - F_c^2)^2]/(n - p)\}^{1/2}$, where *n* = number of reflections and *p* is the total number of parameters refined. ^b*R*¹ = $\sum||F_0| - |F_c||/\sum|F_0|$. ^c*R* indices for data cut off at *I* > 2 σ (*I*). ^d*wR*² = $\{\sum[w(F_0^2 - F_c^2)^2]/\sum[w(F_0^2)^2]\}^{1/2}$, where $w = 1/[\sigma^2(F_0^2) + (aP)^2 + bP]$, $P = (F_0^2 + 2F_c^2)/3$.

were performed using the Xsophe (Bruker Biospin GmbH) suite.⁴⁴ The elemental analysis was performed by Midwest Microlab, LLC, of Indianapolis, IN.

Calculations. All calculations were performed using the Gaussian 09 package.⁴⁵ Geometry optimizations and Kohn–Sham orbitals were calculated at Perdew–Burke–Ernzerhof (PBE)⁴⁶ (for exchange and correlation) and B3LYP⁴⁷ levels of density functional theory (DFT). The validity of all structures was confirmed by the absence of imaginary frequencies. The basis set chosen for all main group elements, except hydrogen, is the 6-31G(d,p). For tungsten a double- ζ (DZ) basis set with an effective electron core potential (LANL2DZ ECP) was implemented,⁴⁸ and a Gaussian split valence (SV) basis set was used for the hydrogen atoms.⁴⁹ The Mulliken population and atomic orbital composition analyses were calculated via a fragmentation approach using QMForge,⁵⁰ and all molecular orbital images were rendered using Chemcraft⁵¹ and Jmol.⁵²

RESULTS AND DISCUSSION

Reference Compounds. Several criteria governed the choice of compounds for this tungsten L-edge XAS study: (1) Strict adherence to coordination number of six at tungsten. (2) Avoidance of any ligand with potential redox activity. (3) Preference for compounds with a minimal number of strongly perturbing π -acceptor or π -donor ligands. Compounds with varying numbers of chloride and phosphine ligands meet these conditions and are useful as reference compounds because they are readily accessible synthetically. To that end, we assembled a series comprising [W⁰(PMe₃)₆],⁵³ [W^{II}Cl₂(PMePh₂)₄],²⁷ [W^{III}Cl₂(dppe)₂][PF₆], [W^{IV}Cl₄(PMePh₂)₂],²⁹ [W^V(NPh)-Cl₃(PMe₃)₂],³⁰ and [W^{VI}Cl₆], as they cover the full range of formal oxidation states at tungsten (Chart 1). The metal atom in [W⁰(PMe₃)₆] is as pure a zerovalent tungsten as might be conceptualized and thus is very useful as a spectroscopic benchmark compound. It is deemed superior to [W⁰(CO)₆],

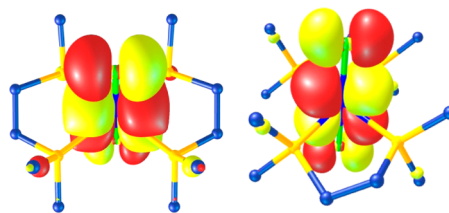
which despite its classification as zerovalent tungsten, experiences significant metal-to-ligand charge transfer via π -backbonding.³ A switch of the number and positions of the chloride and phosphine ligands relates [W^{II}Cl₂(PMePh₂)₄] and [W^{IV}Cl₄(PMePh₂)₂]. The ¹H and ³¹P NMR chemical shifts respond to the paramagnetism of the compounds.²⁸ The W^{II} complex possesses a (t_{2g})⁴ electron configuration, although a spin ground state cannot be applied given the large spin–orbit contribution from the 5d metal. A similar description is pertinent to [W^{IV}Cl₄(PMePh₂)₂], which exhibits a nonzero magnetic moment due to mixing of paramagnetic excited states by spin–orbit coupling,⁵⁴ a situation routinely encountered for octahedral d² species.^{55,56} The W^{III} standard, [W^{III}Cl₂(dppe)₂][PF₆], with fewer π -donor ligands, was favored over [CoCp₂]-[W^{III}Cl₄(PMePh₂)₂].⁵⁷ Although previously generated in a reaction between [W⁰(N₂)₂(dppe)₂] and CH₂Cl₂ in the presence of HFeCo₃(CO)₁₂,⁵⁸ we describe a more direct and better-yielding synthesis via chemical oxidation of [W^{II}Cl₂(dppe)₂] by [FeCp₂][PF₆]. Its constitution was confirmed by X-ray crystallographic analysis (Figure 1); the structure is highly similar to that reported with a [BF₄][−] counterion.⁵⁸ Table 2 contrasts the salient metric parameters

Table 2. Comparison of Averaged Bond Distances (Å) and Angles (deg)

| <i>a</i> | [W ^{II} Cl ₂ (dppe) ₂] ^b | [W ^{III} Cl ₂ (dppe) ₂][PF ₆] |
|--------------------|---|---|
| W–Cl | 2.4228(8) | 2.3207(6) |
| W–P | 2.5008(7) | 2.5518(5) ^c |
| Cl–W–Cl | 173.65(4) | 180 |
| P–W–P ^d | 79.14(3) | 78.27(2) ^c |

^aValues are averaged, where possible, with uncertainty propagations determined according to the general formula for uncertainty in a function of several variables detailed in ref 60. ^bData from ref 59. ^cValues involving the disordered dppe ligand were excluded from the averages. ^dLigand bite angle.

with the precursor, [W^{II}Cl₂(dppe)₂],⁵⁹ and shows a noticeable shortening of the W–Cl distances commensurate with a one-electron depopulation of the degenerate d_{xz,yz} highest occupied molecular orbital (HOMO), which are π -antibonding with the Cl[−] ligands (Figure 2). The breadth of the WP₄ plane expands slightly upon oxidation of the metal ion.

**Figure 2.** Degenerate SOMO comprising the ground state in [W^{III}Cl₂(dppe)₂]⁺ with ~72% d_{xz,yz} character.

The temperature dependence of the magnetic moment of this complex was examined by SQUID magnetometry (Figure S1). The room-temperature effective magnetic moment of 1.2 μ_B is consistent with a low-spin d³ ion (*S* = 1/2) affected by the sizable tungsten spin–orbit coupling constant that reduces the average *g* value to 1.35. Similar room-temperature magnetic moments have been reported for analogous complexes.^{56,58} The axial electron paramagnetic resonance (EPR) spectrum,

measured at 20 K (Figure 3), was simulated using $g = (1.359, 1.266, 1.006)$. The shift to high field leads to significant line

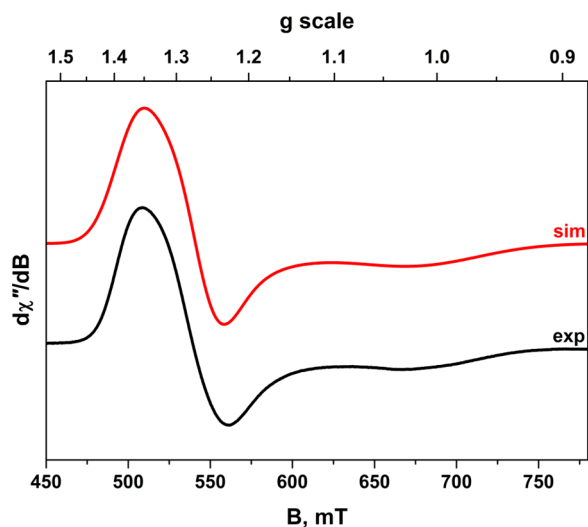


Figure 3. X-band EPR spectrum of $[\text{W}^{\text{III}}\text{Cl}_2(\text{dppe})_2][\text{PF}_6]$ recorded in $\text{CH}_2\text{Cl}_2/\text{THF}$ solution at 20 K (experimental conditions: frequency, 9.6310 GHz; power 0.63 mW; modulation, 0.7 mT). Experimental data are depicted by the black line; the simulation is in red.

broadening that obscures resolution of the ^{183}W ($I = 1/2$, 14.3% abundant) and ^{31}P ($I = 1/2$, 100% abundant) active nuclei. The substantial deviation away from 2.0023 is consistent with a $(d_{xy})^2(d_{xz},d_{yz})^1$ electron configuration.⁶¹ The extent of the g shift, the fast electronic relaxation and low magnetic moment are hallmarks of a near degenerate ground state (2E_g) where orbital angular momentum is partially unquenched. To our knowledge, these are the lowest g values reported for a low-spin W^{III} complex.⁶² $[\text{W}^{\text{V}}(\text{NPh})\text{Cl}_3(\text{PMe}_3)_2]$ is a deliberate choice for W^{V} representative rather than a more obvious $[\text{W}^{\text{V}}\text{OCl}_3(\text{PR}_3)_2]$ compound, which could complicate the spectroscopy should it cocrystallize with the related chloro species $[\text{W}^{\text{IV}}\text{Cl}_4(\text{PR}_3)_2]$.⁶³ The effect of substituting a terminal oxo for an imido ligand was explored by EPR spectroscopy. The simulation of chilled solution (190 K) spectra recorded at S- and X-band frequencies yielded $g_{\text{iso}} = 1.903$ (Figures S2 and S4), a value consistent with an effective magnetic moment lower than the spin-only value.^{30,64} In contrast, $[\text{W}^{\text{V}}\text{OCl}_3(\text{PPh}_3)_2]$ exhibited $g = 1.791$,⁶⁵ which identifies greater covalency within the $\{\text{W}=\text{NPh}\}^{3+}$ unit. This conclusion is supported by a lower tungsten $4f_{7/2}$ binding energy in $[\text{W}^{\text{VI}}(\text{NPh})\text{Cl}_4]_2$ than in $[\text{W}^{\text{VI}}\text{OCl}_4]_x$.⁶⁶ Hyperfine coupling to two equivalent ^{31}P ($26 \times 10^{-4} \text{ cm}^{-1}$) nuclei and the ^{183}W ($60 \times 10^{-4} \text{ cm}^{-1}$) nucleus are observed with improved visibility at S-band (Figures S2 and S3).

$L_{3,2}$ -Edges. The L_{3-} and L_{2-} edges exhibit prominent features referred to as white lines, identified as $2p \rightarrow ns$ and $2p \rightarrow nd$ electronic transitions, which are both dipole-allowed. These edges are separated by $2p$ spin-orbit coupling; the L_{2-} edge for tungsten lies about 1337 eV above the L_{3-} edge. Specifically, the L_{3-} edge represents transitions from $2p_{3/2}$ to both $5d_{3/2}$ and $5d_{5/2}$ states, whereas the L_{2-} edge pertains to excitations from $2p_{1/2}$ solely to the $5d_{3/2}$ state. Changes in the absorption are observed across the series and reflect variation in the number of vacancies in the $5d$ orbitals.^{67,68} The L_{3-} edge is twice as intense as the L_{2-} , though there is deviation from this statistical ratio in the W L -edge study, as previously noted for $5d$ metals.⁶⁹ Excitations to unoccupied s levels are considerably weaker and

can be neglected.⁷⁰ The L_{3-} and L_{2-} edge spectra recorded for each reference compound are presented in Figure 4. Each spectrum has been normalized to the step in the continuum across the absorption edge, and white-line energies are posted in Table 3.

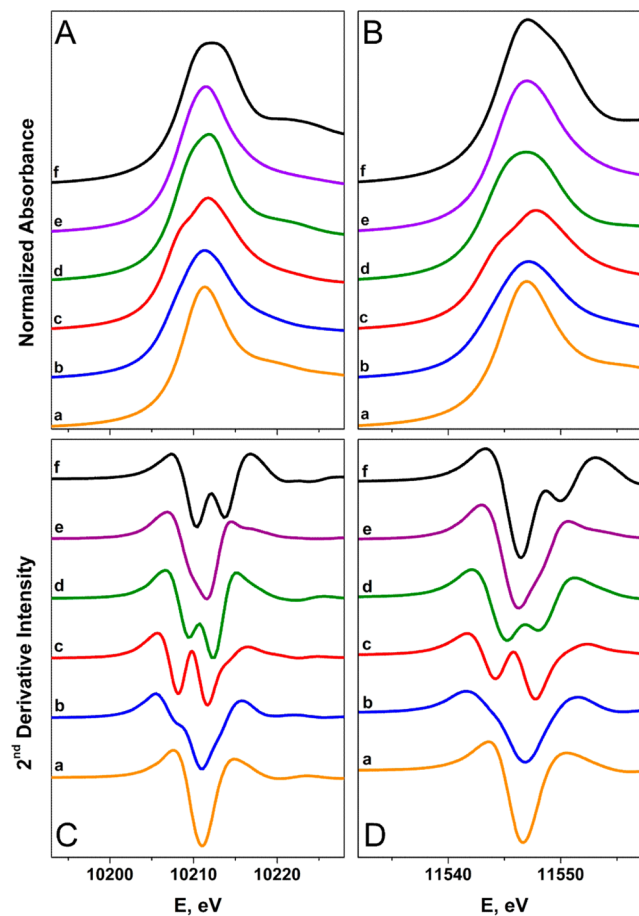


Figure 4. Comparison of the normalized W L_{3-} (panel A) and L_{2-} edge (panel B) and their FFT-smoothed second derivative spectra (panels C and D) of (a) $[\text{W}^0(\text{PMe}_3)_6]$, (b) $[\text{W}^{\text{III}}\text{Cl}_2(\text{PMePh}_2)_4]$, (c) $[\text{W}^{\text{III}}\text{Cl}_2(\text{dppe})_2][\text{PF}_6]$, (d) $[\text{W}^{\text{IV}}\text{Cl}_4(\text{PMePh}_2)_2]$, (e) $[\text{W}^{\text{V}}(\text{NPh})\text{Cl}_3(\text{PMe}_3)_2]$, and (f) $[\text{W}^{\text{VI}}\text{Cl}_6]$.

The white-line energies are invariant at the L_{2-} edge across the series, with only a subtle shift to higher values at the L_{3-} edge for more oxidized W ions. With the exception of $[\text{W}^0(\text{PMe}_3)_6]$ and its $(t_{2g})^6$ electron configuration, each

Table 3. L -Edge Energies for Reference Compounds

| | L_{3-}^a | L_{2-}^a | L_1 | |
|--|------------|------------|----------|--------------------------|
| | | | pre-edge | rising-edge ^b |
| W^c | 10 210.1 | 11 546.3 | | 12 100.0 |
| $[\text{W}^0(\text{PMe}_3)_6]$ | 10 211.3 | 11 547.0 | 12 091.6 | 12 095.2 |
| $[\text{W}^{\text{III}}\text{Cl}_2(\text{PMePh}_2)_4]$ | 10 211.3 | 11 547.1 | 12 092.3 | 12 096.0 |
| $[\text{W}^{\text{III}}\text{Cl}_2(\text{dppe})_2][\text{PF}_6]$ | 10 211.8 | 11 547.9 | 12 092.4 | 12 096.5 |
| $[\text{W}^{\text{IV}}\text{Cl}_4(\text{PMePh}_2)_2]$ | 10 211.8 | 11 547.0 | 12 090.4 | 12 097.6 |
| $[\text{W}^{\text{V}}(\text{NPh})\text{Cl}_3(\text{PMe}_3)_2]$ | 10 211.5 | 11 547.0 | 12 093.1 | 12 097.8 |
| $[\text{W}^{\text{VI}}\text{Cl}_6]$ | 10 212.2 | 11 547.1 | 12 092.1 | 12 099.2 |

^aWhite-line peak maximum. ^bEnergy of first inflection point determined from first derivative of the spectrum. ^cTungsten reference foil.

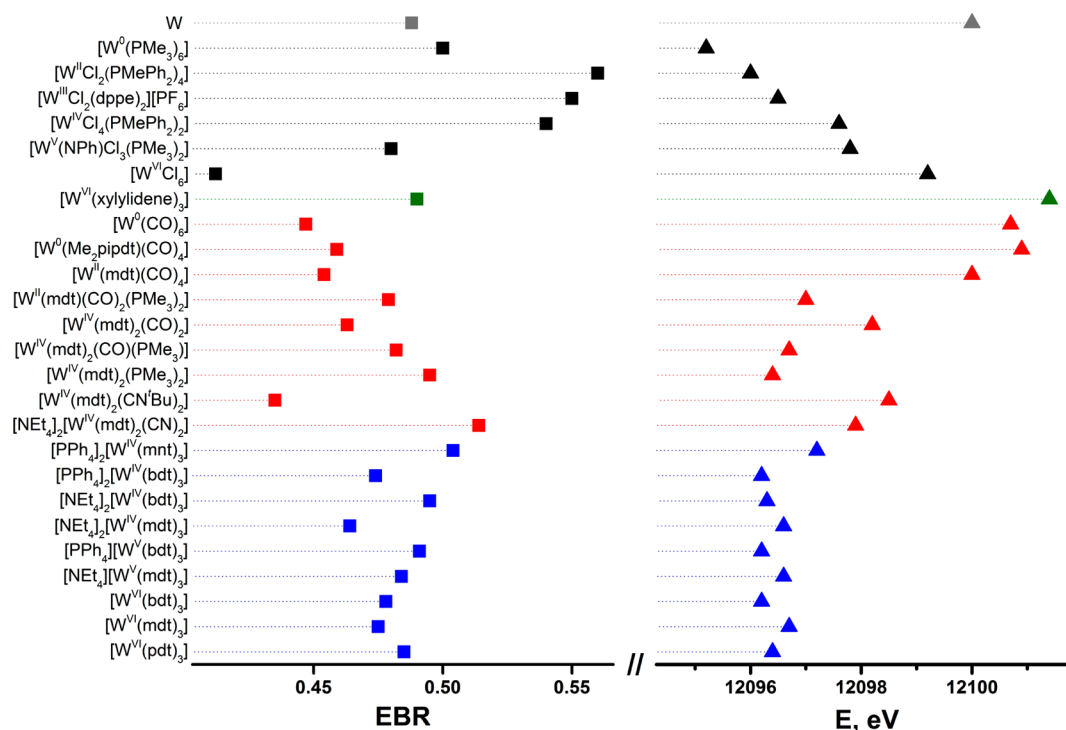


Figure 5. Comparison of EBR (left) and L_1 rising-edge energies (right).

spectrum displays splittings of the white-line peaks due to the ligand-field, which are the same at both edges. Ligand-field splittings, more readily observed in the second derivative plot, range from 1.1 to 3.6 eV (Figure 4) and compare well with those computed by DFT (Table S1). We note that the decreased shielding due to the creation of a core-hole will affect the acceptor orbitals differently, which is an effect not included in the calculations. Assuming near-octahedral geometry for the complexes, the lower energy peak is assigned as a transition to a vacancy in the t_{2g} orbitals, while the higher energy peak is attributed to an excitation to the vacant e_g set. The intensity of the first peak increases relative to the second peak for higher oxidation states of tungsten in a fashion commensurate with the greater number of electron vacancies in the t_{2g} levels. The lower-valent compounds (0 to +IV) of this series are yellow or orange-yellow in color and devoid of low-energy ligand-field transitions consistent with large Δ_{OCT} provided by phosphine ligands. The multiply bonded imido group in $[W^V(NPh)Cl_3(PMe_3)_2]$ leads to a significant departure from an octahedral ligand-field, as reflected by the smaller splitting of the white-line peak. On the whole, this splitting makes precise determination of the peak energy problematic (Table 3).

The white-line intensity at both edges trends well with the assigned oxidation state of tungsten. The peak area was estimated by a simple curve-fitting method, where Gaussian broadened Lorentz functions were used to simulate the white-line peaks after removal of the edge continuum jump modeled by an arctan function (see Supporting Information).^{26,67,71} Because the optimum energy position for the arctan function is somewhat uncertain, they have been fixed to the intersection with the rising-edge of the white-line. The data plotted in Figure S6 show a monotonic increase in the intensity for the L_3 -edge with increasing oxidation state.^{71–73} Departure from linearity is ascribed to variation in the peak profile of the tail of the white line above the ionization threshold due to transitions

to quasi-bound states leading to a deviation of the peak from a pseudo-Voigt line shape.^{69,72} Also $2p-5d$ multiplet effects will affect the spectral shape and peak area.⁷⁴ A similar increase is noted for the L_2 -edge (Figure S6), though $[W^0(PMe_3)_6]$ departs significantly from expectation without an obvious explanation. This leads to a smaller EBR than anticipated based on its L_1 rising-edge energy.

The branching ratio has been used as a metric of metal oxidation state in discrete molecules and extended solids by XAS^{6,23} and electron energy-loss spectroscopy.^{24,75} The EBR has been determined for each reference compound and plotted in Figure 5.^{23–26} Aside from the low value for $[W^0(PMe_3)_6]$, there is a decrease in the EBR as the series of reference compounds is traversed from low to high formal oxidation state. In an effort to assess the validity of assigning a physical oxidation state based on EBR, these values are compared to those obtained for an ensemble of W coordination compounds. The impact of potent π -accepting CO ligands is clearly observed for $[W^0(CO)_6]$, where the computed EBR of 0.45 is indicative of a Z_{eff} more akin to W^V than to W^0 .

At the other end of the scale, $[W^VI(xylylidene)_3]$ has a larger EBR than the corresponding standard, $[W^VICl_6]$. This suggests a somewhat more reduced central ion in $[W^VI(xylylidene)_3]$, which stems from its preference for a C_{3h} symmetric paddle wheel structure as opposed to the octahedron in $[W^VICl_6]$.³¹ Hexamethyltungsten, as an unambiguous W^VI species, is the more ideal candidate for study inasmuch as the charge at tungsten cannot be alleviated by any π donation, there being no lone electron pairs. However, as the synthesis and handling of this compound are attended by some risk of explosion,⁷⁶ $[W^VI(xylylidene)_3]$ is a safer alternative for measurement. A survey of the frontier MOs in an optimized structure of $[W^VI(Me)_6]$ shows a lowest unoccupied molecular orbital (LUMO) that is essentially pure d_z^2 (Figure S42). A pair of deeper lying MOs (HOMO-1,2, e' ; HOMO-3,4, e'') have

moderate metal d character (32% and 34%, respectively), but the bonding is carbon p \rightarrow tungsten d σ donating in character. The metal–ligand bonding description in $[\text{W}^{\text{VI}}(\text{xylylene})_3]$ is qualitatively similar to that in $[\text{W}^{\text{VI}}(\text{Me})_6]$ but is rendered more complicated largely by its lower symmetry (C_{3h}).³¹ The symmetry is produced by a substantial fold of all three xylylene ligands along each intraligand $\text{CH}_2\cdots\text{CH}_2$ vector, which orients the unit almost “side-on.” The methylene carbon atoms are positioned 2.2137(8) Å from the tungsten ion, while their corresponding aromatic carbons are 2.4817(8) Å away. Here the average dihedral angle between the $\text{W}(\text{CH}_2)_2$ and xylylene C_8 planes is 83.8(3)° (Figure S62). This angle is considerably smaller than it is for xylylene complexes with other early transition metals⁷⁷ and is a means by which the ligand further supplements the electron-deficient metal ion. The consequence of this structural distortion is mixing of the ligand-based HOMO with the metal-based LUMO, which transform to the same representation in C_{3h} point symmetry.^{78,79} The mixing deposits metal character into the HOMO (now HOMO-3, Figure S41) and ligand character into the LUMO with overall stabilization of the molecule. Therefore, the EBR is closer to that of $[\text{W}^{\text{V}}(\text{NPh})\text{Cl}_3(\text{PMe}_3)_2]$ than that of $[\text{W}^{\text{VI}}\text{Cl}_6]$ because the metal content of the LUMO is reduced. Tungsten’s preference for high oxidation states establishes $[\text{W}^{\text{VI}}(\text{xylylene})_3]$ as much closer to the limiting description of W^{VI} with alkyl ligands than to a W^0 species with olefinic ligands.

L_3 - and L_2 -edge spectra for a series of tungsten mono-(dithiolene) and bis(dithiolene) compounds with various ancillary ligands (CO , PMe_3 , CN^tBu , CN^-) are collected and overlaid in Figure S8. The white-line energies listed in Table S2 show no discernible correlation with the formal oxidation state. As expected, the comparison of the white-line peak energy with the +II and +IV standards shows no correlation with the formal oxidation state at either the L_3 - or L_2 -edge (Figure S9), demonstrating the dominance of ligand-field over Z_{eff} for excitation from p states. The second series, comprising a set of tris(dithiolene) complexes whose formal oxidation states span +IV to +VI, shows no variation in their white-line energies when plotted against their relevant standards (Figures S10 and S11). The L_2 -edge energies are the same within experimental error, whereas all L_3 -edge values are lower than the three standards. This reflects the similar spectroscopic oxidation state for all tris(dithiolene) complexes as revealed by EPR and S K-edge XAS,^{79,80} in that each member of the electron transfer series is related by predominantly ligand-centered redox steps.⁸¹ The electron-withdrawing effect of the cyano substituents in $[\text{PPh}_4]_2[\text{W}^{\text{IV}}(\text{mnt})_3]$ shifts the peak to an energy higher than for other dithiolene types, as has been observed repeatedly in S K-edge spectra.^{11,14,16,80}

L_1 -Edge. An overlay of the L_1 -edge spectra for six tungsten compounds spanning the formal oxidation state range of 0 to +VI is presented in Figure 6. As further emphasized with the rising-edge energies listed in Table 3, the spectra are qualitatively similar and ordered according to increasing formal oxidation state but are progressively shifted to higher energy by irregular energy increments of 0.2–1.4 eV. Moreover, an inverse correlation between the $L_{3,2}$ -edge EBR and L_1 rising-edge energy is achieved (Figure 5). The correspondence of the rising-edge energy order to the ordering by formal oxidation state is undoubtedly dependent upon juxtaposed members in the series being as alike as possible in the identity and nature of the ligand environment.

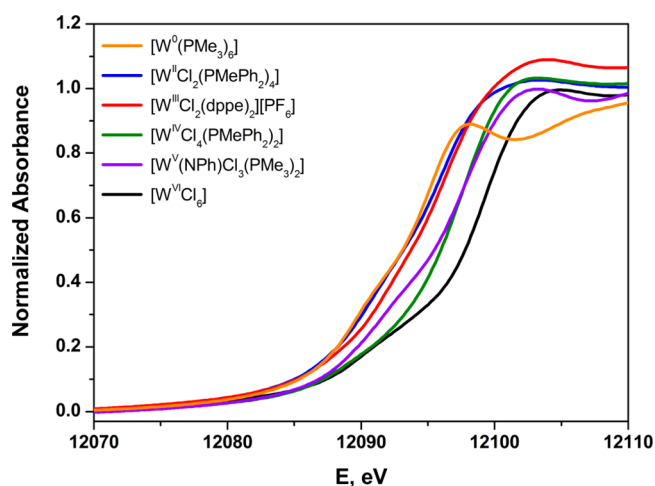


Figure 6. Overlay of the normalized W L_1 -edge X-ray absorption spectra of the reference compounds spanning formal oxidation states 0 \rightarrow +VI.

The comparison between $[\text{W}^{\text{II}}\text{Cl}_2(\text{PMePh}_2)_4]$ and $[\text{W}^{\text{III}}\text{Cl}_2(\text{dppe})_2]^{1+}$ is a noteworthy one, as the two compounds are essentially alike in coordination environment and differ only peripherally in the identity of the supporting phosphine ligand. The one-electron transfer that would oxidize $[\text{W}^{\text{II}}\text{Cl}_2(\text{PMePh}_2)_4]$ to $[\text{W}^{\text{III}}\text{Cl}_2(\text{PMePh}_2)_4]^{1+}$ or reduce $[\text{W}^{\text{III}}\text{Cl}_2(\text{dppe})_2]^{1+}$ to $[\text{W}^{\text{II}}\text{Cl}_2(\text{dppe})_2]$ involves a degenerate pair of MOs that is largely ($\sim 72\%$) tungsten d orbital ($d_{x^2-y^2}$) in character (Figure 2). The inference to which the foregoing leads is that a rising-edge energy change of ~ 0.5 eV might be generally anticipated for a redox process that is metal-based, or largely so, for a third row transition element. For a first row transition metal, a change of ~ 1.0 eV in the K-edge energy is generally associated with a metal-based redox process.¹¹ A somewhat smaller value is plausible for the L_1 -edge energy for a transition metal inasmuch as the 2s core-hole, being further from the nucleus, is conceivably less sensitive to Z_{eff} than a 1s core-hole.

The degree of the departure that can occur between formal and spectroscopic oxidation state is highlighted in spectacular fashion by the contrast between the L_1 -edge spectra for $[\text{W}^0(\text{PMe}_3)_6]$ and $[\text{W}^0(\text{CO})_6]$ (Figure 7). Gauged spectro-

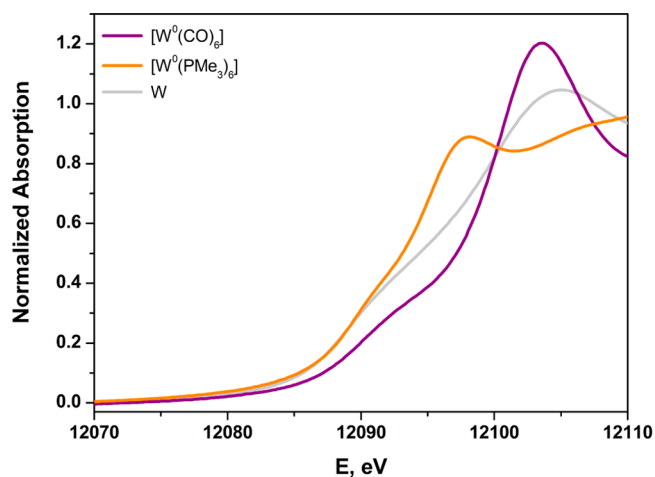


Figure 7. Overlay of the normalized W L_1 -edge X-ray absorption spectra of $[\text{W}^0(\text{PMe}_3)_6]$, $[\text{W}^0(\text{CO})_6]$, and W metal.

scopically by L_1 -edge XAS, $[W^0(CO)_6]$ bears more resemblance to W^V than it does to W^0 . This strongly perturbing effect of CO has been noted previously in a K-edge XAS study of technetium compounds that included $[Tc^{0_2}(CO)_{10}]$.⁸² The spectroscopic difference between $[W^0(CO)_6]$ and $[W^0(PMe_3)_6]$ accords qualitatively with observation: $[W^0(CO)_6]$ is a robust species and stable thermally in the open atmosphere for protracted times; $[W^0(PMe_3)_6]$ is extraordinarily reactive, capable of hard-to-effect C–C and C–H insertion reactions in heteroaromatic molecules,⁸³ and prone to cyclometalation to form $[W^{II}(PMe_3)_4(\eta\text{-}CH_2PMe_2\text{-}H)]$ following dissociation of PMe_3 .⁵³

Although encompassing a broad energy window for formally zerovalent tungsten compounds, $[W^0(PMe_3)_6]$ and $[W^0(CO)_6]$ are nevertheless useful for establishing limiting values for L_1 -edge energies. The former is particularly useful for the definition of a low-energy limit. All other common or conceivable zerovalent compounds would fill in a continuum between these two, in some cases with a plausible estimate of the L_1 -edge energy being possible. For example, $[W^0(CO)_3(\text{triphos})]$ (triphos = $CH_3C(CH_2PPh_2)_3$) would likely have an L_1 rising-edge at the intermediate value of $\sim 12\,098$ eV. We note here that an increasing number of CO ligands in a series of related compounds correlates more strongly with a positive shift of the rising-edge than does the formal oxidation state of the metal ion. This point is exemplified in the series $[W(\text{dithiolene})_n(CO)_{6-2n}]$ ($n = 1\text{--}3$) whose L_1 -edge spectra are compared in Figure S48. The lowest energy corresponds to the compound with the highest formal oxidation state and an absence of CO ligands.

The usefulness of the L_1 -edge XAS data from reference compounds such as the set in Figure 6 is demonstrated by their use as a backdrop for comparison with the L_1 -edge spectra of metallodithiolene compounds $[W^{IV}(\text{mdt})_2(CO)_2]$ and $[W^{IV}(\text{mdt})_2(CN)_2]^{2-}$ (Figure 8). The rising edges of these

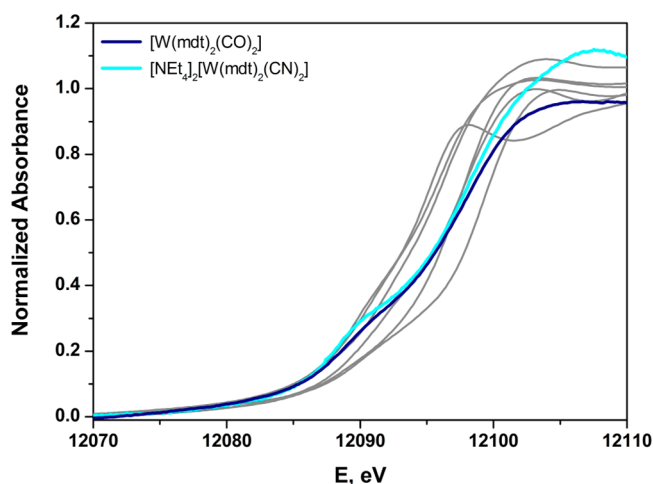


Figure 8. Normalized tungsten L_1 -edge X-ray absorption spectra of $[W^{IV}(\text{mdt})_2(CO)_2]$ and $[W^{IV}(\text{mdt})_2(CN)_2]^{2-}$ overlaid upon the L_1 spectra of the reference compounds in Figure 6.

two metallodithiolene species are similar to one another, suggesting similar Z_{eff} between the two, and roughly comparable to $[W^{IV}Cl_4(PMePh_2)_2]$ and $[W^VCl_3(NPh)(PMe_3)_2]$ (Figure 5). The pre-edge shoulder is also comparable, given the same trigonal prismatic geometry.^{20,22} This observation affirms a description of $[W^{IV}(\text{mdt})_2(CN)_2]^{2-}$

as a W^{IV} species, which is also the conclusion emerging from crystallographic data and a study of its frontier MO composition. The case of $[W^{IV}(\text{mdt})_2(CO)_2]$, however, is considerably more subtle. Sulfur K-edge XAS data and the S–C and C–C bonds within the dithiolene ligands for both $[W^{IV}(\text{mdt})_2(CO)_2]$ and $[W^{IV}(\text{mdt})_2(CN)_2]^{2-}$ indicate appreciably more oxidized sulfur in the former,²² which in the absence of W L -edge XAS data invite the conclusion that its tungsten ion is correspondingly more reduced. However, Figure 8 emphasizes that Z_{eff} in $[W^{IV}(\text{mdt})_2(CO)_2]$ is at least as high as it is in $[W^{IV}(\text{mdt})_2(CN)_2]^{2-}$. Examination of the frontier MOs for $[W^{IV}(\text{mdt})_2(CO)_2]$ shows not only effective W \rightarrow CO π -backbonding, which renders tungsten Z_{eff} relatively high, but also a key MO that is dithiolene π -donating and CO π -accepting via the same tungsten d orbital (Figure S49). This MO bears a strong analogy to *cis*- $W^{IV}O(CO)$ species in which oxo and carbonyl ligands interact in a synergistic way with the same tungsten d orbital as π donor and π acid (Figure S49).⁸⁴ Thus, tungsten L_1 -edge XAS complements the S K-edge data, which only provides the dithiolene sulfur contribution to the frontier MOs.

The position of the rising-edge is also a function of the M–L distances in the first coordination sphere of the central ion.⁸⁵ The distances of the non-dithiolene ancillary ligands from the tungsten ion in complexes of the type $[W^{IV}(\text{mdt})_2(X)(Y)]$ ($X = Y = CO, PMe_3, CN^tBu, CN^-$; $X = CO, Y = PMe_3$; Chart 1) trend as $CO < CN^tBu < CN^- < PMe_3$,²² which correlates directly with the L_1 rising-edge energy; the average W–S bond length remains unchanged across the series. Of course, π -backbonding is also enhanced by a short metal–ligand bond. The W–CO bond lengths in $[W^{IV}(\text{mdt})_2(CO)_2]$ are shorter than the W–CN bond lengths in $[W^{IV}(\text{mdt})_2(CN)_2]^{2-}$ by 0.114(6) Å and contribute to its visibly higher energy tungsten L_1 rising-edge (Figure 8). Substituting CO for PMe_3 shifts the rising-edge to lower energy by the removal of an effective π acceptor and concomitant lengthening of the metal–ligand bond with the larger phosphorus atom. The shift in the L_1 rising-edge corresponds directly to the energy of the first peak in the S K-edge spectrum.²²

An overlay of two formally W^{VI} compounds (Figure 9) similarly illustrates a difference of rising-edge energies, in this case 2.2 eV. While the separation between $[W^{VI}Cl_6]$ and

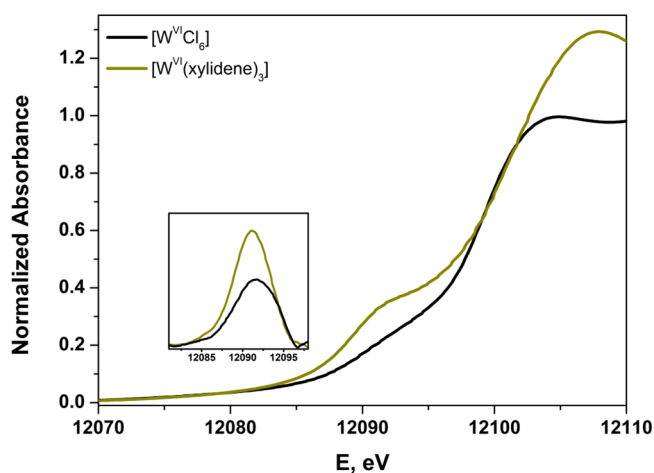


Figure 9. Overlay of the normalized W L_1 -edge X-ray absorption spectra of $[W^{VI}Cl_6]$ and $[W^{VI}(\text{xylidene})_3]$. Inset shows expansion of the pre-edge peak after subtraction of the rising-edge.

$[W^{VI}(\text{xylylene})_3]$ is less stark than that between $[W^0(\text{PMe}_3)_6]$ and $[W^0(\text{CO})_6]$, the order is opposite that given by the EBR. Although chloride is generally a modest π -donor ligand, the chloride lone pairs in $[W^{VI}\text{Cl}_6]$ are situated to form multiple Cl $p \rightarrow W d$ π -type interactions with each of the metal t_{2g} -type atomic orbitals. It is therefore unsurprising that that formal charge in $[W^{VI}\text{Cl}_6]$ is attenuated to the extent that its L_1 rising-edge is markedly lower than it is in $[W^{VI}(\text{xylylene})_3]$, where no ligand lone pairs are present on the atoms immediately coordinated to tungsten. The higher L_1 -edge energy for $[W^{VI}(\text{xylylene})_3]$ can also be ascribed to the shorter W–C distance of 2.2127(8) Å compared with an average W–Cl bond length of 2.276(2) Å in $[W^{VI}\text{Cl}_6]$.⁸⁶

The energy gap between the pre-edge and rising-edge diminishes with increasing metal–ligand bond length because the energy difference between empty metal d orbitals and filled ligand p orbitals decreases. The pre-edge transition for $[W^{VI}(\text{xylylene})_3]$ occurs at slightly lower energy and with appreciably greater intensity than that for $[W^{VI}\text{Cl}_6]$ (Figure 9 inset). Its noncentrosymmetric geometry endows the otherwise weak quadrupole-allowed $2s \rightarrow 5d$ pre-edge excitation with electric dipole character. For trigonal prismatic $[W^{VI}(\text{xylylene})_3]$, the pre-edge feature is defined as the $2s \rightarrow 5d_{xy,x^2-y^2}$ transition, where DFT computed 2.1% W 6p character is admixed into this e' level (LUMO+1 and LUMO+2, Figure S41). The pre-edge peak energy, evaluated after subtracting the rising-edge structure, shows the transition in $[W^{VI}(\text{xylylene})_3]$ is 0.4 eV lower in energy than it is for $[W^{VI}\text{Cl}_6]$. This difference is in keeping with the EBR analysis, which showed the former is slightly more reduced (vide supra).

The impact of three redox-active dithiolene ligands is most apparent comparing the L_1 -edge position of the neutral tris(dithiolene) complexes — $[W^{VI}(\text{bdt})_3]$, $[W^{VI}(\text{mdt})_3]$, and $[W^{VI}(\text{pdt})_3]$ — with the +VI standard, $[W^{VI}\text{Cl}_6]$ (Figure 5). The sulfur-donor species bring about a ~ 2.5 eV shift to lower energy commensurate with the now generally accepted notion that tungsten is not in its highest oxidation state.^{79,87} Each compound forms a three-membered electron transfer series,⁷⁹ where the rising-edge energy remains unresponsive to the successive addition of one electron to give monoanionic and dianionic species. For both the $[W(\text{mdt})_3]^{0/1-/-2-}$ and $[W(\text{bdt})_3]^{0/1-/-2-}$ series, the variation is ± 0.1 eV (Table S4), which approaches the precision limit of the experiment. As noted for the $L_{3,2}$ -edges, the electronically unique $[W^{IV}(\text{mnt})_3]^{2-}$ is 1 eV higher than the other dianions, as the degree of π donation is attenuated by the conjugated CN substituents. These data indicate that Z_{eff} is the same for each member of the series. Recently, we detailed a spectroscopic and theoretical analysis of EPR-active monoanionic tris(dithiolene) complexes of Mo and W.⁷⁹ For the Mo compounds there was an obvious difference between the aromatic and olefinic dithiolene ligand, with the former defined as a Mo^V central ion coordinated by three dianionic dithiolates, while the latter has a Mo^{IV} ion and an oxidized tris(dithiolene) ligand set. In the case of tungsten, the division is considerably more ambiguous, as the electronic structure of compounds with olefinic dithiolenes rests somewhere between the two extremes — W^{IV} and W^V — due to a second-order Jahn–Teller distortion as described for $[W^{VI}(\text{xylylene})_3]$ (vide supra).⁷⁹ Similarly, all neutral complexes adopt the C_{3h} symmetric paddle-wheel structure where the magnitude of the distortion defines the extent of mixing of metal and ligand orbitals, and therein the Z_{eff} of W. Therefore, the ability to apply an integer oxidation state to any tungsten

tris(dithiolene) complex is severely compromised in such strongly covalent systems, as reflected by the uniform rising-edge energies.

The conspicuous pre-edge shoulder in the L_1 -edge spectrum of each tris(dithiolene) compound is a spectroscopic marker for complex geometry. Neutral complexes are noted for their trigonal prismatic WS_6 polyhedron,^{20,32,79,88} as are the complex ions, $[W(\text{mdt})_3]^{1-/-2-}$, but with flat dithiolene ligands.^{35,79} Each member of this series displays an equally intense pre-edge peak (Figure S50 inset), similar to other trigonal prismatic tris(dithiolene) species.^{15,16} The calculated 6p content of the bound $5d_{xy,x^2-y^2}$ is uniform across the series (Table S4). The analogous $[W(\text{bdt})_3]^{0/1-/-2-}$ series shows marked differences in the pre-edge intensity (Figure S51 inset). The degree of p–d mixing increases with decreasing Θ (Table S4); the distorted octahedral $[W^V(\text{bdt})_3]^{1-}$ ($\Theta = 32.3^\circ$) has the weakest pre-edge peak.⁸⁹ Perhaps the most satisfying demonstration rests in the comparison of the two $[W^{IV}(\text{bdt})_3]^{2-}$ complexes with different counterions. Crystallization of this dianion with two tetraethylammonium cations yields a highly trigonal prismatic structure ($\Theta = 1.9^\circ$),⁹⁰ whose pre-edge intensity is noticeably reduced with tetraphenylarsonium counterions, as the structure is shifted to distorted octahedral ($\Theta = 23.0^\circ$).⁹¹ The effect of counterions on the trigonal twist angle has been elegantly detailed in a crystallographic study of $[M^{IV}(\text{bdtCl}_2)_3]^{2-}$ ($M = \text{Mo}, \text{W}$; $\text{bdtCl}_2^{2-} = 3,6\text{-dichlorobenzene-1,2-dithiolate}$),⁹² where those cocrystallized with tetraethylammonium cations are more trigonal prismatic.

CONCLUSIONS

The complete L-edge XAS spectra of a broad set of six-coordinate tungsten compounds have been collected, some compounds being selected for use as references and others because they bear redox-active dithiolene ligands and varying degrees of ambiguity. We are unaware of any comparable study for a 5d metal. For third row metals, for which K-edge XAS spectra are inaccessible, it is clear that L-edge XAS has sufficient spectral resolution to probe metal Z_{eff} when appropriately calibrated. For the reference series employed here, which maintains a rough similarity in ligand environment from one series member to the next, the EBR derived from the relative intensity of the $L_{3,2}$ -edges is anticorrelated with the L_1 rising-edge energy. As anticipated, the white-line peak energies at the L_3 - and L_2 -edges are independent of Z_{eff} . Our observations regarding the EBR and L_1 rising-edge energies demonstrate that these two metrics respond to changes in Z_{eff} for a 5d metal and that the differences are significant as the reference series is traversed from 0 to +VI.

Important contributors to the EBR and rising-edge energy are the metal–ligand bond distances and the π -acid/ π -donor character of the ligands, irrespective of the formal oxidation state classification. This point is underscored by $[W^{II}(\text{mdt})(\text{CO})_4]$ and $[W^{II}(\text{mdt})(\text{CO})_2(\text{PMe}_3)_2]$, where replacement of two strong π -accepting CO ligands with PMe_3 shifts the L_1 rising-edge 3 eV to lower energy. The position of the L_1 rising-edge is strongly influenced by the number of π -acid ligands. Formally zerovalent $[W^0(\text{CO})_6]$ and $[W^0(\text{Me}_2\text{pipdt})(\text{CO})_4]$ have L_1 rising-edge energies that are among the highest shown in Figure 5. As expected, the W^0 atom in these compounds is depleted of charge by the CO ligands, and therefore the physical oxidation state, insofar as one can be gauged, is comparable to the W^V and W^{VI} reference compounds. Here, we

suggest that the effort to assess a spectroscopic oxidation state has meaning as an attempt to rationalize or anticipate some property, such as reactivity, not as a general system for classifying molecules.

For the series of bis(dithiolene) compounds of the type $[\text{W}^{\text{IV}}(\text{mdt})_2\text{XY}]$ ($X = Y = \text{CO}, \text{PMe}_3, \text{CN}^t\text{Bu}, \text{CN}^-$; $X = \text{CO}, Y = \text{PMe}_3$), the EBR and rising-edge energy are consistent with the interpretation of X-ray diffraction and S K-edge XAS studies.²² In the particular case of $[\text{W}^{\text{IV}}(\text{mdt})_2(\text{CO})_2]$ and $[\text{W}^{\text{IV}}(\text{mdt})_2(\text{CN})_2]^{2-}$, their comparable tungsten L_1 rising-edge energies reveal that the strikingly greater degree of dithiolene ligand oxidation in the former compound, where the ligands appear to be at a radical monoanionic redox level, as compared to the latter compound, where the ligands appear to be fully reduced ene-1,2-dithiolates, is offset by transfer of charge density into the CO ligands with little net effect upon tungsten Z_{eff} . For $[\text{W}^{\text{IV}}(\text{mdt})_2(\text{PMe}_3)_2]$, an increase in the bond length between metal and ancillary ligand and a decrease in π backdonation produce an L_1 rising-edge that is ~ 1.5 eV lower in energy than that of $[\text{W}^{\text{IV}}(\text{mdt})_2(\text{CO})_2]$. This energy is similar to that found for $[\text{W}^{\text{III}}\text{Cl}_2(\text{dppe})_2]^{1+}$, which has a similar array of σ - and π -donating ligands. The 1.743(1) Å S–C and 1.346(3) Å C–C_{chelate} dithiolene bond distances in $[\text{W}^{\text{IV}}(\text{mdt})_2(\text{PMe}_3)_2]$ are slightly shorter and longer, respectively, than the values typical of fully reduced ene-1,2-dithiolate.²² The covalency between metal and dithiolene implied by these bond lengths, taken with the absence of strongly perturbing π -acid ancillary ligand, makes it understandable that the tungsten ion in $[\text{W}^{\text{IV}}(\text{mdt})_2(\text{PMe}_3)_2]$ appears more reduced than in $[\text{W}^{\text{IV}}(\text{mdt})_2(\text{CO})_2]$ and $[\text{W}^{\text{IV}}(\text{mdt})_2(\text{CN})_2]^{2-}$ (Figure 5).

The invariance of the EBR, L_{3-} , and L_{2-} edge white-line energy and the L_1 rising-edge energy for the tris(dithiolene) electron-transfer series $[\text{W}(\text{mdt})_3]^z$ and $[\text{W}(\text{bdt})_3]^z$ ($z = 0, 1-, 2-$) align with crystallographic, EPR, and S K-edge XAS data, confirming that each member is related by a ligand-centered redox event. The L_1 pre-edge peak intensity correlates with the magnitude of the trigonal twist angle in these compounds. This observation demonstrates that the L_1 -edge has sufficient resolution to identify molecular geometry both in the solid and solution state, which is analogous to, albeit different from, the diagnostic pre-edge peak for tetrahedral sites in tungsten oxide materials.¹⁹ The spectral detail will be improved by using high-energy resolution fluorescence detection XAS, which has enhanced L-edge spectral definition for several third row metals.⁹³ The technique reduces the effect of lifetime broadening, thereby providing a target for time-dependent DFT simulation of the pre-edge and increasing the electronic structure detail obtained from these measurements. Furthermore, it would be advantageous to exploit solution-state studies to assess changes in geometry in the absence of lattice forces.^{16,81} Specifically for monoanionic tris(dithiolene) compounds, such measurements would provide a crucial link between the crystallographic structure and the putative solution-state geometry based on EPR and DFT results.⁷⁹

A primary conclusion emerging from this effort is that the L-edge X-ray absorption spectra of molecular systems containing 5d metals offer useful and relatively direct insight into electronic structure from the perspective of what one may consider to be the compound's most important part — the metal ion. The EBR and L_1 rising-edge energy collectively provide information that is different from other physical methods about the state of the metal ion when it is strongly

perturbed by π -acid or π -donor ligands and/or coordinated by redox-active ligands. Making the most of any third row metal L-edge XAS data is assisted when other data are available for comparison, and toward that end the EBR and L_1 rising-edge values presented here are intended to be useful in establishing whether some tungsten compound of interest has a metal Z_{eff} greater than, less than, or similar to one of the relatively simple calibrant compounds presented in Table 3. With due circumspection for the limitations and uncertainties in the methodology, we emphasize that metal L-edge X-ray absorption spectroscopy is most useful when complemented by data from other physical techniques, not when applied as a stand-alone method.

■ ASSOCIATED CONTENT

📄 Supporting Information

Complete crystallographic data for $[\text{W}^{\text{III}}\text{Cl}_2(\text{dppe})_2][\text{PF}_6]$ and $[\text{W}^{\text{VI}}(\text{xylylene})_3]$ in CIF format. Thermal ellipsoid plots of $[\text{W}^{\text{VI}}(\text{xylylene})_3]$ and of $[\text{W}^{\text{III}}\text{Cl}_2(\text{dppe})_2][\text{PF}_6]$ cation–anion pairs in the unit cell with complete atom labeling. Multi-frequency EPR and magnetic data for $[\text{W}^{\text{III}}\text{Cl}_2(\text{dppe})_2][\text{PF}_6]$ and $[\text{W}^{\text{V}}(\text{NPh})\text{Cl}_3(\text{PMe}_3)_2]$. Pseudo-Voigt deconvoluted and second derivative $L_{3,2}$ -edge spectra; comparative plots of L_1 -edge and first derivative spectra. Computational details and coordinates for optimized structures. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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Notes

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■ REFERENCES

(1) Jørgensen, C. K. *Oxidation Numbers and Oxidation States*; Springer: Heidelberg, Germany, 1969.

- (2) Hegedus, L. S. *Transition Metals in the Synthesis of Complex Organic Molecules*; University Science Books: Mill Valley, CA, 1994.
- (3) Crabtree, R. H. *The Organometallic Chemistry of the Transition Metals*, 3rd ed.; Wiley & Sons: New York, NY, 2001.
- (4) Chaudhuri, P.; Wieghardt, K. *Prog. Inorg. Chem.* **2001**, *50*, 151.
- (5) Palmer, J. H. *Struct. Bonding Berlin, Ger.* **2012**, *142*, 49.
- (6) Palmer, J. H.; Lancaster, K. M. *Inorg. Chem.* **2012**, *51*, 12473.
- (7) Bart, S. C.; Chlopek, K.; Bill, E.; Bouwkamp, M. W.; Lobkovsky, E.; Neese, F.; Wieghardt, K.; Chirik, P. J. *J. Am. Chem. Soc.* **2006**, *128*, 13901.
- (8) Nguyen, A. I.; Blackmore, K. J.; Carter, S. M.; Zarkesh, R. A.; Heyduk, A. F. *J. Am. Chem. Soc.* **2009**, *131*, 3307.
- (9) Heyduk, A. F.; Zarkesh, R. A.; Nguyen, A. I. *Inorg. Chem.* **2011**, *50*, 9849.
- (10) Solomon, E. I.; Hedman, B.; Hodgson, K. O.; Dey, A.; Szilagyi, R. K. *Coord. Chem. Rev.* **2005**, *249*, 97.
- (11) Sproules, S.; Wieghardt, K. *Coord. Chem. Rev.* **2011**, *255*, 837.
- (12) Krause, M. O.; Oliver, J. H. *J. Phys. Chem. Ref. Data* **1979**, *8*, 329.
- (13) Shulman, R. G.; Yafet, Y.; Eisenberger, P.; Blumberg, W. E. *Proc. Natl. Acad. Sci. U.S.A.* **1976**, *73*, 1384.
- (14) (a) Banerjee, P.; Sproules, S.; Weyhermüller, T.; DeBeer George, S.; Wieghardt, K. *Inorg. Chem.* **2009**, *48*, 5829. (b) Milsmann, C.; Sproules, S.; Bill, E.; Weyhermüller, T.; DeBeer George, S.; Wieghardt, K. *Chem.—Eur. J.* **2010**, *16*, 3628.
- (15) Sproules, S.; Benedito, F. L.; Bill, E.; Weyhermüller, T.; DeBeer George, S.; Wieghardt, K. *Inorg. Chem.* **2009**, *48*, 10926.
- (16) Sproules, S.; Weyhermüller, T.; DeBeer, S.; Wieghardt, K. *Inorg. Chem.* **2010**, *49*, 5241.
- (17) Szilagyi, R. K.; Lim, B. S.; Glaser, T.; Holm, R. H.; Hedman, B.; Hodgson, K. O.; Solomon, E. I. *J. Am. Chem. Soc.* **2003**, *125*, 9158.
- (18) (a) Balerna, A.; Bernieri, E.; Burattini, E.; Kuzmin, A.; Lusia, A.; Purans, J.; Cikmach, P. *Nucl. Instrum. Methods Phys. Res., Sect. A* **1991**, *308*, 240. (b) Kuzmin, A.; Purans, J. *J. Phys.: Condens. Matter* **1993**, *5*, 9423. (c) Tougerti, A.; Cristol, S.; Berrier, E.; Briois, V.; La Fontaine, C.; Villain, F.; Joly, Y. *Phys. Rev. B* **2012**, *85*, 125136. (d) Uehara, Y.; Kawase, K.; Tsuchimoto, J.; Shibano, T. *J. Electron Spectrosc. Relat. Phenom.* **2005**, *148*, 75.
- (19) (a) Barton, D. G.; Soled, S. L.; Meitzner, G. D.; Fuentes, G. A.; Iglesia, E. *J. Catal.* **1999**, *181*, 57. (b) Hilbrig, F.; Göbel, H. E.; Knözinger, H.; Schmelz, H.; Lengeler, B. *J. Phys. Chem.* **1991**, *95*, 6973. (c) Horsley, J. A.; Wachs, I. E.; Brown, J. M.; Via, G. H.; Hardcastle, F. D. *J. Phys. Chem.* **1987**, *91*, 4014. (d) Yamamoto, T.; Orita, A.; Tanaka, T. *X-Ray Spectrom.* **2008**, *37*, 226.
- (20) Chandrasekaran, P.; Arumugam, K.; Jayarathne, U.; Pérez, L. M.; Mague, J. T.; Donahue, J. P. *Inorg. Chem.* **2009**, *48*, 2103.
- (21) Yan, Y.; Chandrasekaran, P.; Mague, J. T.; DeBeer, S.; Sproules, S.; Donahue, J. P. *Inorg. Chem.* **2012**, *51*, 346.
- (22) Yan, Y.; Keating, C.; Chandrasekaran, P.; Jayarathne, U.; Mague, J. T.; DeBeer, S.; Lancaster, K. M.; Sproules, S.; Rubtsov, I. V.; Donahue, J. P. *Inorg. Chem.* **2013**, *52*, 6743.
- (23) Cho, D.-Y.; Park, J.; Yu, J.; Park, J.-G. *J. Phys.: Condens. Matter* **2012**, *24*, 055503.
- (24) (a) Daulton, T. L.; Little, B. J. *Ultramicroscopy* **2006**, *106*, 561. (b) Zhang, Z. *Ultramicroscopy* **2007**, *107*, 598.
- (25) (a) Koshino, M.; Kurata, H.; Isoda, S.; Kobayashi, T. *Micron* **2000**, *31*, 373. (b) Thole, B. T.; van der Laan, G. *Phys. Rev. B* **1988**, *38*, 3158.
- (26) Qi, B.; Perez, I.; Ansari, P. H.; Lu, F.; Croft, M. *Phys. Rev. B* **1987**, *36*, 2972.
- (27) Sharp, P. R. *Organometallics* **1984**, *3*, 1217.
- (28) Over, D. E.; Critchlow, S. C.; Mayer, J. M. *Inorg. Chem.* **1992**, *31*, 4643.
- (29) Sharp, P. R.; Bryan, J. C.; Mayer, J. M.; Templeton, J. L.; Feng, S. *Inorg. Synth.* **1990**, *28*, 326.
- (30) Nielson, A. J.; Waters, J. M. *Aust. J. Chem.* **1983**, *36*, 243.
- (31) Lappert, M. F.; Raston, C. L.; Rowbottom, G. L.; Skelton, B. W.; White, A. H. *J. Chem. Soc., Dalton Trans.* **1984**, 883.
- (32) Goddard, C. A.; Holm, R. H. *Inorg. Chem.* **1999**, *38*, 5389.
- (33) McCleverty, J. A.; Locke, J.; Wharton, E. J.; Gerloch, M. *J. Chem. Soc. A* **1968**, 816.
- (34) Sellmann, D.; Kern, W.; Moll, M. *J. Chem. Soc., Dalton Trans.* **1991**, 1733.
- (35) Fomitchev, D. V.; Lim, B. S.; Holm, R. H. *Inorg. Chem.* **2001**, *40*, 645.
- (36) Stiefel, E. I.; Eisenberg, R.; Rosenberg, R. C.; Gray, H. B. *J. Am. Chem. Soc.* **1966**, *88*, 2956.
- (37) APEX2, Version 2009.11–0; Bruker-AXS, Inc.: Madison, WI, 2009.
- (38) SAINT, Version 7.68A; Bruker-AXS, Inc.: Madison, WI, 2009.
- (39) Sheldrick, G. M. SADABS, Version 2008/2; Universität Göttingen: Göttingen, Germany, 2008.
- (40) Sheldrick, G. M. SHELXS-97, Universität Göttingen: Göttingen, Germany, 2008.
- (41) Sheldrick, G. M. SHELXL-97, Universität Göttingen: Göttingen, Germany, 2008.
- (42) SHELXTL, Version 2008/4; Bruker-AXS, Inc.: Madison, WI, 2008.
- (43) George, G. N. EXAFSPAK & EDG_FIT; Stanford Synchrotron Radiation Laboratory, Stanford Linear Accelerator Center, Stanford University: Stanford, CA, 2000.
- (44) Hanson, G. R.; Gates, K. E.; Noble, C. J.; Griffin, M.; Mitchell, A.; Benson, S. *J. Inorg. Biochem.* **2004**, *98*, 903.
- (45) Frisch, M. J.; Trucks, G. W.; Schlegel, H. B.; Scuseria, G. E.; Robb, M. A.; Cheeseman, J. R.; Scalmani, G.; Barone, V.; Mennucci, B.; Petersson, G. A.; Nakatsuji, H.; Caricato, M.; Li, X.; Hratchian, H. P.; Izmaylov, A. F.; Bloino, J.; Zheng, G.; Sonnenberg, J. L.; Hada, M.; Ehara, M.; Toyota, K.; Fukuda, R.; Hasegawa, J.; Ishida, M.; Nakajima, T.; Honda, Y.; Kitao, O.; Nakai, H.; Vreven, T.; Montgomery, J. A., Jr.; Peralta, J. E.; Ogliaro, F.; Bearpark, M.; Heyd, J. J.; Brothers, E.; Kudin, K. N.; Staroverov, V. N.; Kobayashi, R.; Normand, J.; Raghavachari, K.; Rendell, A.; Burant, J. C.; Iyengar, S. S.; Tomasi, J.; Cossi, M.; Rega, N.; Millam, N. J.; Klene, M.; Knox, J. E.; Cross, J. B.; Bakken, V.; Adamo, C.; Jaramillo, J.; Gomperts, R.; Stratmann, R. E.; Yazyev, O.; Austin, A. J.; Cammi, R.; Pomelli, C.; Ochterski, J. W.; Martin, R. L.; Morokuma, K.; Zakrzewski, V. G.; Voth, G. A.; Salvador, P.; Dannenberg, J. J.; Dapprich, S.; Daniels, A. D.; Farkas, Ö.; Foresman, J. B.; Ortiz, J. V.; Cioslowski, J.; Fox, D. J. *Gaussian 09*; Gaussian, Inc., Wallingford, CT, 2009.
- (46) Perdew, J. P.; Burke, K.; Ernzerhof, M. *Phys. Rev. Lett.* **1996**, *77*, 3865.
- (47) (a) Becke, A. D. *J. Chem. Phys.* **1993**, *98*, 5648. (b) Lee, C. T.; Yang, W. T.; Parr, R. G. *Phys. Rev. B* **1988**, *37*, 785.
- (48) <https://bse.pnl.gov/bse/portal> (accessed November 15, 2013).
- (49) Schäfer, A.; Horn, H.; Ahlrichs, R. *J. Chem. Phys.* **1992**, *97*, 2571.
- (50) Tenderholt, A. L. QMForge: A Program to Analyze Quantum Chemistry Calculations, Version 2.1; <http://qmforge.sourceforge.net> (accessed November 15, 2013).
- (51) Chemcraft, Version 1.6; <http://chemcraftprog.com> (accessed November 15, 2013).
- (52) Jmol: an open source Java viewer for chemical structures in 3D. <http://jmol.sourceforge.net> (accessed November 15, 2013).
- (53) (a) Rabinovich, D.; Parkin, G. *J. Am. Chem. Soc.* **1990**, *112*, 5381. (b) Rabinovich, D.; Zelman, R.; Parkin, G. *J. Am. Chem. Soc.* **1992**, *114*, 4611.
- (54) Schaefer King, M. A.; McCarty, R. E. *Inorg. Chem.* **1973**, *12*, 1972.
- (55) Blight, D. G.; Kepert, D. L. *J. Chem. Soc. A* **1968**, 534.
- (56) Boorman, P. M.; Greenwood, N. N.; Hildon, M. A. *J. Chem. Soc. A* **1968**, 2466.
- (57) Jayarathne, U.; Chandrasekaran, P.; Jacobsen, H.; Mague, J. T.; Donahue, J. P. *Dalton Trans.* **2010**, 39, 9662.
- (58) Imaeda, M.; Nishihara, H.; Nakano, K.; Ichida, H.; Kobayashi, A.; Saito, T.; Sasaki, Y. *Inorg. Chem.* **1985**, *24*, 1246.
- (59) Filippou, A. C.; Schnakenburg, G.; Philippopoulos, A. I. *Acta Crystallogr., Sect. E: Struct. Rep. Online* **2003**, *59*, m602.
- (60) Taylor, J. R. *An Introduction to Error Analysis*; University Science Books: Sausalito, CA, 1997.

- (61) Mabbs, F. E.; Collison, D. *Electron Paramagnetic Resonance of d Transition Metal Compounds*; Elsevier: Amsterdam, 1992.
- (62) (a) Adams, C. J.; Anderson, K. M.; Connelly, N. G.; Harding, D. J.; Hayward, O. D.; Orpen, A. G.; Patrón, E.; Rieger, P. H. *Dalton Trans.* **2009**, 530. (b) Ceausescu, E.; Cornilescu, A.; Nicolescu, E.; Popescu, M.; Coca, S.; Belloiu, C.; Oprea, C.; Dimonie, M.; Hubca, G.; Dragutan, V.; Chipara, M. *J. Mol. Catal.* **1985**, *28*, 351.
- (63) Yoon, K.; Parkin, G.; Rheingold, A. L. *J. Am. Chem. Soc.* **1991**, *113*, 1437.
- (64) Nielson, A. J.; Boyd, P. D. W.; Clark, G. R.; Hunt, P. A.; Hursthouse, M. B.; Metson, J. B.; Rickard, C. E. F.; Schwerdtfeger, P. *J. Chem. Soc., Dalton Trans.* **1995**, 1153.
- (65) Levason, W.; McAuliffe, C. A.; McCullough, F. P., Jr. *Inorg. Chem.* **1977**, *16*, 2911.
- (66) Nielson, A. J.; Metson, J. B. *Polyhedron* **2012**, *31*, 143.
- (67) Horsley, J. A. *J. Chem. Phys.* **1982**, *76*, 1451.
- (68) Wang, H.; Ge, P.; Riordan, C. G.; Brooker, S.; Woome, C. G.; Collins, T.; Melendres, C. A.; Graudejus, O.; Bartlett, N.; Cramer, S. P. *J. Phys. Chem. B* **1998**, *102*, 8343.
- (69) Wei, P. S. P.; Lytle, F. W. *Phys. Rev. B* **1979**, *19*, 679.
- (70) (a) Drube, W.; Treusch, R.; Sham, T. K.; Bzowski, A.; Soldatov, A. V. *Phys. Rev. B* **1998**, *58*, 6871. (b) Stöhr, J. *J. Electron Spectrosc. Relat. Phenom.* **1995**, *75*, 253.
- (71) Choi, Y. G. *Met. Mater. Int.* **2009**, *15*, 993.
- (72) Kosog, B.; La Pierre, H. S.; Denecke, M. A.; Heinemann, F. W.; Meyer, K. *Inorg. Chem.* **2012**, *51*, 7940.
- (73) (a) Shimizu, K.; Oda, T.; Sakamoto, Y.; Kamiya, Y.; Yoshida, H.; Satsuma, A. *Appl. Catal., B* **2012**, *111–112*, 509. (b) Yoshida, H.; Nonoyama, S.; Yazawa, Y.; Hattori, T. *Phys. Scr.* **2005**, *T115*, 813.
- (74) (a) de Groot, F. *Coord. Chem. Rev.* **2005**, *249*, 31. (b) de Groot, F. M. F.; Hu, Z. W.; Lopez, M. F.; Kaindl, G.; Guillot, F.; Tronc, M. *J. Chem. Phys.* **1994**, *101*, 6570. (c) Sham, T. K. *J. Am. Chem. Soc.* **1983**, *105*, 2269.
- (75) (a) Keast, V. J.; Scott, A. J.; Brydson, R.; Williams, D. B.; Bruley, J. *J. Microscopy* **2001**, *203*, 135. (b) Leapman, R. D.; Grunes, L. A. *Phys. Rev. Lett.* **1980**, *45*, 397. (c) Pearson, D. H.; Ahn, C. C.; Fultz, B. *Phys. Rev. B* **1993**, *47*, 8471. (d) Sparrow, T. G.; Williams, B. G.; Rao, C. N. R.; Thomas, J. M. *Chem. Phys. Lett.* **1984**, *108*, 547.
- (76) (a) Galyer, A. L.; Wilkinson, G. *J. Chem. Soc., Dalton Trans.* **1976**, 2235. (b) Galyer, L.; Mertis, K.; Wilkinson, G. *J. Organomet. Chem.* **1975**, *85*, C37.
- (77) Bristow, G. S.; Lappert, M. F.; Martin, T. R.; Atwood, J. L.; Hunter, W. F. *J. Chem. Soc., Dalton Trans.* **1984**, 399.
- (78) Campbell, S.; Harris, S. *Inorg. Chem.* **1996**, *35*, 3285.
- (79) Sproules, S.; Banerjee, P.; Weyhermüller, T.; Yan, Y.; Donahue, J. P.; Wieghardt, K. *Inorg. Chem.* **2011**, *50*, 7106.
- (80) Sproules, S.; Weyhermüller, T.; Goddard, R.; Wieghardt, K. *Inorg. Chem.* **2011**, *50*, 12623.
- (81) Sproules, S. *Prog. Inorg. Chem.* **2014**, *58*, 1.
- (82) Almahamid, L.; Bryan, J. C.; Bucher, J. J.; Burrell, A. K.; Edelstein, N. M.; Hudson, E. A.; Kaltsoyannis, N.; Lukens, W. W.; Shuh, D. K.; Nitsche, H.; Reich, T. *Inorg. Chem.* **1995**, *34*, 193.
- (83) Sattler, A.; Parkin, G. *Nature* **2010**, *463*, 523.
- (84) (a) Thomas, S.; Tiekink, E. R. T.; Young, C. G. *Organometallics* **1996**, *15*, 2428. (b) Thomas, S.; Tiekink, E. R. T.; Young, C. G. *Inorg. Chem.* **2006**, *45*, 352.
- (85) Kutzler, F. W.; Natoli, C. R.; Misemer, D. K.; Doniach, S.; Hodgson, K. O. *J. Chem. Phys.* **1980**, *73*, 3274.
- (86) Cotton, F. A.; Kibala, P. A.; Sandor, R. B. W. *Acta Crystallogr., Sect. C* **1989**, *45*, 1287.
- (87) Kapre, R. R.; Bothe, E.; Weyhermüller, T.; DeBeer George, S.; Wieghardt, K. *Inorg. Chem.* **2007**, *46*, 5642.
- (88) Huynh, H.; Lügger, T.; Hahn, F. E. *Eur. J. Inorg. Chem.* **2002**, 3007.
- (89) Burrow, T. E.; Morris, R. H.; Hills, A.; Hughes, D. L.; Richards, R. L. *Acta Crystallogr., Sect. C* **1993**, *49*, 1591.
- (90) Lorber, C.; Donahue, J. P.; Goddard, C. A.; Nordlander, E.; Holm, R. H. *J. Am. Chem. Soc.* **1998**, *120*, 8102.
- (91) Knoch, F.; Sellmann, D.; Kern, W. *Z. Kristallogr.* **1993**, *205*, 300.
- (92) Sugimoto, H.; Furukawa, Y.; Tarumizu, M.; Miyake, H.; Tanaka, K.; Tsukube, H. *Eur. J. Inorg. Chem.* **2005**, 3088.
- (93) (a) Glatzel, P.; Singh, J.; Kvashnina, K. O.; van Bokhoven, J. A. *J. Am. Chem. Soc.* **2010**, *132*, 2555. (b) Hämäläinen, K.; Siddons, D. P.; Hastings, J. B.; Berman, L. E. *Phys. Rev. Lett.* **1991**, *67*, 2850. (c) Makosch, M.; Kartusch, C.; Sá, J.; Bessa Duarte, R.; van Bokhoven, J. A.; Kvashnina, K.; Glatzel, P.; Fernandes, D. L. A.; Nachttegaal, M.; Kleymenov, E.; Szlachetko, J.; Neuhold, B.; Hungerbühler, K. *Phys. Chem. Chem. Phys.* **2012**, *14*, 2164. (d) Safonova, O. V.; Tromp, M.; van Bokhoven, J. A.; de Groot, F. M. F.; Evans, J.; Glatzel, P. *J. Phys. Chem. B* **2006**, *110*, 16162. (e) Swarbrick, J. C.; Skyllberg, U.; Karlsson, T.; Glatzel, P. *Inorg. Chem.* **2009**, *48*, 10748.