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Isolation of Ce(IV) centered polyoxoalkoxide sandwich-type complexes allows comparison of metal–oxygen bond covalency

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The new Ce^{III} centered sandwich-type complex (TBA)₃[Ce(W₄O₁₃(OMe)₄MoNO)₂] is reported. The redox properties of this molecule, and its all-molybdenum analogue, (TBA)₃[Ce(Mo₅O₁₃(OMe)₄NO)₂], were investigated using cyclic voltammetry. The data reveals the presence of reversible Ce^{IV}/Ce^{III} redox couples at modest potentials. One electron oxidation of the complexes provides facile access to the corresponding Ce^{IV} derivatives, which were fully characterized. ¹⁷O NMR spectroscopy reveals that the chemical shifts of the oxygen nuclei directly bound to Ce^{IV} are much higher than the corresponding signals in isostructural, diamagnetic, Zr^{IV}, Hf^{IV}, or Th^{IV} centered complexes. Density functional theory (DFT) calculations indicate that the increase in chemical shift correlates with an increase in the covalency of the M^{IV}–O bonds, illustrating that ¹⁷O NMR spectroscopy is a powerful experimental tool for interrogating the nature of metal oxygen bonding in diamagnetic complexes.

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Introduction

The manipulation of lanthanide (Ln) and actinide (An) redox states is exploited in separation processes, where the ability to selectively oxidize certain metal cations facilitates their isolation from complex mixtures.^{1–3} In this context, cerium is unique amongst the lanthanides due to its ability to access a [Xe]4f⁰ electronic configuration upon oxidation to the +4 oxidation state.^{4,5} Other high valent lanthanides are rarely observed, with only a handful of molecular complexes known.^{6–11} The ability to readily cycle between Ce^{III} and Ce^{IV} has been exploited extensively in organic chemistry, with cerium complexes being applied as oxidants in a range of organic transformations, while cerium oxide has been used catalytic converters in petrol cars to support the oxidation of CO to CO₂.^{12–18}

Trivalent lanthanides are often considered as possessing localized, core-like, valence 4f orbitals and therefore are said to primarily form ionic bonds.^{19,20} However, recent work has shown that oxidation of Ce^{III} to Ce^{IV} can “turn on” mixing of metal 4f orbitals with ligand orbitals, leading to the formation of bonds with a higher degree of covalent character.²¹ This phenomenon is well documented for [Ce^{III}Cl₆]^{3–} and [Ce^{IV}Cl₆]^{2–}, where Cl K-edge X-ray absorption spectroscopy (XAS) demonstrates a substantial increase in Ce–Cl orbital mixing upon oxidation of the lanthanide from Ce^{III} to Ce^{IV}.^{20–22} These results were compared to a series of [M^{IV}Cl₆]^{2–} (M = Ti, Zr, Hf, and U) and, interestingly, the 4f-orbital

participation in the Ce–Cl bonds of [Ce^{IV}Cl₆]^{2–} was observed to be more than twice that of the 5f-orbital contribution to the corresponding U–Cl bonds of [U^{IV}Cl₆]^{2–}. This was attributed to the lower energy of the Ce 4f vs. U 5f-orbitals, implying that matching the energy of Cl 3p orbitals with the f-orbitals is a stronger contributor to the covalency of these types of bonds. A number of studies utilizing XAS experiments and/or computational calculations to assess the covalency of Ln–E/An–E bonds (E = C, N, O, S) exist.^{4,23–35} Nuclear magnetic resonance (NMR) spectroscopy has also proved useful in many studies where simulation of NMR spectra is often used as a tool to verify the accuracy of computational studies.^{36–39} Furthermore, the spin–orbit contributions (σ_{SO}) to the nuclear shielding constant (σ) often provide valuable insights when assessing bond covalency.^{40–43}

To carry out investigations of this type, isostructural An and Ln containing complexes are required to make like for like comparisons. In this context, polyoxometalates (POMs), anionic molecular metal oxide clusters typically based on W^{VI}, Mo^{VI}, or V^V, have become popular ligands in lanthanide and actinide chemistry.^{44–61} While some studies with Ce–POM systems focus on catalytic^{62,63} and biological^{64–67} applications, many of the studies in this field limit analysis to solid state characterization. Indeed, despite the fact that a growing number of isostructural Ln–POM/An–POM systems are known, there are no studies which employ these complexes as frameworks for systematic comparison of Ln–O/An–O bonding.

Previously, our group utilized lacunary polyoxoalkoxide complexes, (TBA)₂[Mo₅O₁₃(OMe)₄NO][Na(MeOH)] (1-NaMo₅) and (TBA)₂[W₄O₁₃(OMe)₄MoNO][Na(MeOH)] (1-NaW₄Mo), for the synthesis of a series of M^{IV} centered sandwich-type

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assemblies with the general formula $(\text{TBA})_2[\text{M}^{\text{IV}}\{\text{M}'_4\text{O}_{13}(\text{OMe})_4\text{MoNO}\}_2]$ ($\text{M}^{\text{IV}} = \text{Zr, Hf, Th, Np, U}$ and $\text{M}' = \text{Mo}$ or W).^{68–70} Detailed structural analysis was performed by single crystal X-ray diffraction (SCXRD). Additionally, the solubility of these complexes renders solution phase analysis possible. Indeed, characterization of the $(\text{TBA})_2[\text{M}^{\text{IV}}\{\text{M}'_4\text{O}_{13}(\text{OMe})_4\text{MoNO}\}_2]$ complexes by ^{17}O NMR spectroscopy offers a direct spectroscopic handle for the $\text{M}^{\text{IV}}\text{--O}$ bond. Other groups have previously isolated several isostructural lanthanide centered complexes with the general formula $(\text{TBA})_3[\text{Ln}^{\text{III}}\{\text{Mo}_5\text{O}_{13}(\text{OMe})_4\text{NO}\}_2]$, where $\text{Ln}^{\text{III}} = \text{Ce, Eu, Tb, Dy, Ho, and Er}$. These were obtained either by treatment of **1-NaMo₅** with an appropriate lanthanide salt in methanol⁷¹ or by treatment of $(\text{TBA})_4[\alpha\text{-Mo}_8\text{O}_{26}]$ with hydroxylamine, dicyclohexylcarbodiimide (DCC), and the appropriate lanthanide nitrate in methanol.⁷² The related complexes $(\text{TBA})_3[\text{Ln}^{\text{III}}\{\text{Mo}_5\text{O}_{13}(\text{OMe})_4\text{NNC}_6\text{H}_4\text{-}p\text{-NO}_2\}_2]$ (where $\text{Ln}^{\text{III}} = \text{Tb, Dy, Ho, Er, Yb, and Nd}$) have also been reported and their magnetic properties were investigated.⁷³ Interestingly, no attempts were made to interrogate the electrochemistry of any of these lanthanide complexes and, consequently, Ce^{IV} centered complexes remain unknown.

Herein, we extend Proust and Villaneau's method for the synthesis of $(\text{TBA})_3[\text{Ce}^{\text{III}}\{\text{Mo}_5\text{O}_{13}(\text{OMe})_4\text{NO}\}_2]$ (**2-Ce(Mo₅)₂**) to access the tungsten containing analogue $(\text{TBA})_3[\text{Ce}^{\text{III}}\{\text{W}_4\text{O}_{13}(\text{OMe})_4\text{MoNO}\}_2]$ (**2-Ce(W₄Mo)₂**).⁷¹ Both complexes were fully characterized by ^1H NMR spectroscopy, ^{17}O NMR spectroscopy, electronic absorption spectroscopy, SCXRD and cyclic voltammetry. Importantly, the cyclic voltammograms (CVs) of **2-Ce(Mo₅)₂** and **2-Ce(W₄Mo)₂** show the presence of a reversible $\text{Ce}^{\text{IV}}/\text{Ce}^{\text{III}}$ redox couple at *ca.* 0.3–0.4 V *vs.* $\text{Fc}^{+/0}$. With this knowledge in hand, we carried out one electron oxidation to isolate the Ce^{IV} centered complexes $(\text{TBA})_2[\text{Ce}^{\text{IV}}\{\text{Mo}_5\text{O}_{13}(\text{OMe})_4\text{NO}\}_2]$ (**3-Ce(Mo₅)₂**) and $(\text{TBA})_2[\text{Ce}^{\text{IV}}\{\text{W}_4\text{O}_{13}(\text{OMe})_4\text{MoNO}\}_2]$ (**3-Ce(W₄Mo)₂**). Electronic absorption spectroscopy of the oxidized complexes reveals the presence of a new ligand-to-metal charge transfer (LMCT) process between the polyoxoalkoxo metalloligand and the empty $4f^0$ orbital of Ce^{IV} , with this assignment supported by time-dependent density functional theory (TD-DFT) calculations. ^{17}O NMR spectroscopy of the now diamagnetic **3-Ce(Mo₅)₂** and **3-Ce(W₄Mo)₂** complexes shows that the chemical shifts of the oxygen nuclei directly bound to Ce^{IV} are much higher than the corresponding signals in the other diamagnetic M^{IV} centered complexes we have reported.^{68–70} This increase in chemical shift correlates with an increase in metal orbital contribution to the $\text{M}^{\text{IV}}\text{--O}$ bonds and delocalization index (DI), both of which were obtained from DFT calculations and can be used as a measure for bond covalency. Collectively, these data show that ^{17}O NMR chemical shift acts as an experimental handle for M--O bond covalency in diamagnetic polyoxometalate complexes.³³

Experimental

General considerations

Air- and moisture-sensitive manipulations with all complexes were carried out using a standard high-vacuum

line, Schlenk techniques, or an MBraun inert atmosphere drybox containing an atmosphere of purified dinitrogen. The MBraun glovebox was equipped with a cold well designed for freezing samples in liquid nitrogen, as well as a $-35\text{ }^\circ\text{C}$ freezer for cooling samples and crystallizations. Solvents for sensitive manipulations were dried and deoxygenated using literature procedures with a Seca solvent purification system or a glass contour solvent purification system (Pure Process Technology, LLC) and stored over activated 4 Å molecular sieves (Fisher Scientific) prior to use. Deuterated solvents were purchased from Cambridge Isotope Laboratories, dried with molecular sieves and degassed by three freeze-pump-thaw cycles. 40% ^{17}O enriched H_2O was purchased from CortecNet and used as received. $(\text{TBA})_4[\text{Mo}_8\text{O}_{26}]$,⁷⁴ $(\text{TBA})_2[\text{WO}_4]$,⁷⁵ $(\text{TBA})_2[\text{Mo}_5\text{O}_{13}(\text{OMe})_4\text{NO}][\text{Na}(\text{MeOH})]$ (**1-NaMo₅**)⁷⁶ and, $(\text{TBA})_2[\text{W}_4\text{O}_{13}(\text{OMe})_4\text{MoNO}][\text{Na}(\text{MeOH})]$ (**1-NaW₄Mo**)^{68,71} were synthesized according to literature procedures. The synthesis of **2-Ce(Mo₅)₂** and **2-Ce(W₄Mo)₂** was adapted from the previously reported procedure.⁷¹ $\text{Ce}(\text{OTf})_3$ was purchased from Strem Chemicals, while all other reagents were purchased from commercial sources (Fisher Scientific, VWR, and Sigma-Aldrich) and used without further purification.

General procedure for the synthesis of

$(\text{TBA})_3[\text{Ce}^{\text{III}}\{\text{M}_4\text{O}_{13}(\text{OMe})_4\text{MoNO}\}_2]$ ($\text{M} = \text{Mo}$, **2-Ce(Mo₅)₂**, $\text{M} = \text{W}$, **2-Ce(W₄Mo)₂**)

In a 15 mL pressure vessel, $(\text{TBA})_2[\text{M}_4\text{O}_{13}(\text{OMe})_4\text{MoNO}][\text{Na}(\text{MeOH})]$ (0.36 mmol, 2 eq.) was dissolved in MeOH (4 mL) forming a purple solution. $\text{Ce}(\text{OTf})_3$ ($\text{OTf} = \text{O}_3\text{SCF}_3$; 117 mg, 0.20 mmol, 1.1 eq.) was dissolved in MeOH (4 mL) and added slowly to the mixture. The pressure vessel was sealed, and the solution was heated at $50\text{ }^\circ\text{C}$ for 2 hours with stirring. The hot reaction mixture was passed through a filter paper to remove any precipitate. The filtered reaction mixture was then cooled to $-30\text{ }^\circ\text{C}$ in a freezer and stored overnight, after which block-shaped crystals formed. The mother liquor was decanted, and the crystals were washed with cold MeOH (2 mL) and Et_2O (10 mL \times 2). The crystals were then dried under vacuum.

2-Ce(Mo₅)₂. Red/violet solid (317 mg, 68% yield). ^1H NMR (500 MHz, CD_3CN): δ (ppm) 1.17 (t, 36H), 1.66 (m, 24H), $-\text{CH}_2$ peak overlap with reference peak at 1.94, 3.34 (t, 24H), 3.84 (s, 24H). ^{17}O NMR (67.8 MHz, CD_3CN): δ (ppm) -41.6 ($\mu_5\text{-O}$), 479.7 (Mo–O–Mo), 879.8 (Mo=O), 886.8 (Ce–O–Mo). λ_{max} (MeCN) = 552 nm ($\epsilon = 144\text{ mol}^{-1}\text{ dm}^3\text{ cm}^{-1}$). Anal. Calcd. for $\text{C}_{56}\text{H}_{132}\text{N}_5\text{Mo}_{10}\text{O}_{36}\text{Ce}$ (mol. wt 2551.287 g mol⁻¹): C, 26.36%; H, 5.22%; N, 2.75%. Found: C, 26.452%; H, 4.927%; N, 2.802%.

2-Ce(W₄Mo)₂. Blue/purple solid (334 mg, 57% yield). ^1H NMR (500 MHz, CD_3CN): δ (ppm) 1.42 (t, 36H), 2.06 (m, 24H), 2.38 (m, 24H), 3.18 (s, 24H), 3.92 (t, 24H). ^{17}O NMR (67.8 MHz, CD_3CN): δ (ppm) -76.8 ($\mu_5\text{-O}$), 356.4 (W–O–W), 715.7 (W=O), 740.4 (Ce–O–W). λ_{max} (MeCN) = 564 nm ($\epsilon = 129\text{ mol}^{-1}\text{ dm}^3\text{ cm}^{-1}$). Anal. Calcd. for $\text{C}_{56}\text{H}_{132}\text{N}_5\text{W}_8\text{Mo}_2\text{O}_{36}\text{Ce}$ (mol. wt 3254.407 g mol⁻¹): C, 20.67%; H, 4.09%; N, 2.15%. Found: C, 20.580%; H, 3.815%; N, 2.146%.



General procedure for the synthesis of

(TBA)₂[Ce^{IV}{M₄O₁₃(OME)₄MoNO}₂] (M = Mo, 3-Ce(Mo₅)₂, M = W, 3-Ce(W₄Mo)₂)

In a 20 mL scintillation vial, (TBA)₃[Ce{M₄O₁₃(OME)₄MoNO}₂] (0.039 mmol, 1 eq.) was dissolved in MeCN (2–3 mL) forming a homogenous solution. The solution was transferred to a vial containing solid [N(C₆H₄Br-4)₃][SbCl₆] (35 mg, 0.043 mmol, 1.1 eq.). The solution immediately turns dark orange/brown when using 2-Ce(Mo₅)₂ or dark yellow/brown when using 3-Ce(W₄Mo)₂. The mixture was stirred 15 minutes, and subsequently dried under vacuum leaving a dark residue. The residue was suspended in MeOH (5 mL) and the resultant suspension filtered through a bed of Celite (*ca.* 1 cm) and the eluent was discarded. The solid was washed with MeCN : Et₂O (1 : 5, 2 mL) and then extracted with DCM until eluent ran colorless (*ca.* 5 mL). This solution was then dried under vacuum to give crude product.

3-Ce(Mo₅)₂. Orange solid (70 mg, 78% yield). Dark orange block-shaped single crystals were obtained by vapor diffusion of Et₂O into a saturated solution of the product in MeCN at room temperature. ¹H NMR (500 MHz, CD₂Cl₂): δ (ppm) 1.12 (t, 24H), 1.58 (m, 16H), 1.83 (m, 16H), 3.34 (s, 24H), 4.73 (t, 24H). ¹⁷O NMR (67.8 MHz, CD₂Cl₂): δ (ppm) 16.3 (μ₅-O), 534.9 (Mo-O-Mo), 789.7 (Ce-O-Mo), 925.1 (Mo=O). Anal. Calcd. for C₄₀H₉₆N₄Mo₁₀O₃₆Ce_{0.5} Et₂O (mol. wt 2345.878 g mol⁻¹): C, 21.50%; H, 4.34%; N, 2.39%. Found: C, 21.748%; H, 4.389%; N, 2.486%.

3-Ce(W₄Mo)₂. Yellow solid (81 mg, 69% yield). Yellow block-shaped single crystals were obtained by vapor diffusion of Et₂O into a saturated solution of the product in MeCN at room temperature. ¹H NMR (500 MHz, CD₂Cl₂): δ (ppm) 1.12 (t, 24H), 1.59 (m, 16H), 1.85 (m, 16H), 3.37 (s, 24H), 4.87 (t, 24H). ¹⁷O NMR (67.8 MHz, CD₂Cl₂): δ (ppm) -12.8 (μ₅-O), 395.8 (W-O-W), 660.8 (Ce-O-W), 755.9 (W=O). λ_{max} (MeCN) = 568 nm (ε ≈ 105 mol⁻¹ dm³ cm⁻¹). Anal. Calcd. for C₄₀H₉₆N₄Mo₁₀O₃₆Ce.MeOH (mol. wt 3043.978 g mol⁻¹): C, 16.18%; H, 3.31%; N, 1.84%. Found: C, 16.407%; H, 2.994%; N, 1.627%.

Physical measurements

¹H NMR spectra for all other compounds were recorded at room temperature on a 400 MHz Bruker AVANCE spectrometer, a 500 MHz Bruker AVANCE spectrometer, or a JEOL 500 spectrometer and locked on the signal of deuterated solvents. All chemical shifts are reported relative to tetramethylsilane using the chosen deuterated solvent as a standard. ¹⁷O NMR spectra were collected at room temperature on a JEOL 500 spectrometer or a 500 MHz Bruker AVANCE spectrometer (both at 67.8 MHz), with the spectrometer locked on the signal of the deuterated solvent and all chemical shifts given relative to an external standard of D₂O. Cyclic voltammetry (CV) was performed using a three-electrode setup inside a glove box (MBraun UniLab, USA) using a Bio-Logic SP 150 potentiostat/galvanostat. The concentration of the cluster and the supporting electrolyte (TBAPF₆) were kept at 1 mM and 100 mM respectively throughout all measurements. CVs were recorded using a 3 mm diameter glassy carbon working electrode (CH Instruments,

USA), a Pt wire auxiliary electrode (CH Instruments, USA), and a Ag/Ag⁺ non-aqueous reference electrode with 0.01 M AgNO₃ in 0.1 M TBAPF₆ in acetonitrile (BASi, USA). Ferrocene was used as an internal standard after completion of the measurements, and potentials were referenced *versus* the Fc⁺⁰ couple. Electronic absorption measurements were recorded at room temperature in anhydrous MeCN or DCM in sealed 1 cm quartz cuvettes using an Agilent Cary 6000i UV-vis-NIR spectrophotometer. Elemental analysis data were obtained from the Elemental Analysis Facility at the University of Rochester. Microanalysis samples were weighed with a PerkinElmer model AD6000 autobalance, and their compositions were determined with a PerkinElmer 2400 series II analyzer. Air-sensitive samples were handled in a VAC Atmospheres glovebox.

X-ray crystallography

Crystals were placed onto a nylon loop and mounted on a Rigaku XtaLAB Synergy-S Dualflex diffractometer equipped with a HyPix-6000HE HPC area detector for data collection at 100.00(10) K. A preliminary set of cell constants and an orientation matrix were calculated from a small sampling of reflections.⁷⁷ A short pre-experiment was run, with both CuKα and MoKα radiation, from which an optimal data collection strategy was determined. All data for the reported crystal structures were ultimately collected with CuKα radiation, as the results of the pre-experiments indicated that using MoKα radiation did not offer any significant improvement in structure quality, but greatly increased the collection time. After the intensity data were corrected for absorption, the final cell constants were calculated from the full dataset from the xyz centroids of the strong reflections.⁷⁷ The structures were solved using SHELXT and refined using SHELXL.^{78,79} Full-matrix least squares/difference Fourier cycles were performed to assign the non-hydrogen atoms. All non-hydrogen atoms were first refined isotropically, followed by using anisotropic displacement parameters. All hydrogen atoms were placed in ideal positions and refined as riding atoms with relative isotropic displacement parameters. Additional refinement details are given in the SI, Section S5.

Computational methods

All DFT calculations were performed in the gas phase using ORCA 6.0.0.⁸⁰ The initial geometry for all complexes studied was taken from SCXRD structures. Solvent molecules and counter cations were removed. Relevant bond distances were compared to the experimental crystal structure, resulting in absolute average errors of 0.65% for Ce^{IV}-O bonds and -1.33% for M^{VI}-O bonds. All calculations presented here were performed using the hybrid exchange-correlation functional PBE0,⁸¹ and scalar relativistic corrections were added with the ZORA method as developed in ORCA.^{82,83} An all-electron basis set, SARC/ZORA-def2TZVP was used for all the atoms.⁸⁴ Fine tolerances on energy (10⁻¹¹), density matrix (10⁻⁸), and integrations were used. To account for solvent effects, the electronic structure was then obtained with the conductor-like polarizable continuum model (CPCM) using acetonitrile as the solvent.



Natural bond order (NBO) analyses were performed within the ORCA interface.⁸⁵ From the NBO, it was possible to obtain the metal atom contribution to the natural localized molecular orbitals (NLMOs). The obtained NBO Lewis structure and shape of the relevant orbitals were checked against various DFT functionals and basis sets (see SI) and were found to be consistent across methods. QTAIM analysis was performed using the Basin Analysis function within the program Multiwfn.^{86,87} Using the electron density, it was possible to extract the Delocalization Index (DI) value for the studied systems.

Results & discussion

Synthesis and characterization of Ce^{III} complexes

Villanneau and co-workers have previously reported the synthesis of (TBA)₃[Ce^{III}{Mo₅O₁₃(OMe)₄NO}₂] (**2-Ce(Mo₅)₂**) by refluxing **1-NaMo₅** with half an equivalent of Ce^{IV}(SO₄)₂.⁷¹ They verified the reaction also works when using Ce^{III}(NO₃)₂·6H₂O in place of Ce^{IV}(SO₄)₂, though no experimental description is provided.⁷¹ To gain familiarity with these Ce containing systems, we first attempted to repeat the synthesis of **2-Ce(Mo₅)₂** using a slightly modified procedure. A purple solution of **1-NaMo₅** in MeOH was added directly to a vial containing half an equivalent of Ce^{III}(OTf)₃ and the mixture was stirred at 50 °C for two hours. Cooling this solution to -30 °C induced the formation of red/violet crystals (see experimental for more

details). Characterization of these crystals using ¹H NMR spectroscopy (Fig. 1a, S1 and S2) reveals five resonances, four of which are assigned to the TBA cations (at 1–3.5 ppm), and an additional signal observed at 3.84 ppm assigned to the -OMe groups of **2-Ce(Mo₅)₂**. This signal is shifted in comparison to the corresponding resonance of the analogous Bi^{III} centered complex (*i.e.*, (TBA)₃[Bi^{III}{Mo₅O₁₃(OMe)₄NO}₂]; δ = 4.60 ppm in CD₃CN),⁷⁰ likely due to the influence of the paramagnetic Ce^{III} (f⁴) center.

We have recently reported that the use of ¹⁷O enriched **1-NaMo₅** in the synthesis of M(Mo₅)₂ sandwich-type complexes provides access to complexes which are readily characterized by ¹⁷O NMR spectroscopy. This was pursued for **2-Ce(Mo₅)₂** (see experimental section), with the ¹⁷O NMR spectrum of the obtained product shown in Fig. 1b and S7. The spectrum contains four resonances, consistent with the four oxygen environments. The signals associated with the bridging Mo–O–Mo groups (green) and the μ₅-O centers (magenta) are assigned based on literature precedent.^{68,69,88,89} Assignment of the Ce–O–Mo and terminal Mo=O groups is more difficult due to the small chemical shift between these two signals (*ca.* 7 ppm). We note that bridging M–O–Mo groups are typically observed upfield (*ca.* 400–700 ppm);^{88–91} the unusually high chemical shift of the oxygen nuclei present in the Ce–O–Mo bridges is likely a result of the influence of the paramagnetic Ce^{III} center.

In the interpretation of signals of NMR spectra that are paramagnetically broadened or shifted, the observed chemical shift (δ) can be decomposed into three components. As described in eqn (1), these are orbital shift (δ^{orb}), the Fermi contact shift (δ^{FC}), and the pseudocontact shift (δ^{PC}).^{92,93}

$$\delta = \delta^{\text{orb}} + \delta^{\text{FC}} + \delta^{\text{PC}} \quad (1)$$

The orbital shift can be thought of as the typical chemical shift in the corresponding diamagnetic compound and is approximately temperature independent. Together, the Fermi contact shift and pseudocontact shift, which represent through bond and through space interactions between the nuclear spins and the unpaired electrons of the paramagnetic center respectively, cause deviations in the observed chemical shifts of paramagnetic complexes from that of their diamagnetic counterparts. Importantly, the Fermi contact shift and pseudocontact shift are inversely proportional to distance from the paramagnetic center and to temperature.^{92–94} Considering these factors in the context of the ¹⁷O NMR spectrum **2-Ce(Mo₅)₂**, it is fair to assume that the oxygen nuclei of the Ce–O–Mo bridges will be more strongly affected by the paramagnetic Ce(III) center than the oxygen nuclei of Mo=O groups. Additionally, it is also expected that as temperature increases the contributions of δ^{FC} and δ^{PC} to the observed chemical shift should decrease, resulting in a spectrum that appears more similar to that of a diamagnetic complex. In the case of **2-Ce(Mo₅)₂**, this means that the peak corresponding to the Ce–O–Mo bridges should be much more sensitive to temperature than that of the Mo=O groups, moving upfield (*i.e.*, to a position more typical of M–O–Mo bridges) as temperature increases.

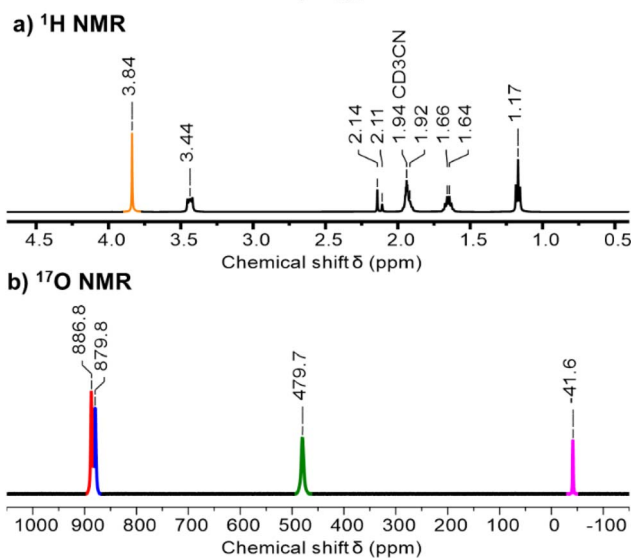
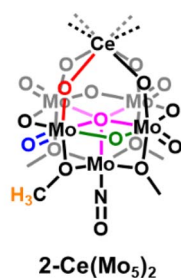


Fig. 1 ¹H NMR spectrum (a) and ¹⁷O NMR spectrum (b) of **2-Ce(Mo₅)₂**. Spectra were obtained in CD₃CN at room temperature.



To verify this, variable temperature (VT) ^{17}O NMR spectroscopy was performed on a sample of $2\text{-Ce}(\text{Mo}_5)_2$ in CD_3CN (Fig. S11). As expected, the signals observed at 886.6 ppm and 879.6 ppm in Fig. 1b (collected 19.4 °C) behave differently during VT experiments. The resonance that was originally observed at 886.8 ppm in Fig. 1b shows a strong temperature dependence, moving from 905.6 ppm at -17.5 °C to 863.1 ppm at 80.1 °C. This *ca.* 40 ppm shift upfield over the temperature range studied is consistent with the expected behavior of the oxygen nuclei of the Ce–O–Mo bridges. Conversely, the peak observed at 879.6 ppm in Fig. 1b is less sensitive to temperature, moving only *ca.* 3.5 ppm downfield as temperature is increased from -20 °C \rightarrow 80 °C. As such, we assign this signal to the Mo=O groups. The other oxygen nuclei present in $2\text{-Ce}(\text{Mo}_5)_2$ behave similarly to the Mo=O groups, showing downfield shifts of *ca.* 6 ppm and 36 ppm for the Mo–O–Mo and $\mu_5\text{-O}$ groups respectively. We note that the magnitude of these shifts increases as the distance of the respective oxygen nuclei to the Ce^{III} center decreases.

We next set out to extend the family of Ce sandwich-type complexes through the synthesis of tungsten-containing analogue $(\text{TBA})_3[\text{Ce}^{\text{III}}\{\text{W}_4\text{O}_{13}(\text{OMe})_4\text{MoNO}\}_2]$ ($2\text{-Ce}(\text{W}_4\text{Mo})_2$). Following the methods described above, $1\text{-NaW}_4\text{Mo}$ was treated with 0.55 eq. of $\text{Ce}^{\text{III}}(\text{OTf})_3$ in MeOH at 50 °C, resulting in the formation of a blue/purple solution. Cooling the solution to -30 °C led to the formation of blue/purple crystals. The crystals

were analyzed by ^1H NMR spectroscopy (Fig. 2a, S4 and S5). The spectrum is very similar to that of $2\text{-Ce}(\text{Mo}_5)_2$, with five signals in the correct ratio to be assigned to the expected resonances of the TBA cations and –OMe groups of the product. Interestingly, the signals assigned to the TBA cations of $2\text{-Ce}(\text{W}_4\text{Mo})_2$ are systematically shifted downfield compared to the corresponding resonances in the ^1H NMR spectrum of $2\text{-Ce}(\text{Mo}_5)_2$, while an upfield shift is observed for the –OMe signal (3.18 ppm for $2\text{-Ce}(\text{W}_4\text{Mo})_2$ vs. 3.84 ppm for $2\text{-Ce}(\text{Mo}_5)_2$). This observation suggests that the influence of the paramagnetic Ce^{III} center changes as a function of the framework metal present in the system.

^{17}O NMR spectroscopy was performed on ^{17}O enriched $2\text{-Ce}(\text{W}_4\text{Mo})_2$ (Fig. 2b and S9). The spectrum is very similar to that of $2\text{-Ce}(\text{Mo}_5)_2$, with four peaks observed that can be readily assigned to the oxygen nuclei of the $\{\text{W}_4\text{Mo}\}$ units which can undergo isotopic enrichment during the synthesis, these being the Ce–O–W bridges (red), the W–O–W bridges (green), the terminal W=O groups (blue), and the central $\mu_5\text{-O}$ nuclei (magenta) (Fig. 2b). There is less ambiguity in the assignment of the Ce–O–W groups vs. the W=O groups in $2\text{-Ce}(\text{W}_4\text{Mo})_2$ considering the larger chemical shift difference between the two peaks (*ca.* 26 ppm) in the ^{17}O NMR spectrum and the VT ^{17}O NMR study performed on $2\text{-Ce}(\text{Mo}_5)_2$. The major difference between the ^{17}O NMR spectra of $2\text{-Ce}(\text{W}_4\text{Mo})_2$ and $2\text{-Ce}(\text{Mo}_5)_2$ is the systematic shifting of all peaks upfield, with the effect more pronounced for the resonances observed at higher chemical shifts. This observation is common when comparing the ^{17}O NMR spectra of isostructural polyoxotungstates (POTs) and polyoxomolybdates (POMOs), and is attributed to the longer, more ionic, W–O bonds of these compounds.^{88,89}

Single crystals of $2\text{-Ce}(\text{W}_4\text{Mo})_2$ were grown by vapor diffusion of Et_2O into a saturated solution of the complex dissolved in MeOH. Analysis by single crystal X-ray diffraction (SCXRD) gave, after data refinement, the structure shown in Fig. 3. The structure of $2\text{-Ce}(\text{Mo}_5)_2$ was previously reported by Villanneau and co-workers, though this structure features an unusually long terminal nitrosyl bond of 1.35(4) Å. This is *ca.* 0.14–0.15 Å longer than the corresponding bonds in $2\text{-Ce}(\text{W}_4\text{Mo})_2$ and analogous M^{IV} centered complexes (*i.e.* $(\text{TBA})_2[\text{M}\{\text{M}'_4\text{O}_{13}(\text{OMe})_4\text{MoNO}\}_2]$ $\text{M} = \text{Zr}^{\text{IV}}, \text{Hf}^{\text{IV}}, \text{Th}^{\text{IV}}, \text{U}^{\text{IV}}, \text{M}' = \text{Mo}^{\text{VI}}$ or W^{VI}). We therefore re-acquired structural data for $2\text{-Ce}(\text{Mo}_5)_2$. The obtained structure is also shown in Fig. 3 and features terminal nitrosyl bond lengths that are the same, within error, as those in $2\text{-Ce}(\text{W}_4\text{Mo})_2$. In both structures, the cerium center occupies an approximately square antiprismatic coordination environment, with average Ce–O bonds lengths of *ca.* 2.48 Å. This illustrates that framework metal substitution has practically no influence on the local coordination environment at Ce. Other Ce^{III} centered sandwich-type polyoxometalate complexes present in the literature (*e.g.* $[\text{Ce}(\text{W}_5\text{O}_{18})_2]^{6-}$, $[\text{Ce}(\alpha\text{-PW}_{11}\text{O}_{39})_2]^{11-}$, *cis-* or *trans-* $[\text{Ce}(\alpha_2\text{-P}_2\text{W}_{17}\text{O}_{61})]^{17-}$) also feature a square antiprismatic coordination environment at Ce, with average Ce–O bond lengths of *ca.* 2.47–2.49 Å.^{53,54,95–98} These distances resemble those of $2\text{-Ce}(\text{Mo}_5)_2$ and $2\text{-Ce}(\text{W}_4\text{Mo})_2$ which, along with the presence of three TBA cations per sandwich complex, support the presence of trivalent cerium center.

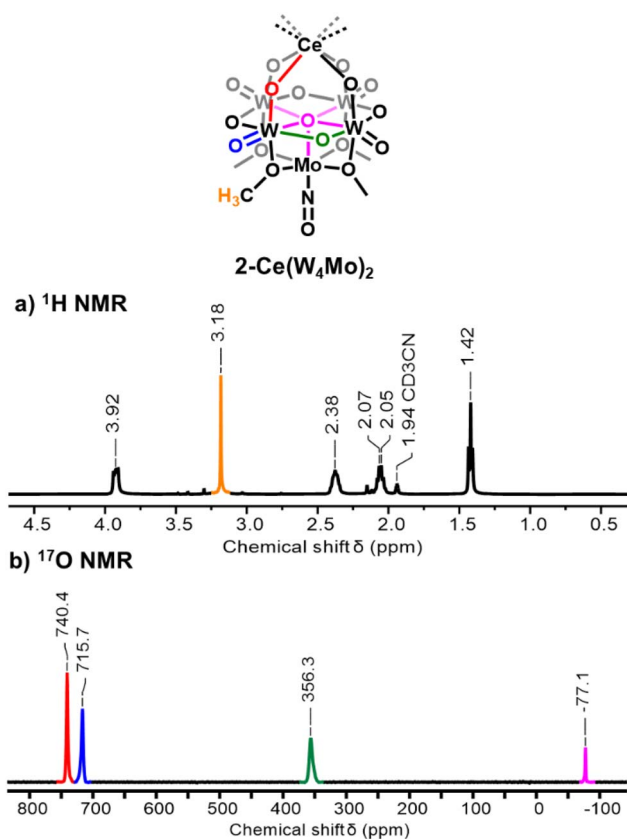


Fig. 2 ^1H NMR spectrum (a) and ^{17}O NMR spectrum (b) of $2\text{-Ce}(\text{W}_4\text{Mo})_2$. Spectra were obtained in CD_3CN at room temperature.



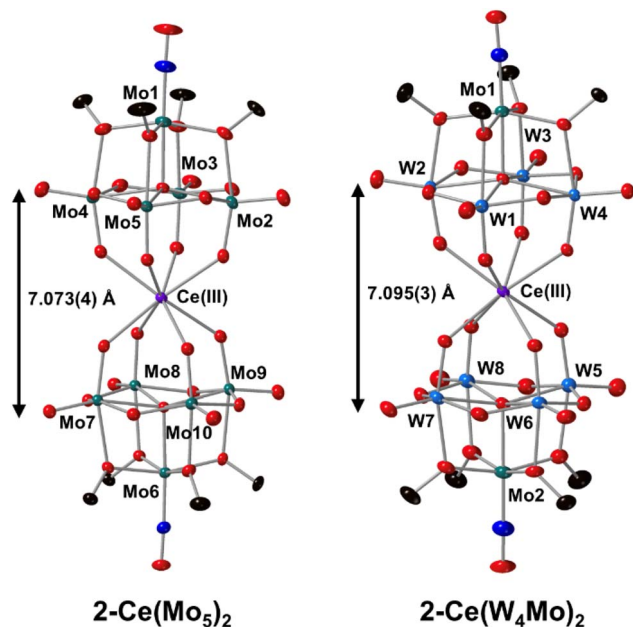


Fig. 3 SCXRD structures of $2\text{-Ce}(\text{Mo}_5)_2$ and $2\text{-Ce}(\text{W}_4\text{Mo})_2$ with probability ellipsoids set at 50%. The tetrabutylammonium cations, solvent molecules and some disorder has been masked for clarity.

With full structural characterization in hand, we turned our attention to examination of the electronic structure of the Ce(III)-centered sandwich-type complexes. We first recorded the electronic absorption spectra of $2\text{-Ce}(\text{Mo}_5)_2$ and $2\text{-Ce}(\text{W}_4\text{Mo})_2$, with the obtained spectra shown in Fig. S13 and S16 (and in Fig. 5, black lines). The main feature in both spectra is a broad low intensity transition centered at 552 nm ($\epsilon = 144 \text{ dm}^3 \text{ cm}^{-1}$) for $2\text{-Ce}(\text{Mo}_5)_2$ and 564 nm ($\epsilon = 129 \text{ dm}^3 \text{ cm}^{-1}$) for $2\text{-Ce}(\text{W}_4\text{Mo})_2$. This peak has been observed in 1-NaMo_5 , $1\text{-NaW}_4\text{Mo}$, and a number of the corresponding M^{II} , M^{III} , and M^{IV} sandwich-type complexes.^{68–71,76} We have previously shown this peak can be attributed to transitions originating from occupied orbitals localized on the $\{\text{Mo}-\text{NO}\}^4$ units.⁶⁸ However, in our previous work, we have shown that the energy of this transition is not sensitive to the framework metal, observing close to identical λ_{max} values for this transition in pairs of isostructural sandwich-type complexes.⁶⁸ Therefore, the 12 nm difference between the λ_{max} values of $2\text{-Ce}(\text{Mo}_5)_2$ and $2\text{-Ce}(\text{W}_4\text{Mo})_2$ is surprising. Both spectra also show an intense absorption below 400 nm. This intense absorption is often observed in the electronic absorption spectra of polyoxometalate compounds and is attributed to $\text{O}(2p) \rightarrow \text{M}(4d/5d)$ ($\text{M} = \text{Mo}$ or W) ligand-to-metal charge transfer (LMCT).^{68,99,100} This feature likely obscures resolution of characteristic Ce(III) $4f \rightarrow 5d$ transitions, which typically occur at *ca.* 300–400 nm.^{101–107}

The redox properties of the Ce^{III} centered complexes were assessed using cyclic voltammetry on $2\text{-Ce}(\text{Mo}_5)_2$ and $2\text{-Ce}(\text{W}_4\text{Mo})_2$ (Fig. 4). The CVs of both complexes possess multiple reduction events. The most obvious differences are the potentials of the first two reduction processes of $2\text{-Ce}(\text{Mo}_5)_2$ ($E_{1/2} = -1.44 \text{ V}$, $-1.59 \text{ V vs. Fc}^{+/0}$) compared to $2\text{-Ce}(\text{W}_4\text{Mo})_2$ ($E_{1/2} = -1.21 \text{ V}$, $-1.33 \text{ V vs. Fc}^{+/0}$) and the presence of an additional

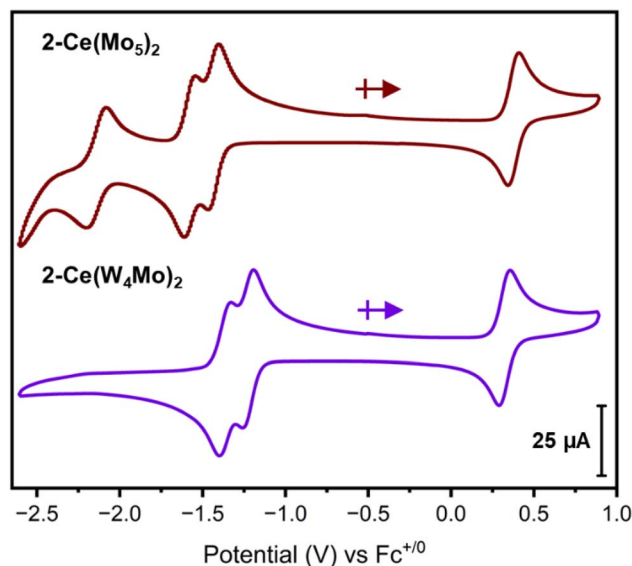


Fig. 4 Cyclic voltammograms of $2\text{-Ce}(\text{Mo}_5)_2$ (maroon) and $2\text{-Ce}(\text{W}_4\text{Mo})_2$ (purple). The data was acquired in MeCN with 0.1 M TBA(PF₆) supporting electrolyte, 1 mM of cluster, and a scan rate of 200 mV s^{-1} .

pseudo-reversible reduction event at $-2.18 \text{ V vs. Fc}^{+/0}$ in the CV of $2\text{-Ce}(\text{Mo}_5)_2$. The observation of a third reduction event in the CV of $2\text{-Ce}(\text{Mo}_5)_2$ suggests that a higher energy LUMO (*e.g.* LUMO+2) is more accessible for $2\text{-Ce}(\text{Mo}_5)_2$ than in $2\text{-Ce}(\text{W}_4\text{Mo})_2$.

When comparing these CV's to those of An^{IV} centered sandwich-type complexes with the general formula $(\text{TBA})_2[\text{M}'_4\text{O}_{13}(\text{OMe})_4\text{MoNO}]_2$ ($\text{M} = \text{Th}$, U , Np and $\text{M}' = \text{Mo}$ or W), both $2\text{-Ce}(\text{Mo}_5)_2$ and $2\text{-Ce}(\text{W}_4\text{Mo})_2$ have access to one less redox event than their An^{IV} counterparts and the observed redox events are shifted cathodically by *ca.* 0.5–0.75 V.^{68–70} These discrepancies are likely a result of the increased overall negative charge of complex, caused by incorporation of Ce^{III} vs. An^{IV} . Interestingly, the potentials of the 1st and 2nd reduction events are also closer together ($\Delta E = 0.15 \text{ V}$ and 0.12 V for $2\text{-Ce}(\text{Mo}_5)_2$ and $2\text{-Ce}(\text{W}_4\text{Mo})_2$) than the corresponding reduction events of the An^{IV} centered analogues ($\Delta E = 0.44\text{--}0.53 \text{ V}$ and 0.19 V for $\text{An}(\text{Mo}_5)_2$ and $\text{An}(\text{W}_4\text{Mo})_2$ systems, respectively). This may be a result of the increased distance between the two halves of the sandwich-type complexes in $2\text{-Ce}(\text{Mo}_5)_2$ ($\mu_5\text{-O} \rightarrow \mu_5\text{-O} = 7.073(4) \text{ \AA}$) and $2\text{-Ce}(\text{W}_4\text{Mo})_2$ ($\mu_5\text{-O} \rightarrow \mu_5\text{-O} = 7.095(3) \text{ \AA}$) compared to the $\text{An}(\text{Mo}_5)_2$ (average $\mu_5\text{-O} \rightarrow \mu_5\text{-O}$ *ca.* 6.87 \AA) and $\text{An}(\text{W}_4\text{Mo})_2$ (average $\mu_5\text{-O} \rightarrow \mu_5\text{-O}$ *ca.* 6.88 \AA) systems.^{68–70} If the 1st and 2nd reduction events correspond to sequential addition of an electron to each polyoxoalkoxide unit, then increasing the distance between these units should serve to electronically decouple the events, leading to smaller potential differences between the 1st and 2nd reduction processes. The higher sensitivity to this distance observed in the case of the all-molybdenum system can be rationalized by considering the nature of the LUMO/LUMO+1 in these sandwich-type complexes, as previously reported.⁶⁸ In all molybdenum complexes (like $2\text{-Ce}(\text{Mo}_5)_2$), these orbitals are localized on the



equatorial planes formed by the molybdenum centers either side of the central metal.⁶⁸ Given these orbitals are spatially close, changing the separation between the polyoxoalkoxide units is likely to drive large changes in the energy difference between the 1st and 2nd reduction events. Conversely, in tungsten-containing systems (like $2\text{-Ce}(\text{W}_4\text{Mo})_2$), the LUMO and LUMO+1 are localized on the peripheral Mo centers of the $\{\text{Mo}-\text{NO}\}_4$ units.⁶⁸ The large innate spatial separation between these orbitals caused by the change in electronic structure means these orbitals are already more electronically decoupled and therefore less sensitive to changes in the $\mu_5\text{-O} \rightarrow \mu_5\text{-O}$ distance.

The CVs both possess a chemically reversible $\text{Ce}^{\text{IV}}/\text{Ce}^{\text{III}}$ redox couple, occurring at 0.38 V vs. $\text{Fc}^{+/0}$ ($\Delta E_p = 65$ mV, $I_{p,a}/I_{p,c} = 1.08$) for $2\text{-Ce}(\text{Mo}_5)_2$ and 0.32 V vs. $\text{Fc}^{+/0}$ ($\Delta E_p = 65$ mV, $I_{p,a}/I_{p,c} = 1.07$) $2\text{-Ce}(\text{W}_4\text{Mo})_2$. Other $\text{Ce}^{\text{IV}}/\text{Ce}^{\text{III}}$ couples reported for $\text{Ce}(\text{POM})_2$ sandwich-type complexes range from 0.28 V to 0.86 V vs. SCE (saturated calomel electrode), however direct comparison of these values to those of $2\text{-Ce}(\text{Mo}_5)_2$ and $2\text{-Ce}(\text{W}_4\text{Mo})_2$ is difficult given the different reference electrode and the fact all of these values are measured in water (typically 0.1 M KCl or buffered pH 4.5 solutions).^{5,55,108,109} Comparing the potentials of the $\text{Ce}^{\text{IV}}/\text{Ce}^{\text{III}}$ redox couples of $2\text{-Ce}(\text{Mo}_5)_2$ and $2\text{-Ce}(\text{W}_4\text{Mo})_2$ to those of other Ce complexes supported by organic (anionic) oxygen bearing ligands where the redox couples were reported in MeCN shows the values for $2\text{-Ce}(\text{Mo}_5)_2$ and $2\text{-Ce}(\text{W}_4\text{Mo})_2$ are more positive. For these organic complexes, $\text{Ce}^{\text{IV}}/\text{Ce}^{\text{III}}$ redox couples can occur anywhere from -0.56 V vs. $\text{Fc}^{+/0}$ to -1.18 V vs. $\text{Fc}^{+/0}$.^{5,110,111} This suggests that polyanionic organic ligands are better at stabilizing Ce^{IV} than the polyoxoalkoxide clusters present in $2\text{-Ce}(\text{Mo}_5)_2$ and $2\text{-Ce}(\text{W}_4\text{Mo})_2$. However, the observed reversibility of the $\text{Ce}^{\text{IV}}/\text{Ce}^{\text{III}}$ redox couples and the stability of the Ce^{IV} complexes (see below) suggest $2\text{-Ce}(\text{Mo}_5)_2$ and $2\text{-Ce}(\text{W}_4\text{Mo})_2$ could potentially be applied as redox mediators or electrocatalysts in the future.^{18,112–114}

One-electron oxidation of $2\text{-Ce}(\text{Mo}_5)_2$ and $2\text{-Ce}(\text{W}_4\text{Mo})_2$

Given the CVs of $2\text{-Ce}(\text{Mo}_5)_2$ and $2\text{-Ce}(\text{W}_4\text{Mo})_2$ possess a reversible $\text{Ce}^{\text{IV}}/\text{Ce}^{\text{III}}$ redox couple, we sought to investigate one electron oxidation of these complexes to give access to the corresponding Ce^{IV} centered sandwich-type assemblies. Initially, we attempted electrochemical oxidation. Performing bulk oxidation on 1 mM solutions of $2\text{-Ce}(\text{Mo}_5)_2$ or $2\text{-Ce}(\text{W}_4\text{Mo})_2$ in MeCN/0.1 M TBA(PF_6) at ca. 0.7 V vs. $\text{Fc}^{+/0}$ leads to a color change, with the red/violet and blue/purple solutions of $2\text{-Ce}(\text{Mo}_5)_2$ and $2\text{-Ce}(\text{W}_4\text{Mo})_2$ turning orange and yellow respectively. Inspections the CVs of the solutions after bulk oxidation (Fig. S24 and S25) shows they are almost identical to those of pristine $2\text{-Ce}(\text{Mo}_5)_2$ and $2\text{-Ce}(\text{W}_4\text{Mo})_2$, suggesting the complex is stable under these conditions. The only major difference is that the position of the open circuit potential has moved to 0.47 V and 0.42 V respectively, suggesting successful formation of the target Ce^{IV} centered complexes, referred to as $3\text{-Ce}(\text{Mo}_5)_2$ and $3\text{-Ce}(\text{W}_4\text{Mo})_2$. The changes in color observed upon oxidation lead to changes in the electronic absorption spectra of the materials, as shown in Fig. 5a and b (red lines). The broad peak centered at 552 nm in the electronic absorption spectrum of $2\text{-Ce}(\text{Mo}_5)_2$

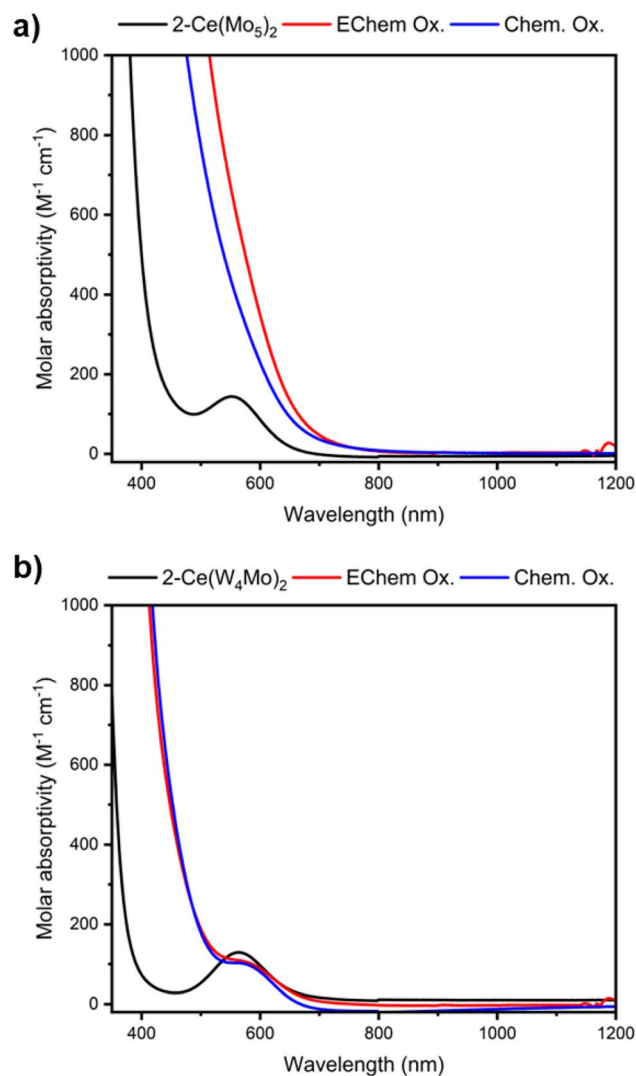


Fig. 5 UV-vis spectra of pristine $2\text{-Ce}(\text{Mo}_5)_2$ (a, black) and $2\text{-Ce}(\text{W}_4\text{Mo})_2$ (b, black) compared to those obtained after electrochemical oxidation (red) or chemical oxidation by tris(4-bromophenyl) ammoniumyl hexachloroantimonate (blue). All spectra were recorded in MeCN at room temperature.

$\text{Ce}(\text{Mo}_5)_2$ is obscured in by a new charge transfer process that onsets at ca. 700 nm. Similarly, the peak that was present in the electronic absorption spectrum of $2\text{-Ce}(\text{W}_4\text{Mo})_2$ (564 nm) appears to be slightly red shifted in the spectrum $3\text{-Ce}(\text{W}_4\text{Mo})_2$ and is now a shoulder on the side of a new intense absorption feature. These intense absorptions are likely caused by a LMCT process between filled orbitals of the polyoxoalkoxide ligands and the new, low-lying 4f based LUMO centered on Ce.

To confirm this, TD-DFT calculations were performed on $3\text{-Ce}(\text{Mo}_5)_2$ and $3\text{-Ce}(\text{W}_4\text{Mo})_2$ after structural optimization. For $3\text{-Ce}(\text{Mo}_5)_2$, the simulated electronic absorption spectrum shows intense absorption below 500 nm which is caused primarily by two distinct types of transitions (Fig. S26). The dominant contribution is confirmed to be a LMCT process involving transitions from occupied molecular orbitals localized on the POM cage (e.g. HOMO-7) to unoccupied Ce^{IV} 4f



orbitals (Fig. S27). The secondary contribution involves transitions from orbitals localized on the $\{\text{Mo}-\text{NO}\}^4$ unit (e.g. HOMO-1) to vacant orbitals largely localized on the other Mo centers (Fig. S28). For the $3\text{-Ce}(\text{W}_4\text{Mo})_2$, the primary contribution is also a LMCT to unoccupied Ce^{IV} 4f orbitals but originates from occupied orbitals more delocalized across the whole polyoxoalkoxide cage (Fig. S32 and S33). This transition occurs at higher energy than in the all-molybdenum system explaining the difference in the onset of the intense absorption observed in the UV-vis spectra of $3\text{-Ce}(\text{Mo}_5)_2$ and $3\text{-Ce}(\text{W}_4\text{Mo})_2$. Both simulated spectra also feature less intense absorptions at ca. 600 nm (Fig. S29 and S34). These are readily assigned to LMCT between filled orbitals localized on the $\{\text{Mo}-\text{NO}\}^4$ units and empty Ce^{IV} 4f orbitals (Fig. S30 and S35). This is likely the cause for shoulder observed at ca. 568 nm on the side of the intense absorption present in the electronic absorption spectrum of $3\text{-Ce}(\text{W}_4\text{Mo})_2$ (Fig. 5b). A separate feature is not seen in the electronic absorption spectrum $3\text{-Ce}(\text{Mo}_5)_2$ (Fig. 5a), as it is likely obscured by the more intense LMCT process.

We next pursued isolation of the Ce^{IV} centered complexes. Treatment of either $2\text{-Ce}(\text{Mo}_5)_2$ and $2\text{-Ce}(\text{W}_4\text{Mo})_2$ with 1.1 equivalents of tris(4-bromophenyl)ammoniumyl hexachloroantimonate in MeCN ($E_{\text{Ox}} = 0.67 \text{ V vs. Fc}^{+/0}$ in MeCN)

leads to an immediate color change.¹¹⁵ Following work-up, orange and yellow solutions of $3\text{-Ce}(\text{Mo}_5)_2$ and $3\text{-Ce}(\text{W}_4\text{Mo})_2$, respectively, are obtained. Characterization of the crude products by ^1H NMR spectroscopy result in the spectra shown in Fig. 6 (Fig. S3 and S6). The spectra of $3\text{-Ce}(\text{Mo}_5)_2$ and $3\text{-Ce}(\text{W}_4\text{Mo})_2$ are much more similar than the ^1H NMR spectra of the respective starting materials, with the peaks associated with the TBA cations (ca. 1–3.5 ppm) appearing in almost identical positions. The signals associated with the $-\text{OMe}$ groups of the sandwich-type complexes are observed at 4.73 ppm and 4.87 ppm respectively for $3\text{-Ce}(\text{Mo}_5)_2$ and $3\text{-Ce}(\text{W}_4\text{Mo})_2$. These resonances are now at identical positions to the corresponding peaks in the series of closed-shell M^{IV} centered sandwich-type complexes with the general formula $(\text{TBA})_2[\text{M}'_4\text{O}_{13}(\text{OMe})_4\text{MoNO}]_2$ ($\text{M} = \text{Zr}, \text{Hf}, \text{Th}$ and $\text{M}' = \text{Mo}$ or W), supporting successful formation of the desired Ce^{IV} sandwich-type complexes.^{68,70} This is further supported by electronic absorption spectroscopy, where spectra of $3\text{-Ce}(\text{Mo}_5)_2$ and $3\text{-Ce}(\text{W}_4\text{Mo})_2$ obtained from chemical oxidation were almost identical to those obtained from electrochemical oxidation experiments (Fig. 5a and b, blue lines). The Ce^{IV} complexes are relatively stable, with a half-life of more than one week when left in solution under inert atmosphere (Fig. S12).

To confirm the structures of $3\text{-Ce}(\text{Mo}_5)_2$ and $3\text{-Ce}(\text{W}_4\text{Mo})_2$, single crystals were grown by vapor diffusion of Et_2O in to saturated solutions of the complexes dissolved in MeCN. The obtained structures are shown in Fig. 7. The sandwich-type complexes are left mostly unchanged upon oxidation, with the bond distances within the polyoxoalkoxide ligands showing

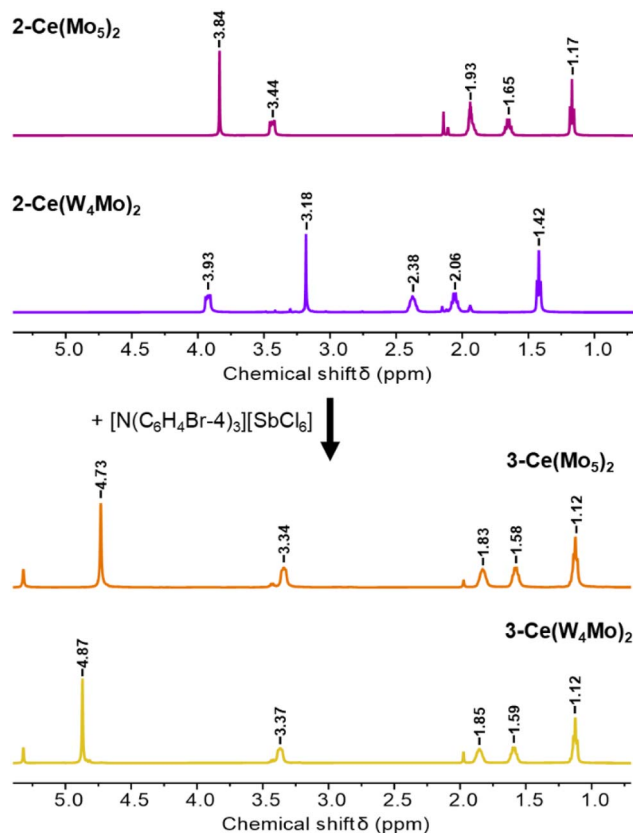


Fig. 6 ^1H NMR spectra following the one electron oxidation of $2\text{-Ce}(\text{Mo}_5)_2$ and $2\text{-Ce}(\text{W}_4\text{Mo})_2$ by tris(4-bromophenyl)ammoniumyl hexachloroantimonate to give the Ce^{IV} centered complexes $3\text{-Ce}(\text{Mo}_5)_2$ and $3\text{-Ce}(\text{W}_4\text{Mo})_2$. Spectra of the Ce^{III} complexes were obtained in CD_3CN , while the Ce^{IV} complexes were obtained in CD_2Cl_2 . All spectra were recorded at room temperature.

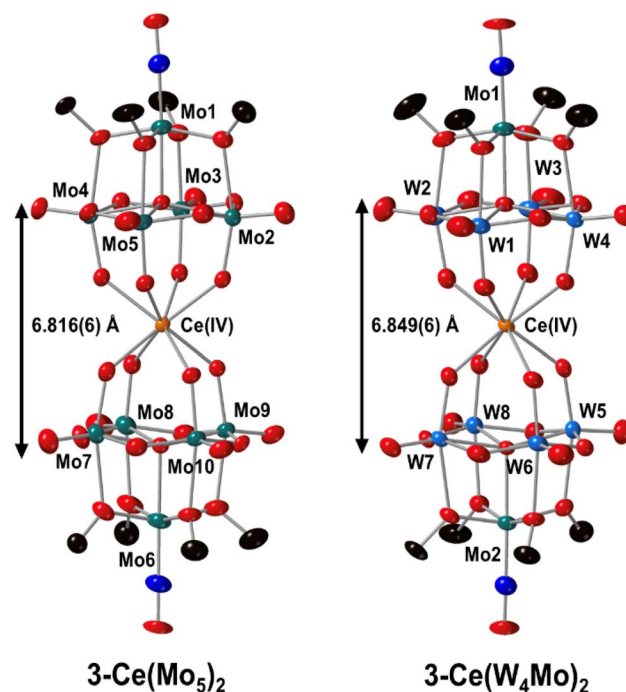


Fig. 7 SCXRD structures of the Ce^{IV} centered complexes $3\text{-Ce}(\text{Mo}_5)_2$ and $3\text{-Ce}(\text{W}_4\text{Mo})_2$ with probability ellipsoids set at 50%. The tetrabutylammonium cations, solvent molecules, and some disorder has been masked for clarity.



only very minor variations (Table S5). As expected, the main consequence of oxidation of Ce^{III} to Ce^{IV} is a contraction of the Ce–O bonds, with average Ce–O bond distances of *ca.* 2.34 Å and 2.35 Å respectively for $3\text{-Ce}(\text{Mo}_5)_2$ and $3\text{-Ce}(\text{W}_4\text{Mo})_2$. The 0.13–0.14 Å decrease in bond lengths compared to $2\text{-Ce}(\text{Mo}_5)_2$ and $2\text{-Ce}(\text{W}_4\text{Mo})_2$ can be attributed to the decrease in ionic radius accompanied with oxidation of Ce^{III} (1.14 Å) to Ce^{IV} (0.97 Å).¹¹⁶ This Ce–O bond contraction translates to a drop in the distance between the two polyoxoalkoxide units of the sandwich-type complex (approximated by the $\mu_5\text{-O} \rightarrow \mu_5\text{-O}$ distance). To quantify the deviation from square antiprismatic geometry, continuous shape measurements (CShM) were calculated (Table S6).^{117,118} Both $3\text{-Ce}(\text{Mo}_5)_2$ and $3\text{-Ce}(\text{W}_4\text{Mo})_2$ display nearly ideal square antiprismatic about the Ce centers (CShM values = 0.20960 and 0.22048, where a value of 0 reflects the ideal geometry). These values are lower than the corresponding values for $3\text{-Ce}(\text{Mo}_5)_2$ and $3\text{-Ce}(\text{W}_4\text{Mo})_2$ (CShM values = 0.93012 and 0.42435), illustrating how oxidation of the Ce center influences the overall geometry of the complexes.

¹⁷O NMR spectroscopy and M–O bond covalency in M^{IV} centered sandwich-type complexes

After successful isolation the Ce^{IV} centered sandwich-type complexes, $3\text{-Ce}(\text{Mo}_5)_2$ and $3\text{-Ce}(\text{W}_4\text{Mo})_2$, we sought to isolate ¹⁷O enriched analogues of the complexes to verify how the ¹⁷O spectra of these complexes change upon oxidation and how they compare to the spectra of other isostructural M^{IV} centered sandwich-type complexes.^{68,69} Following the procedures discussed above, ¹⁷O enriched samples of $2\text{-Ce}(\text{Mo}_5)_2$ and $2\text{-$

$\text{Ce}(\text{W}_4\text{Mo})_2$ were oxidized by one-electron, purified, and their ¹⁷O NMR spectra were recorded. The obtained spectra are shown in Fig. 8a and b, along with the corresponding spectra of other isostructural, diamagnetic, M^{IV} centered sandwich-type complexes. The peaks assigned to the terminal M = O groups (blue), bridging M–O–M groups (green), and central $\mu_5\text{-O}$ groups (magenta) are all shifted downfield in the ¹⁷O NMR spectra of $3\text{-Ce}(\text{Mo}_5)_2$ and $3\text{-Ce}(\text{W}_4\text{Mo})_2$ compared to the corresponding signals in the spectra of $2\text{-Ce}(\text{Mo}_5)_2$ and $2\text{-Ce}(\text{W}_4\text{Mo})_2$ ($\text{M} = \text{Mo}$ or W , Fig. 1b and 2b). The deshielding of these nuclei can be attributed to the reduction in the anionic charge of the system which accompanies the oxidation of Ce^{III} to Ce^{IV} . The only nuclei that do not follow this logical behavior are the Ce–O–M groups (red). The resonances assigned to these nuclei instead shift upfield. This is likely because oxidation of Ce^{III} (f^1) to Ce^{IV} (f^0) allows these peaks, which were observed at unusually high chemical shifts in the ¹⁷O NMR spectra of $2\text{-Ce}(\text{Mo}_5)_2$ and $2\text{-Ce}(\text{W}_4\text{Mo})_2$ due to interactions with the paramagnetic Ce^{III} center, to move back to positions more typical of bridging oxygen nuclei.^{88–91}

When comparing the ¹⁷O NMR spectra of $3\text{-Ce}(\text{Mo}_5)_2$ and $3\text{-Ce}(\text{W}_4\text{Mo})_2$ with those of other M^{IV} sandwich-type complexes (Fig. 8a and b, $\text{M}^{\text{IV}} = \text{Zr}, \text{Hf}, \text{Th}$)^{68,69} we can see that the positions of the peaks associated with the terminal M = O groups (blue), the bridging M–O–M groups (green), and the central $\mu_5\text{-O}$ groups (magenta) are insensitive to the identity of the M^{IV} ion present at the center of the sandwich-type complexes. Indeed, these peaks all occur within *ca.* 10 ppm of each other within each series of sandwich-type complexes (*i.e.* $\text{M}^{\text{IV}}(\text{Mo}_5)_2$ or $\text{M}^{\text{IV}}(\text{W}_4\text{Mo})_2$). As may be expected, this contrasts the behavior of

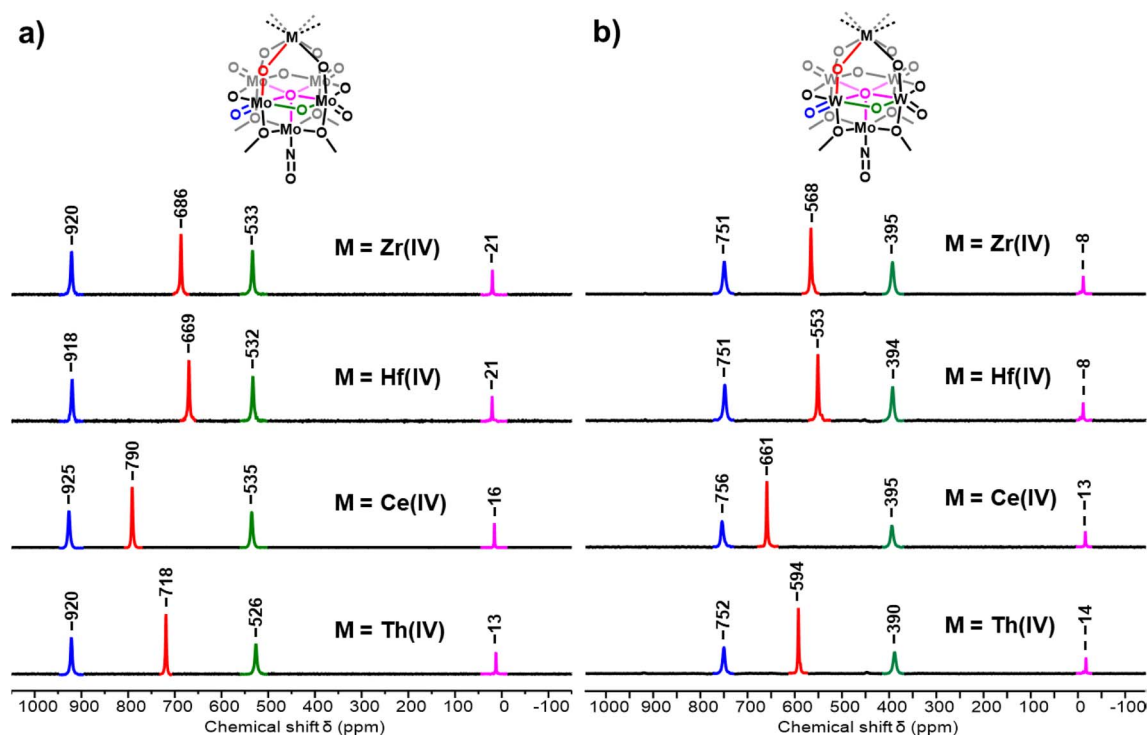


Fig. 8 ¹⁷O NMR spectrum of (a) $3\text{-Ce}(\text{Mo}_5)_2$ and (b) $3\text{-Ce}(\text{W}_4\text{Mo})_2$ compared to other isostructural M^{IV} centered sandwich-type complexes. All spectra were obtained in CD_2Cl_2 at room temperature.



the M^{IV} -O-M groups (red), with the peak assigned to these nuclei varying by >100 ppm across the series. In both series of complexes (*i.e.* $M^{IV}(\text{Mo}_5)_2$ or $M^{IV}(\text{W}_4\text{Mo})_2$), the magnitude of the observed chemical shift increases in the order $\text{Hf-O-M} < \text{Zr-O-M} < \text{Th-O-M} < \text{Ce-O-M}$ ($M = \text{Mo}$ or W). Given this trend persists between the series, it is likely that this relative ordering of the complexes is caused by differences in the M^{IV} -O bonding, while the 110–130 ppm difference in chemical shift between the signals of the pairs of $M^{IV}(\text{Mo}_5)_2$ and $M^{IV}(\text{W}_4\text{Mo})_2$ complexes can be attributed to differences in the Mo^{VI} -O and W^{VI} -O bonds.

Given the nature of both the M^{IV} -O bonds and M^{VI} -O bonds present in the sandwich-type complexes appear to strongly influence the experimentally observed ^{17}O NMR chemical shifts, we sought to gain further insights into these bonding interactions using DFT calculations. Optimized structures of the M^{IV} centered sandwich complexes discussed in Fig. 8 were first obtained and frequency calculations were performed to ensure the structures represent a minimum on their respective potential energy surfaces. Natural Bond Order (NBO) analysis

was then used to obtain metal atom contributions to the natural localized molecular orbitals (NLMOs) (Tables S12–15), specifically focusing on the nature of the M^{IV} -O- M^{VI} bonds ($M^{IV} = \text{Zr}, \text{Hf}, \text{Ce},$ or Th and $M^{VI} = \text{Mo}$ or W). The M^{IV} -O bonds in our systems are defined by two NLMOs, one σ -type and one π -type (Fig. 9a and b). Both NLMOs are dominated by the s and p orbitals of oxygen, with all metal orbitals contributing <15% in all cases. Of the M^{IV} centers present, Ce^{IV} orbitals (primarily f and d orbitals) contribute more to the NLMOs than any other M^{IV} , followed by Th , then Zr , and finally Hf (full details in Fig. S36–S39). This is largely in-line with the results obtained for $[\text{M}^{IV}\text{Cl}_6]^{2-}$ ($M^{IV} = \text{Zr}, \text{Hf},$ and Ce) complexes, where Ce^{IV} contributes much more to M-Cl bonds than Zr^{IV} or Hf^{IV} due to increased f -orbital contribution.^{21,24} Moreover, this follows exactly the order of ^{17}O NMR chemical shifts of the M^{IV} -O- M^{VI} nuclei presented in Fig. 8, suggesting a correlation between the M^{IV} orbital contribution to M^{IV} -O bonds (*i.e.* M-O bond covalency) and the observed chemical shifts.

Investigation of the specific contributions to the NLMOs of the Ce^{IV} -O bonds highlights that the d -orbital contributions are similar to those of the Th^{IV} -O bonds (see Fig. S40 and Table S17), and thus the overall increase in metal orbital contributions to the Ce^{IV} -O bonds is result of increased f -orbital contributions. This finding is consistent with previous studies on f -element-oxide systems, which establish that favorable energy matching between metal- f and oxygen- p orbitals can be a more dominant factor in determining f -orbital contribution to bonding than what the greater spatial extent of orbitals in heavier actinides might imply.^{119,120} The electronic origin of this trend was elucidated by analyzing the Projected Density of States (PDOS). While the highly delocalized electronic structure of the POM framework complicates the analysis of individual canonical molecular orbitals, the PDOS provides a clear picture. Fig. S41 shows that the unoccupied Ce $4f$ states are energetically closer to the O $2p$ -dominated valence band than the Th $5f$ states (Fig. S42). This favorable energy matching for Ce^{IV} promotes more effective orbital mixing, thus explaining the greater f -orbital character observed in the NBO analysis. In contrast, the d -orbital contributions are more comparable for both metals, highlighting that the f -orbitals are the primary differentiators of M-O covalency in these two systems.

This interpretation aligns with the work of Minasian *et al.* on simple lanthanide dioxides, which demonstrated that an 8-coordinate environment is crucial for enabling f -orbital participation in bonding.³¹ While the perfect cubic symmetry in LnO_2 allows for a clear analysis of specific symmetry-allowed orbital interactions, the highly delocalized valence molecular orbitals and slightly distorted structures (*i.e.* not perfectly D_{4d}) present in our study make it difficult to identify the specific MOs with $\text{O}(2p)$ character that are responsible for symmetry allowed overlap with the metal d/f orbitals. Even if the complex nature of the POM cage orbitals makes it difficult to isolate specific symmetry allowed overlap effects, a qualitative agreement with Minasian *et al.*, supported by NBO analysis and QTAIM DI delocalization indices (see below) provides insight into the importance of 8-coordinate geometry for M-O orbital mixing, and the specific nature of the metal.

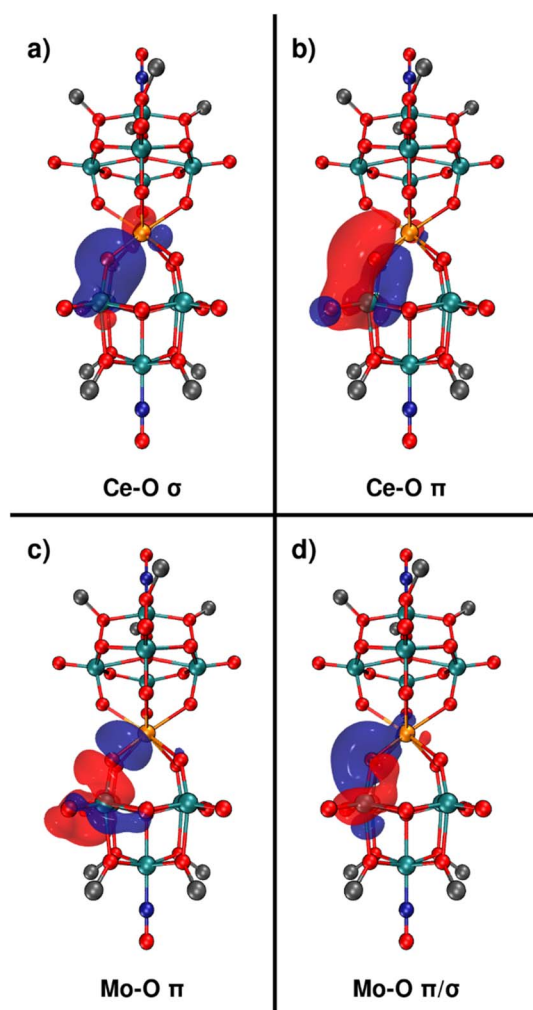


Fig. 9 NLMOs which define the Ce^{IV} -O bonds (a and b) and Mo^{VI} -O bonds (c and d) of the Ce^{IV} -O- Mo^{VI} bridges present in $3\text{-Ce}(\text{Mo}_5)_2$. Similar graphics for all other complexes discussed in Fig. 8 can be found in Fig. S43–49.



The M^{VI} -O bonds ($M^{VI} = Mo$ or W) of the M^{IV} -O- M^{VI} bridges are similarly defined by two NLMOs, one π -type and one σ/π hybrid (Fig. 9c and d). Again, these orbitals have primarily O character with minor contributions from the M^{VI} center. These orbitals remain consistent across the series of complexes studied, with Mo^{VI} orbitals always contributing more to the NLMOs than W^{VI} orbitals, regardless of the M^{IV} center present. This could explain the fact that ^{17}O NMR chemical shifts of the oxygen nuclei of the M^{IV} -O- M^{VI} bridges in the $M^{IV}(Mo_5)_2$ complexes are 110–130 ppm higher than in the corresponding signal in the ^{17}O NMR spectra of the $M^{IV}(W_4Mo)_2$ complexes, suggesting that an overall increase in metal orbital contribution to the bonds of the M^{IV} -O- M^{VI} bridges drives an increase in the observed ^{17}O NMR chemical shift.

Though the intuitive understanding of orbital mixing is provided by NBO analysis, other metrics have proved effective in the literature for assessing bonding.^{121–123} Specifically, delocalization indices (DI) obtained from Quantum Theory of Atoms in Molecules (QTAIM) analyses, which provide a measure of the number of electron pairs exchanged in an interaction, have been reported as an effective measure of metal bond covalency.^{23,25,27,33,122,124,125} The DI values for the atoms involved in both the M^{IV} -O and M^{VI} -O bonds of the M^{IV} -O- M^{VI} bridges present in the series of complexes studied are given in Table S16. Focusing on the M^{IV} -O bonds, plotting DI vs. ^{17}O NMR chemical shift of the oxygen nuclei present in the M^{IV} -O- M^{VI} bridges gives the graph shown in Fig. 10. It is immediately apparent that as DI increases, which implies an increase in bond covalency, the observed ^{17}O NMR chemical shift also increases, with very good linear fits observed. The observed deshielding of the ^{17}O nuclei as M-O bond covalency increases can be attributed to Spin-Orbit Heavy Atom on the Light Atom (SO-HALA) effects.^{42,120,126–132} Specifically, the presence of empty low-lying valence d- or f-orbitals allows the deshielding $\sigma_{HA-LA}^* \leftrightarrow \eta_{HA}^*$ coupling mechanism to dominate. This leads to relatively small deshielding effects in $4d^0/5d^0$ transition metals,

but the magnitude increases drastically when moving to $4f^0$ Ce^{IV} .¹²⁰ While even larger effects are expected for $5f^0$ systems, moderate deshielding effects (comparable to $5d$ systems) are often reported for Th^{IV} as bonding is typically dominated by $6d/7s$ orbitals with only minor contributions of $5f$ orbitals.¹²⁰ This is in line with the NBO and PDOS analysis presented above and thus the relationship between ^{17}O NMR chemical shift and M^{IV} -O bond covalency shown in Fig. 10 reproduces reported trends in the magnitude of SO-HALA effects well.

The vertical offset between the two series can be attributed to difference in covalency of the M^{VI} -O bonds of the M^{IV} -O- M^{VI} bridges. On average, DI values of 1.30 for Mo^{VI} -O bonds and 1.24 for W^{VI} -O bonds were obtained. This is in line with literature precedent that W^{VI} -O bonds are typically more ionic than Mo^{VI} -O bonds.^{88,89} Plotting the sum of the DIs of the M^{IV} -O and M^{VI} -O bonds vs. ^{17}O NMR chemical shift of the oxygen nuclei present in the M^{IV} -O- M^{VI} bridges (Fig. S50) allows the two series (*i.e.* $M^{IV}(Mo_5)_2$ and $M^{IV}(W_4Mo)_2$ complexes) to be treated collectively. A strong correlation between the total DI and ^{17}O NMR chemical shift is still observed, however there are some anomalies which suggest that this simplistic approach cannot completely capture the influence that varying the framework metal has on metal-oxygen bonding. The bond covalency trends obtained from our combined experimental and computational analysis are in line with the computational studies from Kaltsoyannis and co-workers, who assessed M-O bond covalency in a series of $M(OC_6H_5)_4$ ($M = Ti^{IV}, Zr^{IV}, Hf^{IV}, Ce^{IV}, Th^{IV}, Pa^{IV}, U^{IV},$ and Np^{IV}), and the experimental study from Schelter, who observed $Ce^{IV}=N$ bonds are more covalent than $Th^{IV}=N$ bonds.^{33,34,123}

Conclusions

In conclusion, we have built on the original methods of Proust and Villanneau to access the Ce^{III} centered sandwich-type complexes **2-Ce(Mo₅)₂** and **2-Ce(W₄Mo)₂**.⁷¹ Cyclic voltammetry revealed the complexes possess a reversible Ce^{IV}/Ce^{III} redox couples at *ca.* 0.35 V vs. $Fc^{+/0}$ in MeCN. One-electron chemical oxidation using tris(4-bromophenyl)ammoniumyl hexachloroantimonate provides convenient access to the corresponding Ce^{IV} complexes **3-Ce(Mo₅)₂** and **3-Ce(W₄Mo)₂**. While the optical properties of **2-Ce(Mo₅)₂** and **2-Ce(W₄Mo)₂** are very similar to the reported sodium complexes **1-NaMo₅** and **1-NaW₄Mo**,^{68,70,71,76} oxidation to Ce^{IV} leads to the emergence of a new charge transfer process assigned to a LMCT between filled orbitals of the polyoxoalkoxide ligands and the newly empty $4f$ orbitals of Ce^{IV} . ^{17}O enriched analogues of the complexes were readily prepared from the corresponding ^{17}O enriched starting materials. While analysis of the ^{17}O NMR spectra of the Ce^{III} centered complexes shows the influence of the paramagnetic Ce^{III} ($4f^1$) center, the ^{17}O NMR spectra of the diamagnetic Ce^{IV} centered complexes are in line with the spectra of the previously prepared Zr^{IV} , Hf^{IV} , and Th^{IV} containing complexes.^{68,69} The only major deviation in the spectra of the Ce^{IV} complexes is the high chemical shift of the peaks assigned to the oxygen nuclei of the Ce -O- M^{VI} bridges ($M^{VI} = Mo$ or W). Computational calculations clearly indicate that this increase in chemical shift can be attributed to an increase in the M^{IV} -O bond covalency. Further

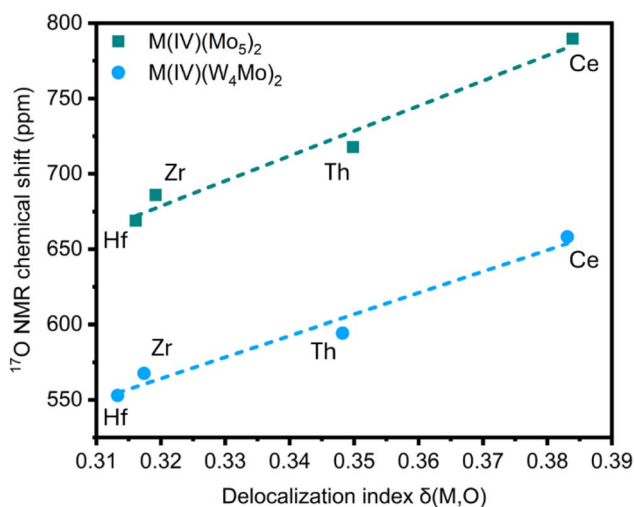


Fig. 10 Plot of M^{IV} -O delocalization index obtained from QTAIM analysis vs. ^{17}O NMR chemical shift (ppm) for the complexes discussed in this work.



investigations revealed a strong correlation between ^{17}O NMR chemical shift and M–O delocalization indices (which act as measure of bond covalency) obtained from QTAIM analysis, in this series of isostructural diamagnetic complexes. These results display the utility of ^{17}O NMR spectroscopy as an experimental tool for investigating metal–oxygen bonding and covalency.

Author contributions

E. M. M. and D. S. conceived of the project. D. S., N. G., and A. C. B. synthesized and characterized all compounds. W. W. B. determined the crystal structures and carried out elemental analysis measurements. Calculations were performed by M. P. and M. T. R., while E. M. M. directed the project. The manuscript was written through contributions of all authors. All authors have given approval to the final version of the manuscript.

Conflicts of interest

The authors declare no competing financial interest.

Data availability

The data supporting this article, including the coordinates of the optimized geometries of the structures discussed in this work obtained in computational studies, have been included as part of the supplementary information (SI). All other data is available upon requests made to the corresponding author.

CCDC 2481875–2481878 contain the supplementary crystallographic data for this paper.^{133a–d}

Supplementary information: additional spectroscopic, crystallographic, and computational data (PDF). See DOI: <https://doi.org/10.1039/d5sc06415e>.

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References

- R. J. Ellis and M. R. Antonio, *ChemPlusChem*, 2012, **77**, 41–47.
- M.-H. Chiang, C. W. Williams, L. Soderholm and M. R. Antonio, *Eur. J. Inorg. Chem.*, 2003, **2003**, 2663–2669.
- Y. Wei, M. Kumagai, Y. Takashima, M. Asou, T. Namba, K. Suzuki, A. Maekawa and S. Ohe, *J. Nucl. Sci. Technol.*, 1998, **35**, 357–364.
- T. A. Pham, A. B. Altman, S. C. E. Stieber, C. H. Booth, S. A. Kozimor, W. W. Lukens, D. T. Olive, T. Tyliczszak, J. Wang, S. G. Minasian and K. N. Raymond, *Inorg. Chem.*, 2016, **55**, 9989–10002.
- N. A. Piro, J. R. Robinson, P. J. Walsh and E. J. Schelter, *Coord. Chem. Rev.*, 2014, **260**, 21–36.
- A. C. Boggiano, C. M. Studvick, S. Roy Chowdhury, J. E. Niklas, H. Tateyama, H. Wu, J. E. Leisen, F. Kleemiss, B. Vlasisavljevich, I. A. Popov and H. S. La Pierre, *Nat. Chem.*, 2025, **17**, 1005–1010.
- N. T. Rice, I. A. Popov, D. R. Russo, J. Bacsá, E. R. Batista, P. Yang, J. Telser and H. S. La Pierre, *J. Am. Chem. Soc.*, 2019, **141**, 13222–13233.
- A. R. Willauer, I. Douair, A.-S. Chauvin, F. Fadaei-Tirani, J.-C. G. Bünzli, L. Maron and M. Mazzanti, *Chem. Sci.*, 2022, **13**, 681–691.
- T. P. Gomba, A. Ramanathan, N. T. Rice and H. S. La Pierre, *Dalton Trans.*, 2020, **49**, 15945–15987.
- A. C. Boggiano, S. R. Chowdhury, M. D. Roy, M. G. Bernbeck, S. M. Greer, B. Vlasisavljevich and H. S. La Pierre, *Angew. Chem., Int. Ed.*, 2024, **63**, e202409789.
- A. R. Willauer, C. T. Palumbo, F. Fadaei-Tirani, I. Zivkovic, I. Douair, L. Maron and M. Mazzanti, *J. Am. Chem. Soc.*, 2020, **142**, 5538–5542.
- A. Trovarelli, *Catalysis by Ceria and Related Materials*, Imperial College Press, 2002.
- A. Trovarelli, *Catal. Rev.*, 1996, **38**, 439–520.
- G. Pérez Bedwell, N. Suryadevara, Z. Qi, R. W. Gable, P. Bencok, M. L. Baker and C. Boskovic, *Inorg. Chem.*, 2025, **64**, 6519–6530.
- V. Nair and A. Deepthi, *Chem. Rev.*, 2007, **107**, 1862–1891.
- V. Sridharan and J. C. Menéndez, *Chem. Rev.*, 2010, **110**, 3805–3849.
- V. Nair, L. Balagopal, R. Rajan and J. Mathew, *Acc. Chem. Res.*, 2004, **37**, 21–30.
- H. Tsurugi and K. Mashima, *J. Am. Chem. Soc.*, 2021, **143**, 7879–7890.
- Y. Qiao, H. Yin, L. M. Moreau, R. Feng, R. F. Higgins, B. C. Manor, P. J. Carroll, C. H. Booth, J. Autschbach and E. J. Schelter, *Chem. Sci.*, 2021, **12**, 3558–3567.
- K. D. Shumilov, A. J. Jenkins, H. S. La Pierre, B. Vlasisavljevich and X. Li, *J. Am. Chem. Soc.*, 2024, **146**, 12030–12039.
- M. W. Löble, J. M. Keith, A. B. Altman, S. C. E. Stieber, E. R. Batista, K. S. Boland, S. D. Conradson, D. L. Clark, J. Lezama Pacheco, S. A. Kozimor, R. L. Martin, S. G. Minasian, A. C. Olson, B. L. Scott, D. K. Shuh, T. Tyliczszak, M. P. Wilkerson and R. A. Zehnder, *J. Am. Chem. Soc.*, 2015, **137**, 2506–2523.
- J. A. Branson, P. W. Smith, D.-C. Sergentu, D. R. Russo, H. Gupta, C. H. Booth, J. Arnold, E. J. Schelter, J. Autschbach and S. G. Minasian, *J. Am. Chem. Soc.*, 2024, **146**, 25640–25655.
- D.-C. Sergentu and J. Autschbach, *Chem. Sci.*, 2022, **13**, 3194–3207.
- S. A. Kozimor, P. Yang, E. R. Batista, K. S. Boland, C. J. Burns, D. L. Clark, S. D. Conradson, R. L. Martin, M. P. Wilkerson and L. E. Wolfsberg, *J. Am. Chem. Soc.*, 2009, **131**, 12125–12136.



- 25 R. Beekmeyer and A. Kerridge, *Inorganics*, 2015, **3**, 482–499.
- 26 P. L. Arnold, Z. R. Turner, N. Kaltsoyannis, P. Pelekanaki, R. M. Bellabarba and R. P. Tooze, *Chem. - Eur. J.*, 2010, **16**, 9623–9629.
- 27 N. Kaltsoyannis, *Inorg. Chem.*, 2013, **52**, 3407–3413.
- 28 A. Kerridge, *Dalton Trans.*, 2013, **42**, 16428–16436.
- 29 S. G. Minasian, J. M. Keith, E. R. Batista, K. S. Boland, D. L. Clark, S. A. Kozimor, R. L. Martin, D. K. Shuh and T. Tyliczszak, *Chem. Sci.*, 2014, **5**, 351–359.
- 30 M. Gregson, E. Lu, F. Tuna, E. J. L. McInnes, C. Hennig, A. C. Scheinost, J. McMaster, W. Lewis, A. J. Blake, A. Kerridge and S. T. Liddle, *Chem. Sci.*, 2016, **7**, 3286–3297.
- 31 S. G. Minasian, E. R. Batista, C. H. Booth, D. L. Clark, J. M. Keith, S. A. Kozimor, W. W. Lukens, R. L. Martin, D. K. Shuh, S. C. E. Stieber, T. Tyliczszak and X.-d. Wen, *J. Am. Chem. Soc.*, 2017, **139**, 18052–18064.
- 32 D.-C. Sergentu, T. J. Duignan and J. Autschbach, *J. Phys. Chem. Lett.*, 2018, **9**, 5583–5591.
- 33 V. E. J. Berryman, Z. J. Whalley, J. J. Shephard, T. Ochiai, A. N. Price, P. L. Arnold, S. Parsons and N. Kaltsoyannis, *Dalton Trans.*, 2019, **48**, 2939–2947.
- 34 T. Cheisson, K. D. Kersey, N. Mahieu, A. McSkimming, M. R. Gau, P. J. Carroll and E. J. Schelter, *J. Am. Chem. Soc.*, 2019, **141**, 9185–9190.
- 35 W. W. Lukens, S. G. Minasian and C. H. Booth, *Chem. Sci.*, 2023, **14**, 12784–12795.
- 36 L. A. Seaman, P. Hrobárik, M. F. Schettini, S. Fortier, M. Kaupp and T. W. Hayton, *Angew. Chem., Int. Ed.*, 2013, **52**, 3259–3263.
- 37 S. L. Staun, D.-C. Sergentu, G. Wu, J. Autschbach and T. W. Hayton, *Chem. Sci.*, 2019, **10**, 6431–6436.
- 38 G. T. Kent, X. Yu, G. Wu, J. Autschbach and T. W. Hayton, *Chem. Sci.*, 2021, **12**, 14383–14388.
- 39 H. Tateyama, A. C. Boggiano, C. Liao, K. S. Otte, X. Li and H. S. La Pierre, *J. Am. Chem. Soc.*, 2024, **146**, 10268–10273.
- 40 J. Du, J. A. Seed, V. E. J. Berryman, N. Kaltsoyannis, R. W. Adams, D. Lee and S. T. Liddle, *Nat. Commun.*, 2021, **12**, 5649.
- 41 O. Ordoñez, X. Yu, G. Wu, J. Autschbach and T. W. Hayton, *Inorg. Chem.*, 2024, **63**, 9427–9433.
- 42 D. E. Smiles, G. Wu, P. Hrobárik and T. W. Hayton, *J. Am. Chem. Soc.*, 2016, **138**, 814–825.
- 43 O. Ordoñez, X. Yu, G. Wu, J. Autschbach and T. W. Hayton, *Inorg. Chem.*, 2022, **61**, 15138–15143.
- 44 L. Bion, P. Moisy, F. Vaufray, S. Méot-Reymond and C. Madic, *Radiochim. Acta*, 1997, **78**, 73–82.
- 45 P. J. Subintoro and K. P. Carter, *Inorg. Chem.*, 2025, **64**, 11380–11397.
- 46 M. N. Sokolova, G. B. Andreev and A. B. Yusov, *Inorg. Chem. Commun.*, 2011, **14**, 1089–1092.
- 47 C. M. Tourné, G. F. Tourné and M. C. Brianso, *Acta Crystallogr., Sect. B: Struct. Sci.*, 1980, **36**, 2012–2018.
- 48 I. Colliard and G. J. P. Deblonde, *Inorg. Chem.*, 2024, **63**, 16293–16303.
- 49 C. Zhang, R. C. Howell, K. B. Scotland, F. G. Perez, L. Todaro and L. C. Francesconi, *Inorg. Chem.*, 2004, **43**, 7691–7701.
- 50 C. A. Colla, I. Colliard, A. M. Sawvel, M. Nyman, H. E. Mason and G. J. P. Deblonde, *Inorg. Chem.*, 2023, **62**, 6242–6254.
- 51 I. Colliard, J. R. I. Lee, C. A. Colla, H. E. Mason, A. M. Sawvel, M. Zavarin, M. Nyman and G. J. P. Deblonde, *Nat. Chem.*, 2022, **14**, 1357–1366.
- 52 I. Colliard and G. J. P. Deblonde, *J. Am. Chem. Soc.*, 2025, **147**, 14455–14467.
- 53 L. Fan, L. Xu, G. Gao, F. Li, Z. Li and Y. Qiu, *Inorg. Chem. Commun.*, 2006, **9**, 1308–1311.
- 54 J. Iijima, E. Ishikawa, Y. Nakamura and H. Naruke, *Inorg. Chim. Acta*, 2010, **363**, 1500–1506.
- 55 R. D. Peacock and T. J. R. Weakley, *J. Chem. Soc. A*, 1971, 1836–1839.
- 56 H. Naruke, J. Iijima and T. Sanji, *Inorg. Chem.*, 2011, **50**, 7535–7539.
- 57 J. Iijima and H. Naruke, *Inorg. Chim. Acta*, 2011, **379**, 95–99.
- 58 A. J. Gaunt, I. May, M. J. Sarsfield, D. Collison, M. Helliwell and I. S. Denniss, *Dalton Trans.*, 2003, 2767–2771.
- 59 R. Gupta, M. K. Saini, F. Doungmene, P. de Oliveira and F. Hussain, *Dalton Trans.*, 2014, **43**, 8290–8299.
- 60 I. Colliard and G. J. P. Deblonde, *JACS Au*, 2024, **4**, 2503–2513.
- 61 A. B. Yusov and V. P. Shilov, *Radiochemistry*, 2007, **49**, 144–151.
- 62 H. M. Qasim, W. W. Ayass, P. Donfack, A. S. Mougharbel, S. Bhattacharya, T. Nisar, T. Balster, A. Solé-Daura, I. Römer, J. Goura, A. Materny, V. Wagner, J. M. Poblet, B. S. Bassil and U. Kortz, *Inorg. Chem.*, 2019, **58**, 11300–11307.
- 63 A. Sundar, D. Bajuk-Bogdanović, G. Ćirić-Marjanović, T. Nisar, V. Wagner, A. Materny, G. B. Jameson, B. S. Bassil and U. Kortz, *ChemistryEurope*, 2025, **3**, 2500082.
- 64 J. Moons, L. S. Van Rompuy, A. Rodriguez, S. A. M. Abdelhameed, W. Simons and T. N. Parac-Vogt, *Polyhedron*, 2019, **170**, 570–575.
- 65 K. Stroobants, E. Moelants, H. G. T. Ly, P. Proost, K. Bartik and T. N. Parac-Vogt, *Chem. - Eur. J.*, 2013, **19**, 2848–2858.
- 66 N. Yadav and S. Singh, *Inorg. Chem.*, 2021, **60**, 7475–7489.
- 67 S. A. M. Abdelhameed, L. Vandebroek, F. de Azambuja and T. N. Parac-Vogt, *Inorg. Chem.*, 2020, **59**, 10569–10577.
- 68 D. Shiels, A. C. Berlfein, B. M. T. C. Peluzo, L. M. Lopez, A. W. Mitchell, W. W. Brennessel, M. Zeller, M. R. Crawley, S. C. Bart, M. T. Ruggiero and E. M. Matson, *Inorg. Chem.*, 2025, **64**, 9180–9194.
- 69 L. R. Valerio, D. Shiels, L. M. Lopez, A. W. Mitchell, M. Zeller, S. C. Bart and E. M. Matson, *Inorg. Chem.*, 2024, **63**, 22639–22649.
- 70 D. Shiels, W. W. Brennessel, M. R. Crawley and E. M. Matson, *Chem. Sci.*, 2024, **15**, 11072–11083.
- 71 R. Villanneau, A. Proust, F. Robert and P. Gouzerh, *J. Chem. Soc., Dalton Trans.*, 1999, 421–426.
- 72 S. She, C. Gao, K. Chen, A. Bayaguud, Y. Huang, B.-W. Wang, S. Gao and Y. Wei, *Inorg. Chem.*, 2018, **57**, 963–969.
- 73 J. J. Baldoví, Y. Duan, C. Bustos, S. Cardona-Serra, P. Gouzerh, R. Villanneau, G. Gontard, J. M. Clemente-



- Juan, A. Gaita-Ariño, C. Giménez-Saiz, A. Proust and E. Coronado, *Dalton Trans.*, 2016, **45**, 16653–16660.
- 74 W. G. Klemperer, in *Inorg. Synth.*, 1990, pp. 74–85.
- 75 T. M. Che, V. W. Day, L. C. Francesconi, M. F. Fredrich, W. G. Klemperer and W. Shum, *Inorg. Chem.*, 1985, **24**, 4055–4062.
- 76 A. Proust, P. Gouzerh and F. Robert, *Inorg. Chem.*, 1993, **32**, 5291–5298.
- 77 *CrysAlisPro*, Rigaku Corporation, Oxford, UK, 2024.
- 78 G. Sheldrick, *Acta Crystallogr., Sect. A: Found. Crystallogr.*, 2015, **71**, 3–8.
- 79 G. Sheldrick, *Acta Crystallogr., Sect. C: Cryst. Struct. Commun.*, 2015, **71**, 3–8.
- 80 F. Neese, *Wiley Interdiscip. Rev.: Comput. Mol. Sci.*, 2022, **12**, e1606.
- 81 C. Adamo and V. Barone, *J. Chem. Phys.*, 1999, **110**, 6158–6170.
- 82 E. van Lenthe, E. J. Baerends and J. G. Snijders, *J. Chem. Phys.*, 1994, **101**, 9783–9792.
- 83 E. v. Lenthe, E. J. Baerends and J. G. Snijders, *J. Chem. Phys.*, 1993, **99**, 4597–4610.
- 84 J. D. Rolfes, F. Neese and D. A. Pantazis, *J. Comput. Chem.*, 2020, **41**, 1842–1849.
- 85 E. D. Glendening, C. R. Landis and F. Weinhold, *J. Comput. Chem.*, 2013, **34**, 1429–1437.
- 86 T. Lu, *J. Chem. Phys.*, 2024, **161**, 082503.
- 87 T. Lu and F. Chen, *J. Comput. Chem.*, 2012, **33**, 580–592.
- 88 M. Pascual-Borràs, X. López, A. Rodríguez-Fortea, R. J. Errington and J. M. Poblet, *Chem. Sci.*, 2014, **5**, 2031–2042.
- 89 M. Filowitz, R. K. C. Ho, W. G. Klemperer and W. Shum, *Inorg. Chem.*, 1979, **18**, 93–103.
- 90 R. J. Errington, L. Coyle, P. S. Middleton, C. J. Murphy, W. Clegg and R. W. Harrington, *J. Cluster Sci.*, 2010, **21**, 503–514.
- 91 L. Coyle, P. S. Middleton, C. J. Murphy, W. Clegg, R. W. Harrington and R. J. Errington, *Dalton Trans.*, 2012, **41**, 971–981.
- 92 J. Novotny, S. Komorovsky and R. Marek, *Acc. Chem. Res.*, 2024, **57**, 1467–1477.
- 93 S. Moon and S. Patchkovskii, in *Calculation of NMR and EPR Parameters*, 2004, pp. 325–338.
- 94 C. N. Reilley, B. W. Good and R. D. Allendoerfer, *Anal. Chem.*, 1976, **48**, 1446–1458.
- 95 X. Ganglin, L. Bin, W. Wenliang, L. Qianding and L. Hengxin, *Acta Chim. Sin.*, 2002, **60**, 2022–2028.
- 96 L. Yuan, C. Qin, X. Wang and E. Wang, *Z. Naturforsch., B: J. Chem. Sci.*, 2007, **62**, 1471–1475.
- 97 H.-Y. Zhao, J.-W. Zhao, B.-F. Yang, H. He and G.-Y. Yang, *CrystEngComm*, 2014, **16**, 2230–2238.
- 98 R. Wan, P. Ma, M. Han, D. Zhang, C. Zhang, J. Niu and J. Wang, *Dalton Trans.*, 2017, **46**, 5398–5405.
- 99 D. Ravelli, D. Dondi, M. Fagnoni, A. Albini and A. Bagno, *Phys. Chem. Chem. Phys.*, 2013, **15**, 2890–2896.
- 100 D. Ravelli, D. Dondi, M. Fagnoni, A. Albini and A. Bagno, *J. Comput. Chem.*, 2011, **32**, 2983–2987.
- 101 P. N. Hazin, C. Lakshminarayan, L. S. Brinen, J. L. Knee, J. W. Bruno, W. E. Streib and K. Folting, *Inorg. Chem.*, 1988, **27**, 1393–1400.
- 102 P. N. Hazin, J. W. Bruno and H. G. Brittain, *Organometallics*, 1987, **6**, 913–918.
- 103 H. Yin, Y. Jin, J. E. Hertzog, K. C. Mullane, P. J. Carroll, B. C. Manor, J. M. Anna and E. J. Schelter, *J. Am. Chem. Soc.*, 2016, **138**, 16266–16273.
- 104 Y. Qiao, D.-C. Sergentu, H. Yin, A. V. Zabula, T. Cheisson, A. McSkimming, B. C. Manor, P. J. Carroll, J. M. Anna, J. Autschbach and E. J. Schelter, *J. Am. Chem. Soc.*, 2018, **140**, 4588–4595.
- 105 Y. Wang, J. Liang, C. Deng, R. Sun, P.-X. Fu, B.-W. Wang, S. Gao and W. Huang, *J. Am. Chem. Soc.*, 2023, **145**, 22466–22474.
- 106 Y. Qiao, T. Cheisson, B. C. Manor, P. J. Carroll and E. J. Schelter, *Chem. Commun.*, 2019, **55**, 4067–4070.
- 107 M. Ciampolini, F. Mani and N. Nardi, *J. Chem. Soc., Dalton Trans.*, 1977, 1325–1328, DOI: [10.1039/DT9770001325](https://doi.org/10.1039/DT9770001325).
- 108 M. R. Antonio, L. Soderholm, C. W. Williams, N. Ullah and L. C. Francesconi, *J. Chem. Soc., Dalton Trans.*, 1999, 3825–3830.
- 109 N. Haraguchi, Y. Okaue, T. Isobe and Y. Matsuda, *Inorg. Chem.*, 1994, **33**, 1015–1020.
- 110 D. W. Wester, G. J. Palenik and R. C. Palenik, *Inorg. Chem.*, 1985, **24**, 4435–4437.
- 111 J. Xu, E. Radkov, M. Ziegler and K. N. Raymond, *Inorg. Chem.*, 2000, **39**, 4156–4164.
- 112 L. Cardinale, S. S. Stahl, D. Kalyani and D. Lehnher, in *Advances in Catalysis*, ed. E. Hevia, M. H. Pérez-Temprano and M. Diéguez, Academic Press, 2023, vol. 72, pp. 57–102.
- 113 S. Diyali, N. Diyali, M. Das, M. Joshi, P. P. Ray, M. S. A. Sher Shah, A. Roy Choudhury and B. Biswas, *Cryst. Growth Des.*, 2022, **22**, 7590–7602.
- 114 A. Abbaspour and M. A. Mehrgardi, *Talanta*, 2005, **67**, 579–584.
- 115 N. G. Connelly and W. E. Geiger, *Chem. Rev.*, 1996, **96**, 877–910.
- 116 R. D. Shannon and C. T. Prewitt, *Acta Crystallogr., Sect. B: Struct. Sci.*, 1970, **26**, 1046–1048.
- 117 D. C. M. Llundell, J. Cirera, P. Alemany and S. Alvarez, *SHAPE, version 2.1*, 2013.
- 118 M. Pinsky and D. Avnir, *Inorg. Chem.*, 1998, **37**, 5575–5582.
- 119 J. A. Platts and R. J. Baker, *Dalton Trans.*, 2020, **49**, 1077–1088.
- 120 J. Vicha, J. Novotný, S. Komorovsky, M. Straka, M. Kaupp and R. Marek, *Chem. Rev.*, 2020, **120**, 7065–7103.
- 121 S. Cooper and N. Kaltsoyannis, *Dalton Trans.*, 2022, **51**, 5929–5937.
- 122 F. Weinhold, *J. Comput. Chem.*, 2012, **33**, 2440–2449.
- 123 V. E. J. Berryman, J. J. Shephard, T. Ochiai, A. N. Price, P. L. Arnold, S. Parsons and N. Kaltsoyannis, *Phys. Chem. Chem. Phys.*, 2020, **22**, 16804–16812.
- 124 A. C. Behrle, A. J. Myers, A. Kerridge and J. R. Walensky, *Inorg. Chem.*, 2018, **57**, 10518–10524.
- 125 K. A. Pace, V. V. Klepov, A. A. Berseneva and H.-C. zur Loye, *Chem. - Eur. J.*, 2021, **27**, 5835–5841.



- 126 J. Novotný, J. Vícha, P. L. Bora, M. Repisky, M. Straka, S. Komorovsky and R. Marek, *J. Chem. Theory Comput.*, 2017, **13**, 3586–3601.
- 127 M. Kaupp, V. G. Malkin, O. L. Malkina and D. R. Salahub, *J. Am. Chem. Soc.*, 1995, **117**, 1851–1852.
- 128 P. Hrobárik, V. Hrobáriková, A. H. Greif and M. Kaupp, *Angew. Chem., Int. Ed.*, 2012, **51**, 10884–10888.
- 129 A. H. Greif, P. Hrobárik, J. Autschbach and M. Kaupp, *Phys. Chem. Chem. Phys.*, 2016, **18**, 30462–30474.
- 130 J. Kaminský, J. Vícha, P. Bouř and M. Straka, *J. Phys. Chem. A*, 2017, **121**, 3128–3135.
- 131 V. Gowda, R. S. Laitinen, V.-V. Telkki, A.-C. Larsson, O. N. Antzutkin and P. Lantto, *Dalton Trans.*, 2016, **45**, 19473–19484.
- 132 J. Vícha, S. Komorovsky, M. Repisky, R. Marek and M. Straka, *J. Chem. Theory Comput.*, 2018, **14**, 3025–3039.
- 133 (a) CCDC 2481875: Experimental Crystal Structure Determination, 2025, DOI: [10.5517/ccdc.csd.cc2p9lh3](https://doi.org/10.5517/ccdc.csd.cc2p9lh3); (b) CCDC 2481876: Experimental Crystal Structure Determination, 2025, DOI: [10.5517/ccdc.csd.cc2p9lj4](https://doi.org/10.5517/ccdc.csd.cc2p9lj4); (c) CCDC 2481877: Experimental Crystal Structure Determination, 2025, DOI: [10.5517/ccdc.csd.cc2p9lk5](https://doi.org/10.5517/ccdc.csd.cc2p9lk5); (d) CCDC 2481878: Experimental Crystal Structure Determination, 2025, DOI: [10.5517/ccdc.csd.cc2p9ll6](https://doi.org/10.5517/ccdc.csd.cc2p9ll6).

