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When identification of the reduction sites in mixed Mo/W Keggin-type polyoxometalate hybrids turns out tricky

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ABSTRACT
The mixed Mo/W Kegging-type polyoxometalate hybrid (TBA)$_4$[PW$_9$Mo$_2$O$_{39}$Sn(C$_6$H$_4$I)] has been prepared by reaction between [α-PW$_9$Mo$_2$O$_{39}$]$^{7-}$ and [Cl$_3$Sn(C$_6$H$_4$I)] in dried acetonitrile, in the presence of tetra-n-butylammonium bromide. Further coupling reaction affords the ferrocenyl derivative (TBA)$_4$[PW$_9$Mo$_2$O$_{39}$][Sn(C$_6$H$_4$)C≡C(C$_6$H$_4$)Fc)]. The POM hybrids have been thoroughly characterized by NMR and IR spectroscopies. The electrochemical analysis confirms their ease of reduction when compared to the all-tungsten analog, albeit with a second reduction process occurring at a lower potential than in the all-molybdenum species. It is noteworthy that the second reduction is accompanied by an unusual red-shift of the electronic absorption spectrum. Whereas there is no doubt that the first reduction deals with molybdenum, the location of the second electron in the bi-reduced species, on the second molybdenum or on tungsten, has thus been the subject of a cross-investigation by spectro-electrochemistry, ESR and theoretical calculations. Finally, it came out that the second reduction is also molybdenum-centered with two unpaired and antiferromagnetically coupled extra electrons.
KEYWORDS mixed polyoxometalates, organic-inorganic hybrids, electroactivity, electronic structure, DFT, spectro-electrochemistry, ESR

INTRODUCTION
As molecular metal oxides, polyoxometalates (POMs) have astonishing redox properties, which are highly tunable according to their molecular structure, the nature of the metal addenda, the heteroatom, the counter cations… These redox properties have been harnessed in several fields of applications, ranging from (opto)-electronic devices to molecular batteries. In this context vanadates are especially attractive because of the accessibility of mixed valence-species and, to a lesser extent, molybdates. However molybdates suffer from higher kinetics lability compared to tungstates. This is heightened in the case of vacant species, used as precursors for the preparation of organic-inorganic POM based hybrids. Whereas POM hybrids of the PW_11 and P_2W_17 anions are well represented, hybrids based on the PMo_11 scaffold are scarce. In our previous studies we have shown the positive effect of Mo when associated to a photo-sensitizer. We have also described the immobilization of the diazonium terminated hybrids onto hydrogenated silicon and we have demonstrated the effect of the nature of the metal addenda, Mo versus W, on the electron transport properties of the resulting molecular junctions. Mixed Mo/W heteropolyanions combine the robustness of tungstates to the ease of reduction of molybdates. Furthermore, when the sites accommodating Mo versus W are precisely defined, it offers an original opportunity to play with localized (Mo^V) and delocalized (W^V) spins upon reduction, to potentially design electrically addressable qubits or quantum gates. In line with our previous work, we have thus chosen the mono-vacant [α-PW_9Mo_2O_39]^{7-} to enlarge the family of the Keggin-type POM hybrids. In this contribution we thus report on the synthesis and characterization of the tin derivative (TBA)_4[PW_9Mo_2O_39{Sn(C_6H_4I)}] K^{W9Mo2Sn} as a new platform to be subsequently engaged in post-functionalization reactions (TBA stands for tetra-
n-butylammonium cation). A remote ferrocenyl unit has also been introduced to give 
(TBA)$_4$[PW$_9$Mo$_2$O$_{39}$\{Sn(C$_6$H$_4$)C≡C(C$_6$H$_4$)Fc\}] $K^{W9Mo2}_{Sn}[Fc]$ in order to provide an internal redox reference for the study of the electrochemical behavior of the new mixed-POM hybrid.

The synthetic routes are presented on Scheme 1. In line with our previous studies,$^{28,29,33–36}$ and to enlarge the family of available precursors for POM processing onto surfaces, we have also prepared the diazonium terminated mixed hybrid 
(TBA)$_3$[PW$_9$Mo$_2$O$_{39}$\{Sn(C$_6$H$_4$)C≡C(C$_6$H$_4$)N$_2$\}] $K^{W9Mo2}_{Sn}[N_2^+]$ by deprotection of 
(TBA)$_4$[PW$_9$Mo$_2$O$_{39}$\{Sn(C$_6$H$_4$)C≡C(C$_6$H$_4$)N$_3$Et$_2$\}] $K^{W9Mo2}_{Sn}[N_3Et_2]$. However, its description exceeds the scope of this article and will be reported elsewhere. The electronic structure of the one- and two-electron reduced states of $K^{W9Mo2}_{Sn}$ (I and II) has been unraveled through combined spectro-electrochemistry, ESR investigation and theoretical calculations.

**Scheme 1.** Synthetic routes to the mixed Mo/W POM hybrids $K^{W9Mo2}_{Sn}$ and $K^{W9Mo2}_{Sn}[Fc]$. In this representation, WO$_6$, MoO$_6$ and PO$_4$ centers are depicted by blue, orange and green octahedra and tetrahedra respectively with metal atoms (W and Mo) and heteroatom P located at the center and oxygen atoms located at the apex of polyhedra. (i) CH$_3$CN, TBABr overnight (ii) [Pd(PPh$_3$)$_2$Cl$_2$], Et$_3$N, DMF, CuI (when required) overnight.

**RESULTS AND DISCUSSION**

**Synthesis of mixed-POM hybrids derived from $K_7[\alpha$-PW$_9$Mo$_2$O$_{39}$]**

While mono- or multi-vacant POMs have been extensively exploited for the incorporation of extra transition metal cations (i.e. other than W, Mo or V) or the preparation of organic-inorganic hybrids, to the best of our knowledge transition metal derivatives of $K_7[\alpha$-PW$_9$Mo$_2$O$_{39}$] or the corresponding $K_8[\alpha$-SiW$_9$Mo$_2$O$_{39}]$ have not been reported. During the final writing of this manuscript, the synthesis and characterization of the Dawson-type
organo-phosphonate hybrid $K_6[P_2W_{15}Mo_2O_{61}(POC_6H_5)_2]$ has been described, emphasizing the interest in well-defined and not randomly distributed mixed Mo/W POMs.$^{38}$

*Synthesis of $K_7[\alpha$-$PW_9Mo_2O_{39}$]*

The synthesis of the monovacant $K_7[\alpha$-$PW_9Mo_2O_{39}$] was first reported in 1977,$^{39}$ starting from the sodium salt of $[HPW_9O_{34}]^{8-}$. Note that the later was initially assigned to be a $B,\beta$-isomer but lately corrected to be a $A,\beta$-isomer.$^{40}$ Some years ago, some of us have thus reported the synthesis of $K_7[\alpha$-$PW_9Mo_2O_{39}$] by reaction between $K_9[A,\alpha$-$PW_9O_{34}]$ and sodium molybdate.$^{41}$ As usual in POM synthesis, the pH of the solution should be carefully controlled and kept between 4.5 and 5. Despite these precautions and many attempts, the $^{31}$P NMR spectrum of $K_7[\alpha$-$PW_9Mo_2O_{39}$] displays a small impurity at -10.28 ppm close to the main signal at -9.67 ppm. The degree of purity was estimated at 97%. This confirms the existence of a predominant isomer, assigned to two adjacent corner-shared MoO$_6$ octahedra, or 1,2-isomer according to IUPAC numbering of metal atom positions.$^{42}$

*Functionalization of $K_7[\alpha$-$PW_9Mo_2O_{39}]$- synthesis of the (TBA)$_4[PW_9Mo_2O_{39}Sn(C_6H_4I)]$*

$K^{WMo_2Sn}$ platform The synthesis of $K^{WMo_2Sn}$ follows those of $(TBA)_4[PW_{11}O_{39}\{Sn(C_6H_4I)\}]$ $K^{WSn}$ and $(TBA)_4[PMo_{11}O_{39}\{Sn(C_6H_4I)\}]$ $K^{MoSn}$,$^{43,44}$ by reaction between the monovacant POM and the trichloro-organotin derivative $[Cl_3Sn(C_6H_4I)]$. However, synthesis of POM hybrids is rarely straightforward and some adaptations are required. $K^{WSn}$ is prepared by reaction between $K_{7.8}Na_3[\alpha$-$PW_{11}O_{39}]$ and $[Cl_3Sn(C_6H_4I)]$ in water under pH control, followed by precipitation by addition of tetrabutylammonium bromide. Direct extension to $K_7[\alpha$-$PW_9Mo_2O_{39}]$ only gave mixtures of compounds according to $^{31}$P NMR spectra. As $K^{MoSn}$ is prepared in acetonitrile from the reaction between $(TBA)_4H_3[\alpha$-$PMo_{11}O_{39}]$ and $[Cl_3Sn(C_6H_4I)]$ in the presence of TBABr and NEt$_3$ used to neutralize the release of HCl, we also tried to start from a suspension of $K_7[\alpha$-$PW_9Mo_2O_{39}]$ in acetonitrile in the presence of
[Cl₃Sn(C₆H₄I)], NEt₃ and TBABr, the later being used as a transfer agent. This procedure also led to a mixture of compounds. Finally, we found that the presence of NEt₃ was not required if the amount of water contained in acetonitrile was limited by prior distillation onto CaH₂. After filtration of some unreacted K₇[α-PW₉Mo₂O₃₉], the solvent was evaporated and the resulting oil was dissolved in dichloromethane in the presence of TBABr. Subsequent workup included washing of the organic phase with water to eliminate mineral salts, evaporation of the solvent, redissolution in acetonitrile and final precipitation by addition of ethanol to recover K⁹⁹Mo²Sn as a white solid (yield 46%).

Post-functionalization of the (TBA)₄[PW₉Mo₂O₃₉{Sn(C₆H₄)}] K⁹⁹Mo²Sn platform: synthesis of (TBA)₄[PW₉Mo₂O₃₉{Sn(C₆H₄)C≡C(C₆H₄)Fc}] K⁹⁹Mo²Sn[Fc]

Once functionalized, the K⁹⁹Sn, K⁹⁹Mo and now the current K⁹⁹Mo²Sn platforms are robust enough to be engaged in post-functionalization reactions performed in organic solvents. The Pd-catalyzed C-C cross-coupling reactions offer an unlimited number of possibilities to anchor a remote functional group for further integration of POM hybrids into advanced architectures or processing into molecular materials. In this contribution we have chosen to illustrate Sonogashira-type reactions between K⁹⁹Mo²Sn and alkynes functionalized by a ferrocenyl (Fc) unit, subsequently used as an electrochemical probe, and, as a second example, by a diethyl-triazene group featuring a protected diazonium function. The reaction between K⁹⁹Mo²Sn and an excess of the appropriate alkyne was carried out in anhydrous DMF in the presence of distilled NEt₃ and the cis-[PdCl₂(PPh₃)₂] catalyst (plus if needed CuI). After elimination of non soluble materials, the crude compounds were precipitated by addition of diethylether, redissolved in CH₃CN, in the presence of TBABr and recovered as pure products by addition of ethanol (yield 77.7%).
Characterization by IR and NMR spectroscopies All compounds have been characterized by elemental analysis, mass spectrometry, IR and $^1$H and $^{31}$P NMR spectroscopies. The $^1$H and $^{31}$P NMR spectra together with the ESI-MS spectra are presented in SI. Integration of the signals corresponding to the aromatic protons relative to the signals characteristic of the TBA$^+$ cations is a good indicator of the accuracy of the proposed molecular formula, which is further confirmed by the molecular peak observed for the POM framework part in the ESI-MS spectra. The purity of the compounds was further attested by $^{31}$P NMR. On going from K$_7$[$\alpha$-PW$_9$Mo$_2$O$_{39}$] to K$^{^{W9Mo2}_{Sn}}$ the $^{31}$P chemical shift experienced a small upfield shift from -9.67 to -9.81 ppm in agreement with completion of the POM vacancy. Note however that the corresponding spectra were recorded in different solvents, LiCl/D$_2$O versus CD$_3$CN, which precludes any deeper comparison. In the K$^{^{W12-xMo_{x}}}$ series, the $^{31}$P Chemical shift $\delta_p$ was found to vary linearly with x, the higher x the higher $\delta_p$.$^{39}$ This is also verified here, $\delta_p$ for K$^{^{W9Mo2}_{Sn}}$ (-9.81 ppm) lying in between $\delta_p$ of K$^{^{W11}_{Sn}}$ (-10.76 ppm) and K$^{^{Mo11}_{Sn}}$ (-2.15 ppm). As expected, and previously observed for the full tungstate or molybdate homologues, the value of $^{31}$P chemical shift associated to the K$^{^{W9Mo2}_{Sn}}$ is barely altered after post-functionalization, with a value of -9.85 ppm associated to K$^{^{W9Mo2}_{Sn}[Fc]}$.

The $^{183}$W NMR spectrum of K$^{^{W9Mo2}_{Sn}}$ is compliant with a Cs symmetry with 5 resonance lines at -69.0, -93.4, -114.7, -116.2 and -135.5 ppm of relative intensities 2:2:2:2:1 (see Figure 1). It is very close to the spectrum reported for K$_5$[$\alpha$-SiW$_9$Mo$_2$VO$_{40}$] (-90, -93.4, -105.3, -109.4 and -134.8 ppm), except that the most down-field signal that was attributed to the two tungsten centers sharing edges with vanadium was much more broader.$^{46}$ This further supports a retention of the A-$\alpha$PW$_9$ structure with 2 corner shared MoO$_6$ octahedra and little if any isomerization (also supported by the $^{31}$P NMR spectrum).
The IR spectrum of $K_{W9Mo2Sn}$ displays the characteristic signature of a Keggin structure with three strong bands at 960, 885 and 808 cm$^{-1}$, corresponding to $\nu_{as}(M-O_d)$, $\nu_{as}(M-O_b-M)$ and $\nu_{as}(M-O_c-M)$, respectively. These values are very close to those of $K_{W11Sn}$ (963, 885 and 814 cm$^{-1}$) and higher than those of $K_{Mo11Sn}$ (943, 866, 806 and 785 cm$^{-1}$). They thus reflect both the composition of the polyanion and its total charge, the vibrations of tungstates occurring at higher wave numbers than those of molybdates and are shifted to lower values with an increase of the total charge. The presence of the phosphate group is disclosed by the $\nu_{as}(P-O)$ vibration at 1070 cm$^{-1}$ and additional bands at 2962 (m), 2937 (m), 2873 (m), 1481 (m) and 1379 (w) cm$^{-1}$ correspond to the stretching and bending modes of the CH$_n$ of the tetrabutylammonium cations. The stretching bands of the C=C bonds of the aromatic ring, expected in the 1500-1600 cm$^{-1}$ region, are too weak to be assigned precisely. Finally, at low wave numbers, in the 400-300 cm$^{-1}$ range, the two-band pattern at about 380 cm$^{-1}$ (sharp strong) and 330 cm$^{-1}$ (sharp medium) is characteristic of the $\alpha$-isomer. Post-functionalization has little effect on the vibrations of the metal-oxide scaffold.

The antisymmetric stretching $\nu(P-O)$ mode of the central PO$_4^{3-}$ group deserves a special comment. It is very sensitive to the symmetry of the anion: while TBA$_3$[PW$_{12}$O$_{40}$] and TBA$_3$[PMo$_{12}$O$_{40}$] display a single band at 1080 and 1063 cm$^{-1}$ respectively, the mono-vacant species TBA$_4$H$_3$[PW$_{11}$O$_{39}$] is characterized by a splitting of the previous band into two

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**Figure 1.** $^{183}$W NMR spectrum of (TBA)$_4$[PW$_9$Mo$_2$O$_{39}$(Sn(C$_6$H$_4$I))] $K_{W9Mo2Sn}$ in CH$_3$CN.
components at 1110 and 1060 cm\(^{-1}\) separated by 50 cm\(^{-1}\) (1079 and 1052 cm\(^{-1}\) for TBA\(_4\)H\(_3\)[PMo\(_{11}\)O\(_{39}\)], 1087 and 1050 cm\(^{-1}\) for K\(_7\)\([\alpha\text{-PW}_9\text{Mo}_2\text{O}_{39}]\)).\(^{24}\) In metal-substituted [PW\(_{11}\)O\(_{39}\)ML\(^{7-n}\)], the splitting between the two \(\nu(\text{P-O})\) vibrations is reduced and its variation is indicative of the interaction of the added ML\(^{n+}\) cation with the vacant POM and the central PO\(_4\)\(^{3-}\) group: the lower the splitting the better the cation refills the vacancy and restore a pseudo-tetrahedral geometry like in the complete POMs (\(L = \text{ligand}\)).\(^{49}\) This has been reported for the incorporation of first-row transition metal cations and we have also observed it in the metal-nitrido derivatives [PW\(_{11}\)O\(_{39}\)MN\(^{10-n}\)]\(^{(7-n)}\) (\(M = \text{Ru}^{\text{VI}}, \text{Cr}^{\text{V}}\)).\(^{50,51}\) The ionic radius of the extra cation compared to that of W\(^{\text{VI}}\)/Mo\(^{\text{VI}}\) and its electronic configuration are likely relevant parameters but the composition of the POM scaffold also appears to play a role: the IR spectra of K\(_{\text{Mo}}\)Sn and K\(_{\text{W9Mo2}}\)Sn display a single \(\nu(\text{P-O})\) band at 1070 cm\(^{-1}\) (but a shoulder at 1055 cm\(^{-1}\) is present on the IR spectra of the post-functionnalized species K\(_{\text{W9Mo2Sn[Fc]}}\)), whereas the \(\nu(\text{P-O})\) band is clearly splitted into two components at 1062 and 1035 cm\(^{-1}\) for K\(_{\text{Mo}}\)Sn (see Figures S5, S8 and S11).\(^{44}\) This suggests a decrease of the distortion from K\(_{\text{Mo}}\)Sn to K\(_{\text{W9Mo2}}\)Sn and an increase of the robustness of the tin insertion, as we are looking for.

**Electrochemical characterization**

The cyclic voltammograms of K\(_{\text{W9Mo2Sn}}\) and K\(_{\text{W9Mo2Sn[Fc]}}\) have been recorded in acetonitrile at a glassy carbon electrode. They are depicted on Figure 2 and relevant electrochemical data for these POM hybrids and others are summarized in Table 1. Both display two reduction processes around \(-0.55\) and \(-1.15\) V/SCE assigned to the POM framework and, for K\(_{\text{W9Mo2Sn[Fc]}}\), an oxidation process at \(+0.51\) V/SCE (Ep\(_a\) = +0.54, Ep\(_c\) = +0.48 \(\Delta\text{Ep} = 0.06\) V) attributed to the ferrocene unit. Fc was used as an internal redox reference and allows direct comparison of the intensities of the waves. Fc was used as an internal redox reference and allows direct comparison of the intensities of the waves of K\(_{\text{W9Mo2Sn[Fc]}}\), since a unique
diffusion coefficient is to be considered for the two covalently connected redox active units, $K^{W9Mo2}_{Sn}$ and $Fc$. It thus become clear that all processes correspond to mono-electronic transfers. The reduction potentials of $K^{W9Mo2}_{Sn}$ and $K^{W9Mo2}_{Sn}[Fc]$ are very similar and in the range of experimental uncertainty. As we have previously noted for organo-tin (and organo-silyl) derivatives, post-functionalization has almost no effect on the redox potentials.\textsuperscript{22,52–54} This suggests an electronic decoupling between the POM core and the organic tether, at variance with what was reported on organo-phosphonate POM hybrids.\textsuperscript{55,56}

Reduction of molybdates is known to be easier than reduction of tungstates\textsuperscript{31,57–59} and indeed the reduction potentials of $(TBA)_{4}[PMO_{11}O_{39}\{Sn(C_6H_4I)\}] \ K^{Mo}_{Sn}$ are shifted towards less negative values when compared to those of the $(TBA)_{4}[PW_{11}O_{39}\{Sn(C_6H_4I)\}] \ K^{W}_{Sn}$ homologue (see Table 1). Besides, the first reduction potentials of $(TBA)_{4}[PMO_{11}O_{39}\{Sn(C_6H_4I)\}]$ ($K^{Mo}_{Sn}$) and $K^{W9Mo2}_{Sn}$ are very close to each other. We can thus draw the conclusion that the first reduction process of $K^{W9Mo2}_{Sn}$ and of the related post-functionalized species leads to the reduction of one of the two molybdenum centers. Assignment of the second reduction event is not straightforward: transfer of a second electron on the second molybdenum could be energy favored but at the expense of a strong electrostatic repulsion. Conversely, reduction of one tungsten will bring additional delocalization on the tungstate scaffold. Note also that the difference between both reduction potentials is larger in the heterometallic $K^{W9Mo2}_{Sn}$ (0,60 V) than in the homometallic $K^{W}_{Sn}$ (0,46 V) and $K^{Mo}_{Sn}$ (0,41 V) species, which could either denote a second electron transfer to tungsten or a strong repulsion between two adjacent Mo(V). This issue was further addressed both theoretically and experimentally, through the electrochemical preparation of the one- and two-electron reduced species $1e-K^{W9Mo2}_{Sn}$ (I) and $2e-K^{W9Mo2}_{Sn}$ (II) and their characterization via UV-Vis and ESR spectroscopies. This will be discussed in a following part.
Cyclic voltammograms of 0.1 mM $\text{K}^{\text{W9Mo2Sn}}$ (left) and $\text{K}^{\text{W9Mo2Sn}[\text{Fc}]}$ (right) in CH$_3$CN (0.1 M TBAPF$_6$) at a glassy carbon electrode, potentials given versus SCE electrode, scan rate 0.1 V$s^{-1}$.

Table 1. Electrochemical data: cathodic peak current $\text{Ep}_c$, anodic peak current $\text{Ep}_a$, peak-to-peak separation $\Delta\text{Ep}$, cathodic to anodic peak intensities ratio $\text{ip}_c/\text{ip}_a$, mid-point potential $\text{Ep}_{1/2}$.

| POM | $\text{Ep}_c$ | $\text{Ep}_a$ | $\Delta\text{Ep}$ | $\text{ip}_c/\text{ip}_a$ | $\text{Ep}_{1/2}$ | ref |
|-----|---------------|---------------|-----------------|-----------------|-----------------|-----|
| $\text{K}^{\text{W9Mo2Sn}}$ | -0.99 | -0.50 | -0.62 | -0.47 | 0.15 | 0.97 | -0.55 | -1.20 | -1.09 | 0.11 | 1.13 | -1.15 |
| $\text{K}^{\text{W9Mo2Sn}[\text{Fc}]}$ | -0.58 | -0.47 | 0.11 | 1.04 | -0.53 | -1.19 | -1.09 | 0.10 | 1.08 | -1.14 |

Spectroscopic Insights into the Electronic structure of the 1- and 2-electron reduced (TBA)$_4[\text{PW}_{9}\text{Mo}_2\text{O}_{39}\{\text{Sn(C}_6\text{H}_4\text{I})\}]$, I and II

Spectro-electrochemistry- UV-Vis spectra of the reduced forms of (TBA)$_4[\text{PW}_{9}\text{Mo}_2\text{O}_{39}\{\text{Sn(C}_6\text{H}_4\text{I})\}]$

The stepwise reduction of a 0.2 mM solution of $\text{K}^{\text{W9Mo2Sn}}$ in MeCN (TBAPF$_6$ 0.1 M?) has been monitored by spectro-electrochemistry (see Figure 3). Applying a potential of -0.8 V/SCE induced the growth of a broad band around 560 nm ($\varepsilon = 750$ M$^{-1}$.cm$^{-1}$), which is compliant with the absorption spectra of 1e-$[\text{SiW}_{11}\text{Mo}^{\text{V}}\text{O}_{40}]^5$- ($\lambda = 510$ nm, $\varepsilon = 950$ M$^{-1}$.cm$^{-1}$), $^60$ 1e- $[\text{PW}_{11}\text{Mo}^{\text{V}}\text{O}_{40}]^4$+ ($\lambda = 500$ nm, $\varepsilon = 1150$ M$^{-1}$.cm$^{-1}$) and 5e-$[\text{SiW}_{9}\text{Mo}^{\text{V}}\text{V}^{\text{III}}\text{O}_{40}]^9$ (6e) ($\lambda = 488$ nm, $\varepsilon = 785$ M$^{-1}$.cm$^{-1}$).$^61$ At this step, the solution of 1e-$\text{K}^{\text{W9Mo2Sn}}$ (I) was more violet than blue, like the solution of 1e-$[\text{PW}_{11}\text{Mo}^{\text{V}}\text{O}_{40}]^4$+ which is reported to be red violet. This broad absorption has been variously attributed to a Mo(V) centered d-d transition$^60$ or
heteronuclear Mo(V)\textgreater\textgreater W(VI) intervalence charge transfer IVCT,\textsuperscript{37,61} and might simply conceal a superimposition of several electronic absorptions.

Decreasing the applied potential to -1.3 V/ECS strengthened a contribution around 720 nm (\(\varepsilon = 2950 \text{ M}^{-1}\cdot\text{cm}^{-1}\)) similar to the main contribution observed in the spectra of 1e-[\(\beta\)-SiW\textsubscript{9}Mo\textsubscript{2}O\textsubscript{40}]\textsuperscript{5-} and 1e-[SiMo\textsubscript{12}O\textsubscript{40}]\textsuperscript{5-} or 2e-[PMo\textsubscript{12}O\textsubscript{40}]\textsuperscript{5-} (related to the reduction of Mo),\textsuperscript{60,62,63} but also of 1e-[SiW\textsubscript{12}O\textsubscript{40}]\textsuperscript{5-} and 1e-[PW\textsubscript{12}O\textsubscript{40}]\textsuperscript{4-} (related to the reduction of W).\textsuperscript{64} It underscores that the absorption maxima of such broad bands are not very specific. The two electron reduced 2e-[SiW\textsubscript{12}O\textsubscript{40}]\textsuperscript{6-} and 2e-[PW\textsubscript{12}O\textsubscript{40}]\textsuperscript{5-} are rather characterized by an absorption at lower wavelength (\(\lambda = 625\) and 653 nm respectively).\textsuperscript{64} In these examples homonuclear IVCT, Mo(V)\textgreater\textgreater Mo(VI) or W(V)\textgreater\textgreater W(VI), are possible. Solutions of 2e-K\textsuperscript{W9Mo2}Sn (II) are blue.

The global shift of the absorption spectra to higher wavelength/lower energy upon the successive reduction steps of K\textsuperscript{W9Mo2}Sn is unusual and at variance with the trend indeed observed along the reduction of [\(\alpha\)-SiW\textsubscript{10}Mo\textsubscript{2}O\textsubscript{40}]\textsuperscript{4+}: 1e-[ \(\alpha\)-SiW\textsubscript{10}Mo\textsubscript{2}O\textsubscript{40}]\textsuperscript{5-} and 2e-[ \(\alpha\)-SiW\textsubscript{10}Mo\textsubscript{2}O\textsubscript{40}]\textsuperscript{6-} are respectively characterized by an absorption at \(\lambda = 590\) nm (\(\varepsilon = 930 \text{ M}^{-1}\cdot\text{cm}^{-1}\)) and \(\lambda = 500\) nm (\(\varepsilon = 3300 \text{ M}^{-1}\cdot\text{cm}^{-1}\)) (the two molybdenum are proposed to belong to the same triad in this example, so sharing edges and not corners like in our case).\textsuperscript{65} Similarly, 3e-[SiW\textsubscript{9}Mo\textsubscript{2}V\textsubscript{IV}O\textsubscript{40}]\textsuperscript{7-} is characterized by an absorption at \(\lambda = 490\) nm (the rather large \(\varepsilon = 4070 \text{ M}^{-1}\cdot\text{cm}^{-1}\) value in this case is due to an additive contribution of V(IV)).\textsuperscript{37} Taken all together, these experimental observations would suggest to associate the second reduction event in K\textsuperscript{W9Mo2}Sn to tungsten (II\textsubscript{MoW}) rather than to the second molybdenum (II\textsubscript{MoMo}), with an unexpected effect of the break of symmetry introduced by the tin functionalization. However, UV-Vis spectra of reduced POMs are finally close to one another, be it in the Keggin- or in the Dawson-series,\textsuperscript{55,64,66,67} with large bands resulting from the overlap of multiple absorption events, d-d, homo- and hetero-nuclear IVCT transitions, occurring in the
whole visible-near IR range and with relative intensities sensitive to structural isomerism.\cite{65}

At this stage, drawing a definitive conclusion regarding the location of the second electron in II, on tungsten ($\text{II}_{\text{MoW}}$) or on molybdenum ($\text{II}_{\text{MoMo}}$) is thus gambling. Further insights will be provided by theoretical calculations.

**Figure 3.** Absorbance (A) monitoring of the electronic spectrum of a 0.2 mM solution of $\text{K}^{\text{WMoO}_2}_{\text{80}}$ in CH$_2$CN (TBAPF$_6$, 0.1 M), upon application of a constant potential of -0.8 V/SCE to generate I (black) then -1.3 V/SCE to generate II (blue).

*ESR spectroscopy of the reduced forms of [PW$_6$Mo$_2$O$_{39}${Sn(C$_6$H$_4$I)$_4$}]$^4-$, 1e-[PW$_6$Mo$_2$O$_{39}${Sn(C$_6$H$_4$I)$_4$}]$^3-$ (I) and 2e-[PW$_6$Mo$_2$O$_{39}${Sn(C$_6$H$_4$I)$_4$}]$^2-$ (II, either $\text{II}_{\text{MoW}}$ or $\text{II}_{\text{MoMo}}$)

Whatever the location of the second electron in II, the final spin state, $S = 0$ or $S = 1$ is to be determined. Provided electron delocalization, a singlet state is generally observed for homometallic POM reduced to an even number of electrons.\cite{66,67,68,69} In mixed POMs the picture is to be nuanced.\cite{2} The properties of tungsto-vanadates have been especially investigated.  

Whereas discussing the electronic structure of reduced [PW$_{10}$V$_2$O$_{40}$]$^{5-}$ or [PMo$_{10}$V$_2$O$_{40}$]$^{5-}$ is meaningless because the general formula hides mixtures of 5 geometrical isomers, [SiW$_{10}$V$_2$O$_{40}$]$^{6-}$ yet corresponds to a single 1,2 isomer with adjacent corner shared VO$_6$ octahedra.\cite{42,70} The ESR spectrum of 1e-[SiW$_{10}$V$_2$O$_{40}$]$^{7-}$ displays a 15-line ESR pattern in
agreement with an electron delocalized over the two vanadium (I = 7/2) but magnetic susceptibility measurements carried out on 2e-[SiW_{10}V_2O_{40}]^8^- revealed that there is essentially no magnetic exchange between the two V(IV), thus two isolated spin $\frac{1}{2}$.\(^{72}\) This is in contrast with the behavior of [SiW_{9}V_{3}O_{40}]^7^-, with one electron trapped on one vanadium in 1e-[SiW_{9}V_{3}O_{40}]^7^-; and two antiferromagnetically coupled electrons in 2e-[SiW_{9}V_{3}O_{40}]^8^- (J = -34.9 cm\(^{-1}\)). The striking difference has been ascribed to some variation of the VOV angle values, possibly modulated by protonation.

During the spectro-electrochemical study on [PW_{9}Mo_{2}O_{39}\{Sn(C_6H_4I)\}]^4^-, aliquots of the solution have been frozen (20K) to be probed by ESR spectroscopy. Parallel to the growth of the first UV-Vis feature, one can clearly observed in X-Band ESR the apparition of a quite rhombic signature around g=2 (see Figure 4). Interestingly enough, one can clearly observe on the side of this main feature hyperfine lines that could account for the about 25% of Mo atoms bearing a 5/2 nuclear spin ($^{95}$Mo (15.7%) and $^{97}$Mo (9.6%)). Indeed, satisfactory simulation of this ESR features where obtained with Easy Spin using the naturally abundant isotopes of Mo with a rhombic g-matrix (g_x=1.941, g_y=1.916, g_z=1.899) and A_x=135 MHz (~50G) and A_z=180 MHz (~68G) (A_y could not be resolved).\(^{73}\) These values are quite in line with those obtained by Pope for the monoreduced [PMoVW_{11}O_{40}] anion with an axial spin system (g|| = 1.918; A_3= 81.3; g_\perp = 1.937, A_\perp= 33.2 G)\(^{61,74}\) and confirm that the first reduction is localized on a Mo atom in I. Upon further reduction, a progressive decrease of the signal is observed with no new features growing, indicating that the 2 electron reduced species II is silent in these conditions.
**Figure 4.** X-band ESR spectrum of a frozen solution (20K) of \( \text{[PW}_9\text{Mo}_2\text{O}_{39}\text{Sn(C}_6\text{H}_4\text{I})]^{5-} \) obtained by spectro-electrochemistry (solution in CH$_3$CN, TBAPF$_6$ 0.1 M). inset: amplification of the central part unveiling the hyperfine coupling.

The amount of II obtained by spectro-electrochemistry was too low to allow magnetic susceptibility measurements. The chemical reduction path has also been investigated, using sodium naphtalenide as reducing agent.$^{75}$ However, we have been faced with the (expected) high sensitivity of the reduced species to reoxidation, so that only characterization of I could be carried out (electronic and ESR spectroscopies, see SI, corroborating the previously described features).

**Theoretical calculations**

To better understand the electronic structure of \( \text{K}^{\text{W9Mo2Sn}} \) and the localization of electrons in its one- and two-electron reduced counterparts, we next conducted theoretical DFT calculations. As shown in Figure 5a, the HOMO of the fully-oxidized system is localized on the organic moiety, while the LUMO and the LUMO+1 orbitals are both centered on Mo(VI) ions, accounting for antibonding combinations of d(Mo) and p(O) orbitals. Notably, the first empty MOs with main contribution from d(W) orbitals are the quasi-degenerated LUMO+2 and LUMO+3, which lie ca. 0.20 eV above the LUMO+1. The first reduction of \( \text{K}^{\text{W9Mo2Sn}} \) was thus found to occur in one of the Mo centers (see Figure 5b for spin density
As already inferred from electrochemical studies (vide supra). Most importantly, DFT calculations revealed that the intriguing second reduction is also Mo-centered, as shown in the spin density distribution of II (Figure 5c). Supporting this assignment, the DFT-calculated reduction potentials for the two successive Mo(VI)→Mo(V) reduction steps (-0.54 and -1.21 V vs SCE, B3LYP-D3 level) are in rather good agreement with the experimental values (Table 2). In fact, using a model system whereby one of the Mo centers was replaced by W, we estimated the potential for a W-centered second reduction to be excessively negative (-1.74 V vs SCE) to match the experimental value of -1.14 V. Similar results for Mo(VI)→Mo(V) reduction steps were found using ωB97X-D and HSE06 functionals (Table 2).

In agreement with the ESR measurements, our DFT calculations predict the ground state of II to be an open-shell singlet, whereby the two ‘extra’ electrons are unpaired but antiferromagnetically coupled (Figure 5c, Table 2). The open-shell singlet solution obtained with the broken-symmetry formalism lies 0.8 kcal mol⁻¹ below in energy than the most stable triplet state and 6.8 kcal mol⁻¹ below the closed-shell singlet, in which both electrons are localized in a MO with contribution from d-type orbitals of both Mo centers. This is in line with previous theoretical studies on the diamagnetism of bi-reduced POMs, which showed that the hopping integrals between neighboring centers and the electron-electron repulsion stabilize the singlet over the triplet state.²⁷⁻²⁹ However, it is important to note that the singlet-state stabilization mechanism in systems with localized electrons differs from those involving hopping electrons. Similar behavior was found using ωB97X-D and HSE06 functionals. Spin density values close to one for Mo atoms (positive and negative signs for alpha and beta electrons, respectively) is consistent with the presence of Mo(V) ions. The rest of the spin density is delocalized over the oxo ligands of Mo(V) centers, as shown in Figure...
5c. Finally, the evolution of the DFT-simulated absorption spectra in going from I to II (see Figure S10 and related text in the SI) reproduces rather well the experimental variations observed in Figure 3, further supporting the herein described electronic structure for species II.

Figure 5. a) Schematic frontier molecular orbital diagram for $K^\text{W9Mo2Sn}_\text{a}$ at B3LYP-D3 level. Table S1 in the SI compiles the frontier MO energies for $K^\text{W9Mo2Sn}_\text{a}$ calculated with different density functionals. (b) and (c) Spin density distributions for the one- and the two-electron reduced species I and II, respectively, showing successive reduction steps of Mo centers and the antiferromagnetic nature of II. Color code: W (blue), Mo (orange), Sn (pink), P (green), I (purple), C (gray), O (red), H (white).

Table 2. Electronic and energetic parameters for calculated $K^\text{W9Mo2Sn}_\text{a}$ and its reduced partners with different density functionals.

| Functional | $E^\text{red}_\text{a}$ | $\Delta E^\text{red}_\text{b}$ | Spin density distribution in II$^c$ | $\Delta G$ (triplet-singlet)$^d$ |
|------------|----------------|-----------------|----------------------|------------------------|
|            | 0e/1e          | 1e/2e           | Mo1                  | Mo2                    |                      |
| B3LYP-D3   | -0.54          | -1.21           | 0.67                 | -0.80                  | 0.79                 | +0.8                |
| oB97X-D    | -0.43          | -1.16           | 0.73                 | -0.84                  | 0.85                 | +0.3                |
| HSE06      | -0.55          | -1.30           | 0.75                 | -0.82                  | 0.82                 | +0.6                |
| Exp.       | -0.55          | -1.15           | 0.61                 | -                      | -                     |                      |

$^a$ Redox potentials (in V) were calculated against SCE, taking an absolute potential of 4.67 V for SCE in nonaqueous conditions. $^b$ Computed redox potential difference between first and second electron reductions. $^c$ Mulliken spin densities for the singlet (broken symmetry) state. $^d$ Energy difference (in kcal mol$^{-1}$) between triplet and singlet states computed at different DFT levels.
CONCLUDING REMARKS
The family of Keggin-type POM hybrids has been enlarged with mixed Mo/W species, (TBA)$_4$[PW$_9$Mo$_2$O$_{39}$[Sn(C$_6$H$_4$I)]] and its ferrocenyl derivative. Both species have similar electroactivity with reduction processes facilitated by the incorporation of molybdenum. This widens the range of redox potential tuning often harnessed in POM based molecular materials. The mono-reduced species, [PW$_9$Mo$^{VI}$Mo$^{V}$O$_{39}$[Sn(C$_6$H$_4$I)]$^5$$^-$, is stable below -0.55 V/ECS and characterized by a main electronic transition at 560 nm. According to TD-DFT calculations, this band is assigned to superimposed Mo(V) d-d transition and intervalence Mo(V)→W(VI) charge transfers (IVCT), while Mo(V)→ Mo(VI) IVCT lies around 1000 nm. The intensity-increase and the unusual red-shift of the UV-vis spectrum upon further reduction, are rather well reproduced by the simulation, in agreement with the apparent maximum found experimentally at 720 nm. Once corroborated by the spectroscopic data, DFT calculations furthermore unveils the site of the second reduction. The bi-reduced species should thus be described as [PW$_9$Mo$_2$$^{V}$O$_{39}$[Sn(C$_6$H$_4$I)]$^6$$^-$, with an open shell singlet ground state, consistently silent in ESR. This plural approach, confronting experimental and theoretical data to the abundant literature dealing with reduced POMs points out the intricacy of their electronic structure and the difficulty to properly attribute the observed electronic events on the basis of a sole technique. This is illustrated by the UV-Vis spectra with broad bands that result from the overlap of so many transitions, metal centered or metal-to-metal, that the quoted absorption maxima are finally of low significance.

EXPERIMENTAL SECTION
General. Chemicals and solvents were obtained from Aldrich or Acros and used as received, except triethylamine and acetonitrile that were distilled from CaH$_2$. K$_7$-α[PW$_9$Mo$_2$O$_{39}$] was prepared as previously described.$^{39,41}$ $^1$H (400MHz), $^{31}$P (121.5MHz) and $^{183}$W (25 MHz)
NMR spectra were recorded on a Bruker AvanceIII Nanobay 400 MHz spectrometer equipped with a BBFO probehead (\(^1\)H and \(^{31}\)P NMR) or on a Bruker AvanceIII 600 MHz spectrometer equipped with a BBO probehead (\(^{183}\)W NMR, 10 mm o.d. tube), respectively. Chemical shifts are quoted as parts per million (ppm) relative to tetramethylsilane using the solvent signals as secondary standard for \(^1\)H and relative to 85\% H\(_3\)PO\(_4\) for \(^{31}\)P and to 2M Na\(_2\)WO\(_4\) alkaline solution in D\(_2\)O for \(^{183}\)W. (s: singlet, d: doublet, t: triplet, sex: sextet, m: multiplet) and coupling constants (J) are quoted in Hertz (Hz). IR spectrum of the powder was recorded from a KBr pellet on a Jasco FT/IR 4100 spectrometer. Elemental analyses were performed at the Institut de Chimie des Substances Naturelles, Gif sur Yvette, France. High-resolution ESI mass spectra were recorded using an LTQ Orbitrap hybrid mass spectrometer (Thermofisher Scientific, Bremen, Germany) equipped with an external ESI source operated in the negative ion mode. Spray conditions included a spray voltage of 3 kV, a capillary temperature maintained at 280 °C, a capillary voltage of −30 V, and a tube lens offset of −90 V. Sample solutions in acetonitrile (10 pmol.μL\(^{-1}\)) were infused into the ESI source by using a syringe pump at a flow rate of 180 μ. h\(^{-1}\). Mass spectra were acquired in the Orbitrap analyzer with a theoretical mass resolving power (Rp) of 100 000 at m/z 400, after ion accumulation to a target value of 105 and a m/z range detection from m/z 300 to 2000. All data were acquired using external calibration with a mixture of caffeine, MRFA peptide and Ultramark 1600 dissolved in Milli-Q water/HPLC grade acetonitrile (50/50, v/v).

Cyclic voltammetry were performed in a three electrode cell, with a glassy carbon working electrode, a Pt counter electrode and a saturated Hg\(_2\)Cl\(_2\)/KCl reference electrode fitted with a bridge containing a saturated aqueous LiCl solution.

**Synthesis of TBA\(_4\)[PW\(_9\)Mo\(_2\)O\(_{39}\)Sn(C\(_6\)H\(_4\)I)] (K\(^{W9Mo2Sn}\)).** [Cl\(_3\)SnC\(_6\)H\(_4\)I] (95.5mg, 0.223mmol) and an excess of tetrabutylammonium bromide TBABr (310mg, 0.962mmol) were placed into 20mL of dried acetonitrile. The latest solution was transferred onto
K$_7$[PW$_9$Mo$_2$O$_{39}$] (500mg, 0.132mmol) and the resulting suspension stirred overnight at room temperature. After centrifugation, the filtered supernatant was concentrated under reduced pressure and led to a greenish oil. This oil was then dissolved into 10mL of dichloromethane with an excess of tetrabutylammonium bromide (200mg, 0.620mmol) and the solution stirred for 1 hr. at room temperature. The organic phase was washed by distilled water (3x30mL) and finally it was evaporated under reduced pressure leading to a white solid. The solid was dissolved in 15mL of dried acetonitrile. A non-dissolved part was filtered and the filtrate was concentrated under reduced pressure until a few mL. K$_{W9Mo2}$Sn was precipitated from the latest concentrated solution by adding an excess of absolute ethanol (45mL). After centrifugation, the product was dried by diethylether. After a last centrifugation step, K$_{W9Mo2}$Sn was obtained as a greenish white powder. Yield: 316 mg (46%).

$^1$H NMR (CD$_3$CN): $\delta$ = 7.84 (d+dd, $^3$J$_{H-H}$=8.13Hz, $^4$J$_{Sn-H}$= 31.47Hz, 2H, Ar-H), 7.44 (d+dd, $^3$J$_{H-H}$=8.13Hz, $^3$J$_{Sn-H}$= 95.4Hz, 2H, Ar-H), 3.13 (m, 32H, N-CH$_2$-CH$_2$-CH$_2$-CH$_3$), 1.63 (m, 32H, N-CH$_2$-CH$_2$-CH$_2$-CH$_3$), 1.39 (s, $^3$J$_{H-H}$=7.32Hz, 32H, N-CH$_2$-CH$_2$-CH$_2$-CH$_3$), 0.98 (t, $^3$J$_{H-H}$=7.32Hz, 48H, N-CH$_2$-CH$_2$-CH$_2$-CH$_3$).

$^{31}$P NMR (CD$_3$CN): $\delta$ = -9.87 (s+d, $^2$J$_{Sn-P}$=27.07Hz).

IR (KBr, cm$^{-1}$): $\nu$ = 2962 (m), 2937 (m), 2873 (m), 1481 (m), 1379 (w), 1070 (s), 960 (s), 885 (s), 808 (s), 683(w), 513 (w), 380 (m).

Anal. Calcd for PW$_9$Mo$_2$O$_{39}$SnIC$_7$H$_{148}$N$_4$ (%): C, 22.16; H, 3.93; N, 1.48. Found: C, 21.95; H, 4.00; N, 1.44. HRMS (ESI): $m/z$: [M]$^{4+}$ calc 705.75 found 705.75; [M+TBA]$^{3+}$ calc 1021.76 found 1022.09; [M+2TBA]$^{2+}$ calc 1653.78 found 1654.58 for PW$_9$Mo$_2$O$_{39}$SnIC$_7$H$_{148}$N$_4$.

**Synthesis of TBA$_4$[PW$_9$Mo$_2$O$_{39}$[Sn(C$_6$H$_4$)C≡C(C$_5$H$_4$)Fe(C$_5$H$_5$)]] (K$_{W9Mo2}$Sn[Fe]).** K$_{W9Mo2}$Sn (100mg, 0.026mmol), ethynylferrocenyl (16.8mg, 0.080mmol) and bis-(triphenylphosphine)palladium(II) dichloride (3.1 mg, 0.004mmol) were dissolved into 2mL of anhydrous DMF and the solution was purged with Ar. Freshly distilled triethylamine (70µL) were then added. The mixture stirred overnight after being degassed with Ar 5min
more. An unidentified solid was eliminated by centrifugation. A product was precipitated by addition of diethylether (45mL) to the supernatant. The resulting solid was separated by centrifugation and then was solubilised into a minimum of acetonitrile. TBABr (126 mg, 0.391mmol) was added to the latest concentrated solution and the mixture was filtered to eliminate a non-identified solid. $K^{W9Mo2Sn}[Fe]$ was precipitated by addition of absolute ethanol (45mL) to the filtrate. The product was separated by centrifugation and was dried by addition of diethylether. The pure $K^{W9Mo2Sn}[Fe]$ was obtained after one last centrifugation step. If the number of TBA determined by $^1$H NMR was lower than expected, one more step was required by using TBA-enriched Amberlyst® resin into acetonitrile. The final product was then obtained as an orange powder. Yield: 78.3 mg (77.7%). $^1$H NMR (CD$_3$CN): $\delta = 7.66$ (d+dd, $^3J_{H-H}=8.24$Hz, $^3J_{Sn-H}=95.0$Hz, 2H, Ar-H), 7.53 (d+dd, $^3J_{H-H}=8.24$Hz, $^4J_{Sn-H}=33.4$Hz, 2H, Ar-H), 4.54 (t, $^3J_{H-H}=1.90$Hz, 2H, Cp-H), 4.31 (t, $^3J_{H-H}=1.90$Hz, 2H, Cp-H), 4.27 (s, 5H, Cp-H), 3.13 (m, 32H, N-CH$_2$-CH$_2$-CH$_2$-CH$_3$), 1.63 (m, 32H, N-CH$_2$-CH$_2$-CH$_2$-CH$_3$), 1.39 (sex, $^3J_{H-H}=7.36$Hz, 32H, N-CH$_2$-CH$_2$-CH$_2$-CH$_3$), 0.98 (t, $^3J_{H-H}=7.36$Hz, 48H, N-CH$_2$-CH$_2$-CH$_2$-CH$_3$). $^{31}$P NMR (CD$_3$CN): $\delta = -9.85$ (s+d, $^2J_{Sn-P}=26.87$Hz). IR (KBr, cm$^{-1}$): $\nu =$ 2960 (m), 2933 (m), 2872 (m), 1483 (m), 1379 (w), 1070 (s), 1056 (sh.), 959 (s), 887 (m), 799 (vs), 499 (w), 378 (m), 332 (w). Anal. Calcd for PW$_9$Mo$_2$O$_{39}$SnFe C$_{82}$H$_{157}$N$_4$ (%): C, 25.41; H, 4.08; N, 1.45. Found: C, 25.47; H, 4.12; N, 1.52. HRMS (ESI): $m/z$: [M]$^+$ calcd 726.27 found 725.77; [M+TBA]$^+$ calcd 1049.12 found 1049.46; [M+2TBA]$^2+$ calcd 1694.83 found 1696.84 for PW$_9$Mo$_2$O$_{39}$SnFe C$_{82}$H$_{157}$N$_4$.

**Computational details.** DFT calculations were performed with the Gaussian 16 (rev. A03)$^{79}$ quantum chemistry package using four different hybrid correlation-exchange functionals, namely B3LYP-D3,$^{80-82}$ oB97X-D$^{83}$ and HSE06.$^{84}$ The LANL2DZ basis set$^{85}$ and associated pseudopotentials were used to describe Mo, W, Sn and I atoms, while the Pople-type 6-
31G(d,p) basis set \( ^{86-88} \) was adopted for the remaining atoms. Solvent effects of acetonitrile were included both in geometry optimizations and energy calculations by means of the IEF-PCM \(^{89} \) implicit solvent model as implemented in Gaussian 16. Geometry optimizations were full and without any symmetry restriction and all the minima were characterized by the lack of imaginary frequencies. Absorption spectra were simulated using time-dependent DFT \(^{90,91} \) in combination with the HSE06 functional, solvent effects and the same basis set described above.

ASSOCIATED CONTENT

Supporting Information IR, \(^{1} \)H NMR, \(^{31} \)P NMR and mass spectra, cyclic-voltammograms at various sweep rates, chemical reduction attempts; DFT calculations: energies for frontier molecular orbitals, TD-DFT-simulated absorption spectra and cartesian coordinates.

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REFERENCES

(1) Pope, M. T. *Heteropoly and Isopoly Oxometalates*. Springer Berlin Heidelberg 1983, ISBN 978-3-662-12006-4
(2) Poblet, J. M.; López, X.; Bo, C. Ab Initio and DFT Modelling of Complex Materials:
Towards the Understanding of Electronic and Magnetic Properties of Polyoxometalates. *Chem. Soc. Rev.* **2003**, *32*(5), 297–308. https://doi.org/10.1039/B109928K.

(3) Ueda, T. Electrochemistry of Polyoxometalates: From Fundamental Aspects to Applications. *ChemElectroChem* **2018**, *5*(6), 823–838. https://doi.org/10.1002/celc.201701170.

(4) Gumerova, N. I.; Rompel, A. Synthesis, Structures and Applications of Electron-Rich Polyoxometalates. *Nat Rev Chem* **2018**, *2*(2), 0112. https://doi.org/10.1038/s41570-018-0112.

(5) Sartzi, H.; Long, D.-L.; Sproules, S.; Cronin, L.; Miras, H. N. Directed Self-Assembly, Symmetry Breaking, and Electronic Modulation of Metal Oxide Clusters by Pyramidal Heteroanions. *Chem. Eur. J.* **2018**, *24*(17), 4399–4411. https://doi.org/10.1002/chem.201705711.

(6) Grigoriev, V. A.; Hill, C. L.; Weinstock, I. A. Role of Cation Size in the Energy of Electron Transfer to 1:1 Polyoxometalate Ion Pairs \((M^+)(X^{n−}VW_{11}O_{39})^{(8−)}\) \((M = Li, Na, K)\). *J. Am. Chem. Soc.* **2000**, *122*(14), 3544–3545. https://doi.org/10.1021/ja993862c.

(7) Chen, B.; Neumann, R. On the Effect of Ion Pairing of Keggin Type Polyoxoanions with Quaternary Ammonium Cations on Redox Potentials in Organic Solvents. *Phys. Chem. Chem. Phys.* **2016**, *18*(32), 22487–22493. https://doi.org/10.1039/C6CP03315F.

(8) Busche, C.; Vilà-Nadal, L.; Yan, J.; Miras, H. N.; Long, D.-L.; Georgiev, V. P.; Asenov, A.; Pedersen, R. H.; Gadegaard, N.; Mirza, M. M.; Paul, D. J.; Poblet, J. M.; Cronin, L. Design and Fabrication of Memory Devices Based on Nanoscale Polyoxometalate Clusters. *Nature* **2014**, *515*(7528), 545–549. https://doi.org/10.1038/nature13951.

(9) Li, J.-S.; Sang, X.-J.; Chen, W.-L.; Zhang, L.-C.; Zhu, Z.-M.; Ma, T.-Y.; Su, Z.-M.; Wang, E.-B. Enhanced Visible Photoresponse of TiO\(_2\) Thin Film with an All-Inorganic Donor–Acceptor Type Polyoxometalate. *ACS Applied Materials & Interfaces* **2015**, *7*(24), 13714–13721. https://doi.org/10.1021/acsami.5b03948.

(10) Vasilopoulou, M.; Douvas, A. M.; Palilis, L. C.; Kennou, S.; Argitis, P. Old Metal Oxide Clusters in New Applications: Spontaneous Reduction of Keggin and Dawson Polyoxometalate Layers by a Metallic Electrode for Improving Efficiency in Organic Optoelectronics. *J. Am. Chem. Soc.* **2015**, *137*(21), 6844–6856. https://doi.org/10.1021/jacs.5b01889.

(11) Ji, Y.; Huang, L.; Hu, J.; Streb, C.; Song, Y.-F. Polyoxometalate-Functionalized Nanocarbon Materials for Energy Conversion, Energy Storage and Sensor Systems. *Energy & Environmental Science* **2015**, *8*(3), 776–789. https://doi.org/10.1039/C4EE03749A.

(12) Chen, J.-J.; Symes, M. D.; Cronin, L. Highly Reduced and Protonated Aqueous Solutions of \([P_2W_{18}O_{62}]^{16−}\) for on-Demand Hydrogen Generation and Energy Storage. *Nature Chem* **2018**, *10*(10), 1042–1047. https://doi.org/10.1038/s41557-018-0109-5.

(13) Friedl, J.; Holland-Cunz, M. V.; Cording, F.; Pfanschilling, F. L.; Wills, C.; McFarlane, W.; Schricker, B.; Fleck, R.; Wolfschmidt, H.; Stimming, U. Asymmetric Polyoxometalate Electrolytes for Advanced Redox Flow Batteries. *Energy Environ. Sci.* **2018**, *11*(10), 3010–3018. https://doi.org/10.1039/C8EE00422F.

(14) VanGelder, L. E.; Matson, E. M. Heterometal Functionalization Yields Improved Energy Density for Charge Carriers in Nonaqueous Redox Flow Batteries. *J. Mater. Chem. A* **2018**, *6*(28), 13874–13882. https://doi.org/10.1039/C8TA03312A.

(15) Linnenberg, O.; Moors, M.; Solé-Daura, A.; López, X.; Bäumer, C.; Kentzinger, E.; Pyckhout-Hintzen, W.; Monakhov, K. Yu. Molecular Characteristics of a Mixed-Valence Polyoxovanadate \([V_{14}^{IV, V_6}O_{42}]^{−}\) in Solution and at the Liquid–Surface Interface. *J. Phys. Chem. C* **2017**, *121*(19), 10419–10429. https://doi.org/10.1021/acs.jpcc.7b02138.

(16) Kawasaki, N.; Wang, H.; Nakanishi, R.; Hamanaka, S.; Kitaura, R.; Shinohara, H.; Yokoyama, T.; Yoshikawa, H.; Awaga, K. Nanohybridization of Polyoxometalate Clusters
and Single-Wall Carbon Nanotubes: Applications in Molecular Cluster Batteries. *Angew. Chem. Int. Ed.* **2011**, *50* (15), 3471–3474. https://doi.org/10.1002/anie.201007264.

(17) Yang, H.; Song, T.; Liu, L.; Devadoss, A.; Xia, F.; Han, H.; Park, H.; Sigmund, W.; Kwon, K.; Paik, U. Polyaniline/Polyoxometalate Hybrid Nanofibers as Cathode for Lithium Ion Batteries with Improved Lithium Storage Capacity. *J. Phys. Chem. C* **2013**, *117* (34), 17376–17381. https://doi.org/10.1021/jp401989j.

(18) Hu, J.; Diao, H.; Luo, W.; Song, Y.-F. Dawson-Type Polyoxomolybdate Anions (P₂Mo₁₈O₆₂₆⁶⁻) Captured by Ionic Liquid on Graphene Oxide as High-Capacity Anode Material for Lithium-Ion Batteries. *Chem. Eur. J.* **2017**, *23* (36), 8729–8735. https://doi.org/10.1002/chem.201701121.

(19) Contant, R.; Ciabrini, J.-P. Stereospecific Preparations of New N-Molybdo-(18-n)-Tungsto-2-Phosphates and Related “Defect” Compounds (n = 2, 4 or 5). *Journal of Inorganic and Nuclear Chemistry* **1981**, *43* (7), 1525–1528. https://doi.org/10.1016/0022-1902(81)80330-2.

(20) Li, C.; Mizuno, N.; Yamaguchi, K.; Suzuki, K. Self-Assembly of Anionic Polyoxometalate–Organic Architectures Based on Lacunary Phosphomolybdates and Pyridyl Ligands. *J. Am. Chem. Soc.* **2019**, *141* (19), 7687–7692. https://doi.org/10.1021/jacs.9b02541.

(21) Li, C.; Mizuno, N.; Yamaguchi, K.; Suzuki, K. Self-Assembly of Anionic Polyoxometalate–Organic Architectures Based on Lacunary Phosphomolybdates and Pyridyl Ligands. *J. Am. Chem. Soc.* **2019**, *141* (19), 7687–7692. https://doi.org/10.1021/jacs.9b02541.

(22) Izzet, G.; Volatron, F.; Proust, A. Tailor-Made Covalent Organic-Inorganic Polyoxometalate Hybrids: Versatile Platforms for the Elaboration of Functional Molecular Architectures. *The Chemical Record* **2017**, *17* (2), 250–266. https://doi.org/10.1002/tcr.201600092.

(23) Anyushin, A. V.; Kondinski, A.; Parac-Vogt, T. N. Hybrid Polyoxometalates as Post-Functionalization Platforms: From Fundamentals to Emerging Applications. *Chem. Soc. Rev.* **2020**, *49* (2), 382–432. https://doi.org/10.1039/C8CS00854J.

(24) Combs-Walker, L. A.; Hill, C. L. Stabilization of the Defect (“lacunary”) Complex Polyoximolybphosphate, PMo11O397⁻. Isolation, Purification, Stability Characteristics, and Metalation Chemistry. *Inorg. Chem.* **1991**, *30* (21), 4016–4026. https://doi.org/10.1021/ic00021a010.

(25) Rusu, M.; Muresan, L.; Tomsab, A. R.; Rusuc, D.; Marcua, Gh. New Organotin Derivatives of Keggin Polyoxomolybdates. *Synthesis and Reactivity in Inorganic and Metal-Organic Chemistry* **2000**, *30* (3), 499–511. https://doi.org/10.1080/00945710009351777.

(26) Black, F. A.; Jacquart, A.; Toupalas, G.; Alves, S.; Proust, A.; Clark, I. P.; Gibson, E. A.; Izzet, G. Rapid Photoinduced Charge Injection into Covalent Polyoxometalate–Bodipy Conjugates. *Chem. Sci.* **2018**, *9* (25), 5578–5584. https://doi.org/10.1039/C8SC00862K.

(27) Benazzi, E.; Karlsson, J.; Ben M’Barek, Y.; Chabera, P.; Blanchard, S.; Alves, S.; Proust, A.; Pullerits, T.; Izzet, G.; Gibson, E. A. Acid-Triggering of Light-Induced Charge-Separation in Hybrid Organic/Inorganic Molecular Photoactive Dyads for Harnessing Solar Energy. *Inorg. Chem. Front.* **2021**, *8* (6), 1610–1618. https://doi.org/10.1039/DOQI01368D.

(28) Volatron, F.; Noël, J.-M.; Rinfray, C.; Decorse, P.; Combellas, C.; Kanoufi, F.; Proust, A. Electron Transfer Properties of a Monolayer of Hybrid Polyoxometalates on Silicon. *Journal of Materials Chemistry C* **2015**, *3* (24), 6266–6275. https://doi.org/10.1039/C5TC00074B.

(29) Laurans, M.; Dalla Francesca, K.; Volatron, F.; Izzet, G.; Guerin, D.; Vuillaume, D.; Lenfant, S.; Proust, A. Molecular Signature of Polyoxometalates in Electron Transport of Silicon-Based Molecular Junctions. *Nanoscale* **2018**, *10* (36), 17156–17165. https://doi.org/10.1039/C8NR04946G.

(30) Martin-Sabi, M.; Soriano-López, J.; Winter, R. S.; Chen, J.-J.; Vilà-Nadal, L.; Long,
D.-L.; Galán-Mascarós, J. R.; Cronin, L. Redox Tuning the Weakly-Type Polyoxometalate Archetype for the Oxygen Evolution Reaction. Nat Catal 2018, 1 (3), 208–213. https://doi.org/10.1038/s41929-018-0037-1.

(31) Prabhakaran, V.; Lang, Z.; Clotet, A.; Poblet, J. M.; Johnson, G. E.; Laskin, J. Controlling the Activity and Stability of Electrochemical Interfaces Using Atom-by-Atom Metal Substitution of Redox Species. ACS Nano 2019, 13 (1), 458–466. https://doi.org/10.1021/acsnano.8b06813.

(32) Clemente-Juan, J. M.; Coronado, E.; Gaita-Ariño, A. Magnetic Polyoxometalates: From Molecular Magnetism to Molecular Spintronics and Quantum Computing. Chem. Soc. Rev. 2012, 41 (22), 7464. https://doi.org/10.1039/c2cs35205b.

(33) Rinfray, C.; Izzet, G.; Pinson, J.; Gam Derouich, S.; Ganem, J.-J.; Combellas, C.; Kanoufi, F.; Proust, A. Electrografting of Diazonium-Functionalized Polyoxometalates: Synthesis, Immobilisation and Electron-Transfer Characterisation from Glassy Carbon. Chem. Eur. J. 2013, 19 (41), 13838–13846. https://doi.org/10.1002/chem.201302304.

(34) Gam Derouich, S.; Rinfray, C.; Izzet, G.; Pinson, J.; Gallet, J.-J.; Kanoufi, F.; Proust, A.; Combellas, C. Control of the Grafting of Hybrid Polyoxometalates on Metal and Carbon Surfaces: Toward Submonolayers. Langmuir 2014, 30 (8), 2287–2296. https://doi.org/10.1021/la500067e.

(35) Huder, L.; Rinfray, C.; Rouchon, D.; Benayad, A.; Baraket, M.; Izzet, G.; Lipp-Bregolin, F.; Lapertot, G.; Dubois, L.; Proust, A.; Jansen, L.; Duclairoir, F. Evidence for Charge Transfer at the Interface between Hybrid Phosphomolybdate and Epitaxial Graphene. Langmuir 2016, 32 (19), 4774–4783. https://doi.org/10.1021/acs.langmuir.6b00870.

(36) Rinfray, C.; Brasiliense, V.; Izzet, G.; Volatron, F.; Alves, S.; Combellas, C.; Kanoufi, F.; Proust, A. Electron Transfer to a Phosphomolybdate Monolayer on Glassy Carbon: Ambivalent Effect of Protonation. Inorganic Chemistry 2016, 55 (14), 6929–6937. https://doi.org/10.1021/acs.inorgchem.6b00485.

(37) Cadot, E.; Fournier, M.; Tézé, A.; Hervé, G. Electrochemical Properties and ESR Characterization of Mixed Valence \(\text{a-}[\text{XMO}_{3x}\text{V}_y\text{W}_6\text{O}_{10}]^{x+}\) Heteropolyanions with \(X = \text{P}^v\) and \(\text{Si}^{iv}, x = 1, 2,\) or 3. Inorg. Chem. 1996, 35 (2), 282–288. https://doi.org/10.1021/ic940908o.

(38) Amin, S. S.; Cameron, J. M.; Winslow, M.; Davies, E. S.; Argent, S. P.; Robinson, D.; Newton, G. N. A Mixed- addenda Mo/W Organofunctionalized Hybrid Polyoxometalate. Euro J of Inorganic Chem 2022, ejic.202200019. https://doi.org/10.1002/ejic.202200019.

(39) Massart, R.; Contant, R.; Fruchart, J. M.; Ciabrin, J. P.; Fournier, M. Phosphorus-31 NMR Studies on Molybdic and Tungstic Heteropolyanions. Correlation between Structure and Chemical Shift. Inorg. Chem. 1977, 16 (11), 2916–2921. https://doi.org/10.1021/ic50177a049.

(40) Knoth, W. H.; Domaille, P. J.; Farlee, R. D. Anions of the Type \([\text{RMHM}]_{2x}\text{W}_{18}\text{P}_2\text{O}_{68}^9\) and \([\text{H}_{2}\text{OCO}]_{x}\text{W}_{18}\text{P}_2\text{O}_{68}^{1+}\). A Reinvestigation of “B,Beta.-W_9PO_34^9\text{−}.” Organometallics 1985, 4 (1), 62–68. https://doi.org/10.1021/om00120a012.

(41) Dablemont, C.; Proust, A.; Thouvenot, R.; Afonso, C.; Fournier, F.; Tabet, J.-C. Investigation of the Reactivity of Amines, Organo-Hydrazines and Tolylisocyanate towards \([\text{PW}_{12}\text{X}_2\text{M}_{10}\text{O}_{40}]^{6−}\) Keggin Anions. Dalton Trans. 2005, No. 10, 1831. https://doi.org/10.1039/b500270b.

(42) Leparulo-Loftus, M. A.; Pope, M. T. Vanadium-51 NMR Spectroscopy of Tungstovanadate Polyanions. Chemical Shift and Line-Width Patterns for the Identification of Stereoisomers. Inorg. Chem. 1987, 26 (13), 2112–2120. https://doi.org/10.1021/ic00260a021.

(43) Matt, B.; Moussa, J.; Chamoreau, L.-M.; Afonso, C.; Proust, A.; Amouri, H.; Izzet, G. Elegant Approach to the Synthesis of a Unique Heteroleptic Cyclometalated Iridium(III)-Polyoxometalate Conjugate. Organometallics 2012, 31 (1), 35–38. https://doi.org/10.1021/om200910p.
(44) Rinfray, C.; Renaudineau, S.; Izzet, G.; Proust, A. A Covalent Polyoxomolybdate-Based Hybrid with Remarkable Electron Reservoir Properties. *Chem. Commun.* 2014, 50 (62), 8575–8577. https://doi.org/10.1039/C4CC03779K.

(45) Lorion, M. M.; Matt, B.; Alves, S.; Proust, A.; Poli, G.; Oble, J.; Izzet, G. Versatile Post-Functionalization of Polyoxometalate Platforms By Using An Unprecedented Range of Palladium-Catalyzed Coupling Reactions. *Chem. Eur. J.* 2013, 19 (38), 12607–12612. https://doi.org/10.1002/chem.201301694.

(46) Cadot, E.; Thouvenot, R.; Teze, A.; Herve, G. Syntheses and Multinuclear NMR Characterizations of .Alpha.-[SiMo$_2$W$_9$O$_{39}$]$^{4-}$ and .Alpha.-[SiMo$_3.XV_xW_9O_{40}$]$^{4+x-}$ (x = 1, 2) Heteropolyoxometalates. *Inorg. Chem.* 1992, 31 (20), 4128–4133. https://doi.org/10.1021/ic00046a026.

(47) Rocchiccioli-Deltcheff, C.; Fournier, M.; Franck, R.; Thouvenot, R. Vibrational Investigations of Polyoxometalates. 2. Evidence for Anion-Anion Interactions in Molybdenum(VI) and Tungsten(VI) Compounds Related to the Keggin Structure. *Inorg. Chem.* 1983, 22 (2), 207–216. https://doi.org/10.1021/ic00144a006.

(48) Thouvenot, R.; Fournier, M.; Franck, R.; Rocchiccioli-Deltcheff, C. Vibrational Investigations of Polyoxometalates. 3. Isomerism in Molybdenum(VI) and Tungsten(VI) Compounds Related to the Keggin Structure. *Inorg. Chem.* 1984, 23 (5), 598–605. https://doi.org/10.1021/ic00173a022.

(49) C. Rocchiccioli-Deltcheff; Thouvenot, R. Metal Complexes of Heteropolyanions .Alpha.-XM11O39n- with X = SiIV or PV and M = MoVI or WVI: Study of Structural Modifications of Ligand by Infrared and Raman Spectrometry. *J. Chem. research (S).* 1977, pp 46–47.

(50) Lahootun, V.; Besson, C.; Villanneau, R.; Villain, F.; Chamareau, L.; Boukeur, K.; Blanchard, S.; Thouvenot, R.; Proust, A. Synthesis and Characterization of the Keggin-Type Ruthenium-Nitrido Derivative [PW$_{11}$O$_{39}$ (RuN)]$^{4-}$ and Evidence of Its Electrophilic Reactivity. *J. Am. Chem. Soc.* 2007, 129 (22), 7127–7135. https://doi.org/10.1021/ja071137t.

(51) Lahootun, V.; Karcher, J.; Courillon, C.; Launay, F.; Mijares, K.; Maatta, E.; Proust, A. A (Nitrido)Chromium(V) Function Incorporated in a Keggin-Type Polyoxometalate: [PW$_{11}$O$_{39}$CrN]$^5-$ Synthesis, Characterization and Elements of Reactivity. *Eur. J. Inorg. Chem.* 2008, 2008 (31), 4899–4905. https://doi.org/10.1002/ejic.200800609.

(52) Duffort, V.; Thouvenot, R.; Afonso, C.; Izzet, G.; Proust, A. Straightforward Synthesis of New Polyoxometalate-Based Hybrids Exemplified by the Covalent Bonding of a Polypyridyl Ligand. *Chemical Communications* 2009, No. 40, 6062. https://doi.org/10.1039/b913475a.

(53) Matt, B.; Coudret, C.; Viala, C.; Jouvenot, D.; Loiseau, F.; Izzet, G.; Proust, A. Elaboration of Covalently Linked Polyoxometalates with Ruthenium and Pyrene Chromophores and Characterization of Their Photophysical Properties. *Inorg. Chem.* 2011, 50 (16), 7761–7768. https://doi.org/10.1021/ic200906b.

(54) Matt, B.; Xiang, X.; Kaledin, A. L.; Han, N.; Moussa, J.; Amouri, H.; Alves, S.; Hill, C. L.; Lian, T.; Musaev, D. G.; Izzet, G.; Proust, A. Long Lived Charge Separation in Iridium(ii)-Photosensitized Polyoxometalates: Synthesis, Photophysical and Computational Studies of Organometallic–Redox Tunable Oxide Assemblies. *Chem. Sci.* 2013, 4 (4), 1737. https://doi.org/10.1039/c3sc21998d.

(55) Cameron, J. M.; Fujimoto, S.; Kastner, K.; Wei, R.-J.; Robinson, D.; Sans, V.; Newton, G. N.; Oshio, H. H. Orbital Engineering: Photoactivation of an Organofunctionalized Polyoxotungstate. *Chem. Eur. J.* 2017, 23 (1), 47–50. https://doi.org/10.1002/chem.201605021.

(56) Kibler, A. J.; Newton, G. N. Tuning the Electronic Structure of Organic–Inorganic Hybrid Polyoxometalates: The Crucial Role of the Covalent Linkage. *Polyhedron* 2018, 154, 1–20. https://doi.org/10.1016/j.poly.2018.06.027.
(57) Himeno, S.; Takamoto, M. Difference in Voltammetric Properties between the Keggin-Type [XW_{12}O_{40}]^{n-} and [XMo_{12}O_{40}]^{n-} Complexes. Journal of Electroanalytical Chemistry 2002, 528 (1–2), 170–174. https://doi.org/10.1016/S0022-0728(02)00901-4.

(58) López, X.; Bo, C.; Poblet, J. M. Electronic Properties of Polyoxometalates: Electron and Proton Affinity of Mixed-Addenda Keggin and Wells–Dawson Anions. J. Am. Chem. Soc. 2002, 124 (42), 12574–12582. https://doi.org/10.1021/ja020407z.

(59) Nambu, J.; Ueda, T.; Guo, S.-X.; Boas, J. F.; Bond, A. M. Detailed Voltammetric and EPR Study of Protonation Reactions Accompanying the One-Electron Reduction of Keggin-Type Polyoxometalates, [XV^{I}M_{11}O_{40}]^{n-} (X = P, As; M = Mo, W) in Acetonitrile. Dalton Trans. 2010, 39 (31), 7364. https://doi.org/10.1039/c003248d.

(60) Sanchez, C.; Livage, J.; Launay, J. P.; Fournier, M.; Jeannin, Y. Electron Delocalization in Mixed-Valence Molybdenum Polyanions. J. Am. Chem. Soc. 1982, 104 (11), 3194–3202. https://doi.org/10.1021/ja00375a044.

(61) Altenau, J. J.; Pope, M. T.; Prados, R. A.; So, Hyunsoo. Models for Heteropoly Blues. Degrees of Valence Trapping in Vanadium(IV)- and Molybdenum(V)-Substituted Keggin Anions. Inorg. Chem. 1975, 14 (2), 417–421. https://doi.org/10.1021/ic50144a042.

(62) Sun, H.-R.; Zhang, S.-Y.; Xu, J.-Q.; Yang, G.-Y.; Shi, T.-S. Electrochemical and In-Situ UV-Visible-near-IR and FTIR Spectroelectrochemical Characterisation of the Mixed-Valence Heteropolyanion PMo_{12}O_{40}^{n-} (n=4, 5, 6, 7) in Aprotic Media. Journal of Electroanalytical Chemistry 1998, 455 (1–2), 57–68. https://doi.org/10.1016/S0022-0728(98)00150-8.

(63) Artero, V.; Proust, A. Reduction of the Phosphodecamolybdate Ion by Phosphonium Ylides and Phosphanes. Eur. J. Inorg. Chem. 2000, 2000 (11), 2393–2400. https://doi.org/10.1002/1099-0682(200011)2000:11<2393::AID-EJIC2393>3.0.CO;2-G.

(64) Varga, G. M.; Papaconstantinou, E.; Pope, M. T. Heteropoly Blues. IV. Spectroscopic and Magnetic Properties of Some Reduced Polytungstates. Inorg. Chem. 1970, 9 (3), 662–667. https://doi.org/10.1021/ic50085a045.

(65) Fruchart, J. M.; Herve, G.; Launay, J. P.; Massart, R. Electronic Spectra of Mixed Valence Reduced Heteropolyanions. Journal of Inorganic and Nuclear Chemistry 1976, 38 (9), 1627–1634. https://doi.org/10.1016/0022-1902(76)80649-5.

(66) Matt, B.; Fize, J.; Moussa, J.; Amouri, H.; Pereira, A.; Artero, V.; Izzet, G.; Proust, A. Charge Photo-accumulation and Photocatalytic Hydrogen Evolution under Visible Light at an Iridium(III)-Photosensitized Polyoxytungstate. Energy Environ. Sci. 2013, 6 (5), 1504. https://doi.org/10.1039/c3ee04352a.

(67) Vilà-Nadal, L.; Peuntinger, K.; Busche, C.; Yan, J.; Lüders, D.; Long, D.-L.; Poblet, J. M.; Guldi, D. M.; Cronin, L. Polyoxometalate {W_{18} O_{56} XO_6 } Clusters with Embedded Redox-Active Main-Group Templates as Localized Inner-Cluster Radicals. Angew. Chem. Int. Ed. 2013, 52 (37), 9695–9699. https://doi.org/10.1002/anie.201303126.

(68) Kozik, Mariusz.; Hammer, C. F.; Baker, L. C. W. Direct Determination by Tungsten-183 NMR of the Locations of Added Electrons in ESR-Silent Heteropoly Blues. Chemical Shifts and Relaxation Times in Polysite Mixed-Valence Transition Metal Species. J. Am. Chem. Soc. 1986, 108 (10), 2748–2749. https://doi.org/10.1021/ja00270a040.

(69) Kozik, Mariusz.; Casan-Pastor, Nieves.; Hammer, C. F.; Baker, L. C. W. Ring Currents in Wholly Inorganic Heteropoly Blue Complexes. Evaluation by a Modification of Evans’ Susceptibility Method. J. Am. Chem. Soc. 1988, 110 (23), 7697–7701. https://doi.org/10.1021/ja00231a019.

(70) Suau, N.; Gaita-Ariño, A.; Clemente-Juan, J. M.; Sánchez-Marín, J.; Coronado, E. Electron Delocalization in Mixed-Valence Keggin Polyoxometalates. Ab Initio Calculation of the Local Effective Transfer Integrals and Its Consequences on the Spin Coupling. J. Am. Chem. Soc. 2002, 124 (50), 15134–15140. https://doi.org/10.1021/ja027806e.
(71) Domaille, P. J. The 1- and 2-Dimensional Tungsten-183 and Vanadium-51 NMR Characterization of Isopolymetalates and Heteropolymetalates. J. Am. Chem. Soc. 1984, 106 (25), 7677–7687. https://doi.org/10.1021/ja00337a004.
(72) Mossoba, M. M.; O’Connor, C. J.; Pope, M. T.; Sinn, E.; Herve, G.; Teze, A. Mixed Valence and Magnetically Coupled Vanadate Domains in Heteropoly Tungstate Anions. J. Am. Chem. Soc. 1980, 102 (22), 6864–6866. https://doi.org/10.1021/ja00542a042.
(73) Stoll, S.; Schweiger, A. EasySpin, a Comprehensive Software Package for Spectral Simulation and Analysis in EPR. Journal of Magnetic Resonance 2006, 178 (1), 42–55. https://doi.org/10.1016/j.jmr.2005.08.013.
(74) Prados, R. A.; Meiklejohn, P. T.; Pope, M. T. Nature of Electron Delocalization in a Heteropoly Blue Anion. Evidence for Valence Trapping at Low Temperatures. J. Am. Chem. Soc. 1974, 96 (4), 1261–1263. https://doi.org/10.1021/ja00811a071.
(75) Zhang, T.; Solé-Daura, A.; Hostachy, S.; Blanchard, S.; Paris, C.; Li, Y.; Carbó, J. J.; Poblet, J. M.; Proust, A.; Guillemot, G. Modeling the Oxygen Vacancy at a Molecular Vanadium(III) Silica-Supported Catalyst. J. Am. Chem. Soc. 2018, 140 (44), 14903–14914. https://doi.org/10.1021/jacs.8b09048.
(76) Suaud, N.; Gaita-Ariño, A.; Clemente-Juan, J. M.; Coronado, E. Electron Delocalization and Electrostatic Repulsion at the Origin of the Strong Spin Coupling in Mixed-Valence Keggin Polyoxometalates: Ab Initio Calculations of the One- and Two-Electron Processes. Chem. Eur. J. 2004, 10 (16), 4041–4053. https://doi.org/10.1002/chem.200305628.
(77) Suaud, N.; López, X.; Ben Amor, N.; Bandeira, N. A. G.; de Graaf, C.; Poblet, J. M. Accuracy of Embedded Fragment Calculation for Evaluating Electron Interactions in Mixed Valence Magnetic Systems: Study of 2e-Reduced Lindqvist Polyoxometalates. J. Chem. Theory Comput. 2015, 11 (2), 550–559. https://doi.org/10.1021/ct5010005.
(78) Namazian, M.; Coote, M. L. Accurate Calculation of Absolute One-Electron Redox Potentials of Some Para -Quinone Derivatives in Acetonitrile. J. Phys. Chem. A 2007, 111 (30), 7227–7232. https://doi.org/10.1021/jp0725883.
(79) Frisch, M. J.; Trucks, G. W.; Schlegel, H. B.; Scuseria, G. E.; Robb, M. A.; Cheeseman, J. R.; Scalmani, G.; Barone, V.; Petersson, G. A.; Nakatsuji, H.; Li, X.; Caricato, M.; Marenich, A. V.; Bloino, J.; Janesko, B. G.; Gomperts, R.; Mennucci, B.; Hratchian, H. P.; Ortiz, J. V.; Izmaylov, A. F.; Sonnenberg, J. L.; Williams-Young, D.; Ding, F.; Lipparrini, F.; Egidi, F.; Goings, J.; Peng, B.; Petrone, A.; Henderson, T.; Ranasinghe, D.; Zakrzewski, V. G.; Gao, J.; Rega, N.; Zheng, G.; Liang, W.; Hada, M.; Ehara, M.; Toyota, K.; Fukuda, R.; Hasegawa, J.; Ishida, M.; Nakajima, T.; Honda, Y.; Kitao, O.; Nakai, H.; Vreven, T.; Throssell, K.; Montgomery, J. A., Jr.; Peralta, J. E.; Ogliaro, F.; Bearpark, M. J.; Heyd, J. J.; Brothers, E. N.; Kudin, K. N.; Staroverov, V. N.; Keith, T. A.; Kobayashi, R.; Normand, J.; Raghavachari, K.; Rendell, A. P.; Burant, J. C.; Iyengar, S. S.; Tomasi, J.; Cossi, M.; Millam, J. M.; Klene, M.; Adamo, C.; Cammi, R.; Ochterski, J. W.; Martin, R. L.; Morokuma, K.; Farkas, O.; Foresman, J. B.; Fox, D. J. Gaussian 16, Revision A.03; Gaussian, Inc.: Wallingford CT, 2016.
(80) Becke, A. D. Density- functional Thermochemistry. III. The Role of Exact Exchange. The Journal of Chemical Physics 1993, 98 (7), 5648–5652. https://doi.org/10.1063/1.464913.
(81) Stephens, P. J.; Devlin, F. J.; Chabalowski, C. F.; Frisch, M. J. Ab Initio Calculation of Vibrational Absorption and Circular Dichroism Spectra Using Density Functional Force Fields. J. Phys. Chem. 1994, 98 (45), 11623–11627. https://doi.org/10.1021/j100096a001.
(82) Grimme, S.; Ehrlich, S.; Goerigk, L. Effect of the Damping Function in Dispersion Corrected Density Functional Theory. J. Comput. Chem. 2011, 32 (7), 1456–1465. https://doi.org/10.1002/jcc.21759.
(83) Chai, J.-D.; Head-Gordon, M. Long-Range Corrected Hybrid Density Functionals
with Damped Atom–Atom Dispersion Corrections. *Phys. Chem. Chem. Phys.* **2008**, *10* (44), 6615. https://doi.org/10.1039/b810189b.

(84) Krukau, A. V.; Vydrov, O. A.; Izmaylov, A. F.; Scuseria, G. E. Influence of the Exchange Screening Parameter on the Performance of Screened Hybrid Functionals. *The Journal of Chemical Physics* **2006**, *125* (22), 224106. https://doi.org/10.1063/1.2404663.

(85) Hay, P. J.; Wadt, W. R. *Ab Initio* Effective Core Potentials for Molecular Calculations. Potentials for the Transition Metal Atoms Sc to Hg. *The Journal of Chemical Physics* **1985**, *82* (1), 270–283. https://doi.org/10.1063/1.448799.

(86) Francl, M. M.; Pietro, W. J.; Hehre, W. J.; Binkley, J. S.; Gordon, M. S.; DeFrees, D. J.; Pople, J. A. Self-consistent Molecular Orbital Methods, XXIII. A Polarization-type Basis Set for Second-row Elements. *The Journal of Chemical Physics* **1982**, *77* (7), 3654–3665. https://doi.org/10.1063/1.444267.

(87) Hariharan, P. C.; Pople, J. A. The Influence of Polarization Functions on Molecular Orbital Hydrogenation Energies. *Theoret. Chim. Acta* **1973**, *28* (3), 213–222. https://doi.org/10.1007/BF00533485.

(88) Hehre, W. J.; Ditchfield, R.; Pople, J. A. Self-consistent Molecular Orbital Methods. XII. Further Extensions of Gaussian-type Basis Sets for Use in Molecular Orbital Studies of Organic Molecules. *The Journal of Chemical Physics* **1972**, *56* (5), 2257–2261. https://doi.org/10.1063/1.1677527.

(89) Cancès, E.; Mennucci, B.; Tomasi, J. A New Integral Equation Formalism for the Polarizable Continuum Model: Theoretical Background and Applications to Isotropic and Anisotropic Dielectrics. *The Journal of Chemical Physics* **1997**, *107* (8), 3032–3041. https://doi.org/10.1063/1.474659.

(90) Bauernschmitt, R.; Ahlrichs, R. Treatment of Electronic Excitations within the Adiabatic Approximation of Time Dependent Density Functional Theory. *Chemical Physics Letters* **1996**, *256* (4–5), 454–464. https://doi.org/10.1016/0009-2614(96)00440-X.

(91) Marques, M. A. L.; Gross, E. K. U. TIME-DEPENDENT DENSITY FUNCTIONAL THEORY. *Annu. Rev. Phys. Chem.* **2004**, *55* (1), 427–455. https://doi.org/10.1146/annurev.physchem.55.091602.094449.
The sites for the successive reduction processes of the mixed Mo/W Keggin-type organo-tin hybrid are debated through combined spectro-electrochemical investigation and theoretical calculations.