



HAL
open science

Molecular signature of polyoxometalates in electron transport of silicon-based molecular junctions

Maxime Laurans, Kevin Dalla Francesca, Florence Volatron, Guillaume Izzet, David Guérin, Dominique Vuillaume, Stéphane Lenfant, Anna Proust

► **To cite this version:**

Maxime Laurans, Kevin Dalla Francesca, Florence Volatron, Guillaume Izzet, David Guérin, et al.. Molecular signature of polyoxometalates in electron transport of silicon-based molecular junctions. *Nanoscale*, 2018, 10 (36), pp.17156-17165. 10.1039/C8NR04946G . hal-01981040

HAL Id: hal-01981040

<https://hal.sorbonne-universite.fr/hal-01981040>

Submitted on 14 Jan 2019

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

Molecular Signature of Polyoxometalates in Electron Transport of Silicon-based Molecular Junctions

Maxime Laurans,^a Kevin Dalla Francesca,^a Florence Volatron,^{*a} Guillaume Izzet,^a David Guerin,^b Dominique Vuillaume,^{*b} Stéphane Lenfant ^{*b} and Anna Proust^{*a}

^aSorbonne Université, CNRS, Institut Parisien de Chimie Moléculaire, IPCM, 4 Place Jussieu, F-75005 Paris, France, anna.proust@sorbonne-universite.fr.

^bInstitute for Electronics Microelectronics and Nanotechnology (IEMN), CNRS, Av. Poincaré, Villeneuve d'Ascq, France, stephane.lenfant@iemn.univ-lille1.fr, dominique.vuillaume@iemn.fr.

Abstract Polyoxometalates (POMs) are unconventional electro-active molecules with a great potential for applications in molecular memories, providing efficient processing steps onto electrodes are available. The synthesis of the organic-inorganic polyoxometalate hybrids $[\text{PM}_{11}\text{O}_{39}\{\text{Sn}(\text{C}_6\text{H}_4)\text{C}\equiv\text{C}(\text{C}_6\text{H}_4)\text{N}_2\}]^{3-}$ ($M = \text{Mo}, \text{W}$) endowed with a remote diazonium function is reported together with their covalent immobilization onto hydrogenated n-Si(100) substrates. Electron transport measurements through the resulting densely-packed monolayers contacted with a mercury drop as a top electrode confirms their homogeneity. Adjustment of the current-voltage curves with the Simmons's equation gives a mean tunnel energy barrier Φ_{POM} of 1.8 eV and 1.6 eV, for the Silicon-Molecules-Metal (SMM) junctions based on the polyoxotungstates ($M = \text{W}$) and polyoxomolybdates ($M = \text{Mo}$), respectively. This follows the trend observed in the electrochemical properties of POMs in solution, the polyoxomolybdates being easier to reduce than the polyoxotungstates, in agreement with lowest unoccupied molecular orbitals (LUMOs) of lower energy. The molecular signature of the POMs is thus clearly identifiable in the solid-state electrical properties and the unmatched diversity of POM molecular and electronic structures should offer a great modularity.

Introduction

Polyoxometalates (POMs) are nano-scaled transition metal oxides displaying discrete and reversible multi-reduction processes with minimal structural rearrangement.¹⁻⁵ This makes them promising candidates to be integrated into multi-level non-volatile molecular memories.⁶⁻⁹ The transport properties of self-assembled monolayers or multilayer films of $\text{H}_3[\text{PW}_{12}\text{O}_{40}]$ electrostatically deposited onto 3-aminopropyl-triethoxysilane (APTES)-modified SiO_2/Si surfaces have been characterized in lateral or vertical molecular junctions.^{6, 10, 11} Recently the Dawson type $[\text{W}_{18}\text{O}_{54}(\text{SeO}_3)_2]^{4-}$ POM was incorporated into a flash memory transistor by drop casting around the Si nanowire channel covered with SiO_2 .^{12, 13} Surprisingly, theoretical calculations concluded that the intimate nature of the POMs is not so important since devices integrating the parent $[\text{W}_{18}\text{O}_{56}(\text{WO}_6)_2]^{10-}$ were expected to behave similarly, despite the increase of the total charge of the polyanion. In the reported examples the ultimate electrical properties depend upon the number of POM layers or the POM packing density. To improve the performances a better control of the POM/electrode interface is required. This includes controlling the POM packing density and the homogeneity of the monolayer, the POM ordering and the POM deposition mode onto the electrode (electrostatic versus covalent anchoring).

Ideally, a weak electronic coupling between the molecules and electrodes is required to favor a long data retention time of the memory (low electron transfer rate). In case of weak molecule/electrode coupling, the electronic structure of the molecules in the solid-state molecular junction is not strongly perturbed compared to the one known in solution and the electron transport properties in the junctions is mainly governed by the properties of the molecules.¹⁴⁻¹⁵ A relationship can then be established between the energetics of the molecules in the solid-state devices and in solution.^{16, 17} Similarly, the asymmetries in voltage-dependent electron transport properties (*i.e.* more efficient for one of two voltage polarities) can be related to the spatial and energy position of the molecular orbital involved in the electron transport.¹⁸⁻²⁰ By contrast, a strong molecule/electrode coupling results in severe modification of the electronic structure (molecule orbital energy shift and broadening),²¹ up to the extreme case of electrode Fermi energy pinning,^{22, 23} for which the electron transport in the molecular junction is no longer dependent on the nature of the molecules.²⁴

Owing to their great structural diversity POMs are well-suited to address the issue of the relationship between transport properties and chemical and electronic molecular structures. In this contribution, we have thus chosen two Keggin-type POMs differing only in the nature of their constitutive metals, molybdenum versus tungsten. Both have been functionalized by the same organic tether with a

remote diazonium function for covalent anchoring onto silicon substrates, yielding isostructural and isocharged hybrids. In solution molybdates are known to be reduced more easily than their tungstates analogues, which corresponds to lowest unoccupied molecular orbitals (LUMOs) of lower energy. To which extend is this trend transposable and identifiable in solid-state molecular junctions? Can we control the energetics of the interface and the charge transport by substituting Mo for W? To study these issues, we describe the covalent grafting of two POM hybrids $\text{TBA}_3[\text{PW}_{11}\text{O}_{39}\{\text{Sn}(\text{C}_6\text{H}_4)\text{C}\equiv\text{C}(\text{C}_6\text{H}_4)\text{N}_2\}]$ ($\text{M} = \text{W}$, named $\text{K}^{\text{W}}_{\text{Sn}}[\text{N}_2^+]$ and $\text{M} = \text{Mo}$, named $\text{K}^{\text{Mo}}_{\text{Sn}}[\text{N}_2^+]$) onto hydrogenated n-Si(100). Integration of monolayers of electroactive molecules in electronic memory devices has been a topic of increasing interest²⁵⁻²⁸ and studies that have made use of oxide-free silicon as bottom electrode are particularly relevant to the CMOS approach.²⁹⁻³⁴ Different procedures have been considered to immobilize molecules onto the hydrogenated Si-substrate obtained after etching of the native oxide, among which hydrosilylation,³⁵⁻³⁷ and, to a lesser extent, spontaneous grafting from aromatic diazonium.³⁸⁻⁴⁰ Dediazonation results in the formation of robust Si-C covalent bonds and densely packed monolayers of POMs connected to the substrate by conjugated tethers. These POM monolayers onto Si have been contacted by a Hg drop to form Silicon-Molecules-Metal (SMM) junctions and the current-voltage (I-V) curves have been recorded, from which we have deduced the electronic structure (molecular energy level) of the SMM junctions. We have found that the trend on the extracted LUMO energy level in the solid-state SMM is following the trend of the POM reduction potentials as determined electrochemically in solution.

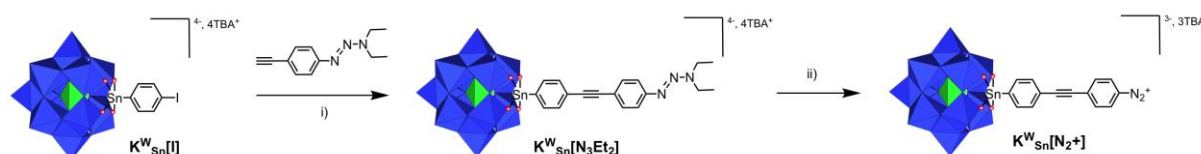
Results and discussion

Synthesis of $\text{TBA}_3[\text{PW}_{11}\text{O}_{39}\{\text{Sn}(\text{C}_6\text{H}_4)\text{C}\equiv\text{C}(\text{C}_6\text{H}_4)\text{N}_2\}]$ ($\text{K}^{\text{W}}_{\text{Sn}}[\text{N}_2^+]$)

Applications of POM-based materials in molecular nanosciences are still limited by their processing steps. Indeed, POMs are polyanions with counter cations, so that most examples of POM deposition onto electrodes rely on cation exchange^{11, 41-45} or entrapment into polymeric matrices.^{46, 47} Materials prepared by drop-casting or spin-coating have also been reported.⁴⁸⁻⁵⁰ Another strategy is to prepare organic-inorganic POM hybrids with a remote anchor able to react either with complementary functions on a pre-assembled SAM (two-step approach) or directly onto the electrode.⁵¹⁻⁵³ The two-step approach has been exemplified by the formation of metal-alkoxide bonds or the more common peptide bonds.⁵⁴⁻⁵⁷ Some of us have investigated the direct grafting and have reported on the grafting of POM hybrids terminated with protected thiols or disulfide onto Au.^{58, 59} Among the possible anchors the diazonium function has especially retained our attention because it (i) allows grafting onto various substrates, Au, carbonaceous materials and Si^{21, 60} and (ii) because dediazonation results in the formation of strong covalent bonds with the substrates, resistant to the application of wide range of external bias.⁶¹⁻⁶³ Yet formation of the diazonium function will introduce an aromatic ring in the organic tether. In previous work, we have thus described the grafting of the diazonium functionalized polyoxotungstate $\text{TBA}_3[\text{PW}_{11}\text{O}_{39}\{\text{Ge}(\text{C}_6\text{H}_4)\text{C}\equiv\text{C}(\text{C}_6\text{H}_4)\text{N}_2\}]$ ($\text{K}^{\text{W}}_{\text{Ge}}[\text{N}_2^+]$, TBA stands for tetra-butylammonium cations) onto glassy carbon and hydrogenated silicon and studied the kinetics of electron transfer at the modified electrode in solution.⁶⁴⁻⁶⁵ As charge storage nodes, polyoxomolybdates offer more perspectives than their tungsten counterparts because the cost of their reduction processes is lower (higher redox potentials).^{47, 66-69} This prompted us to prepare $\text{TBA}_3[\text{PMO}_{11}\text{O}_{39}\{\text{Sn}(\text{C}_6\text{H}_4)\text{C}\equiv\text{C}(\text{C}_6\text{H}_4)\text{N}_2\}]$ ($\text{K}^{\text{Mo}}_{\text{Sn}}[\text{N}_2^+]$), to study its electro-grafting onto glassy carbon and discuss the electrochemical behavior of the corresponding modified electrodes. However $\text{K}^{\text{W}}_{\text{Ge}}[\text{N}_2^+]$ and $\text{K}^{\text{Mo}}_{\text{Sn}}[\text{N}_2^+]$ are differing not only in the nature of the constitutive metal W/Mo but also in the nature of the metalloid substituent Ge/Sn. For synthetic reasons, we failed to prepare the exact analog $\text{K}^{\text{Mo}}_{\text{Ge}}[\text{N}_2^+]$. Although the redox properties of $\text{K}^{\text{W}}_{\text{Sn}}$ derivatives are expected to be almost indistinguishable from those of $\text{K}^{\text{W}}_{\text{Ge}}$ (as verified for example for $\text{TBA}_4[\text{PW}_{11}\text{O}_{39}\{\text{GeC}_6\text{H}_4\}]$ $\text{K}^{\text{W}}_{\text{Ge}}[\text{I}]$ and $\text{TBA}_4[\text{PW}_{11}\text{O}_{39}\{\text{SnC}_6\text{H}_4\}]$ $\text{K}^{\text{W}}_{\text{Sn}}[\text{I}]$) direct comparison of the properties of modified electrodes obtained from $\text{K}^{\text{W}}_{\text{Ge}}[\text{N}_2^+]$ on one hand, and $\text{K}^{\text{Mo}}_{\text{Sn}}[\text{N}_2^+]$ on the other hand is not completely satisfactory. In this contribution we thus describe the synthesis of the missing member of the diazonium-terminated POM family $\text{TBA}_3[\text{PW}_{11}\text{O}_{39}\{\text{Sn}(\text{C}_6\text{H}_4)\text{C}\equiv\text{C}(\text{C}_6\text{H}_4)\text{N}_2\}]$ ($\text{K}^{\text{W}}_{\text{Sn}}[\text{N}_2^+]$). As silicon is technologically relevant to the field of molecular electronics, in a hybrid molecular semi-conductor approach compatible with CMOS (Complementary metal oxide semi-conductor) technology,⁷⁰⁻⁷² $\text{K}^{\text{W}}_{\text{Sn}}[\text{N}_2^+]$ and $\text{K}^{\text{Mo}}_{\text{Sn}}[\text{N}_2^+]$ have been covalently immobilized onto n-Si(100). Probing the charge transport across the POM layer onto silicon by contacting it with a metal electrode to form a SMM junction is the next step on the road to nanoelectronic devices involving POMs. Furthermore, comparison of the electrical behavior of the W/Mo POM junctions will contribute to the understanding of the POM role in the charge transport process.

The iodo-aryl terminated hybrids $\text{K}^{\text{M}}[\text{I}]$ ($\text{M} = \text{W}, \text{Mo}$) are valuable platforms providing a large range of available POM building blocks for subsequent integration in complex architectures and assemblies.⁷³

Synthetic paths exploit the common Sonogashira cross-coupling reaction that has been adapted to the POM hybrids chemistry.⁷³⁻⁷⁶ As depicted on Scheme 1, TBA₄[PW₁₁O₃₉{Sn(C₆H₄)C≡C(C₆H₄)N₃Et₂}] (**K^W_{Sn}[N₃Et₂]**) was first prepared in 55 % yield by reaction between **K^W_{Sn}[I]** and an excess of 3,3-diethyl-1-(4-ethynylphenyl)triaz-1-ene acting as a protected diazonium group in DMF and subsequent work-up. It was characterized by IR, ¹H and ³¹P NMR, ESI-MS and elemental analysis (see experimental part and ESI). The second step consisted in the deprotection of the triazene group to release the diazonium function, achieved by adding an excess of trifluoroacetic acid (TFA) to a solution of **K^W_{Sn}[N₃Et₂]** in acetonitrile. Addition of diethylether was carried out in the presence of tetrabutylammonium hexafluorophosphate to recover the desired product as a tetrabutylammonium salt and limit its protonation. The diazonium-terminated hybrid **K^W_{Sn}[N₂⁺]** was thus obtained as a yellow powder in 79% yield and characterized by IR, ³¹P and ¹H NMR, with integration of the signals of the ¹H NMR spectrum consistent with 8 protons of the aromatic tether for 3 TBA cations. Unfortunately, no reliable ESI-MS spectrum or elemental analysis could be obtained because of the instability of the diazonium function. For this reason it is also recommended to use a freshly prepared sample. Besides the characteristic fingerprint of the [PW₁₁O₃₉{SnAr}]⁴⁻ core, the IR spectrum of **K^W_{Sn}[N₂⁺]** displays a weak band at 2255 cm⁻¹ that we assigned to the N≡N stretch of the diazonium function (a weak band at 2208 cm⁻¹ was attributed to the C≡C bond).



Scheme 1. Synthetic route to the POM hybrid **K^W_{Sn}[N₂⁺]**. In this polyhedral representation, the WO₆ octahedra are depicted with oxygen atoms at the vertices and metal cations buried inside. Color code: WO₆ octahedra, blue; PO₄ tetrahedra, green. i) [Pd(PPh₃)₂Cl₂], CuI, Et₃N, DMF overnight ii) TFA 10 min in MeCN

Grafting of **K^W_{Sn}[N₂⁺]** and **K^{Mo}_{Sn}[N₂⁺]** onto hydrogenated n-Si(100)

The just-etched Si-H substrate was thus dip-coated into a fresh solution of the diazonium-terminated POM **K^W_{Sn}[N₂⁺]** or **K^{Mo}_{Sn}[N₂⁺]**, without any supplied external bias. We have previously shown that the immersion time and rinsing procedure were critical to get a uniform monolayer with a high packing density and to avoid the formation of multilayer by competitive electrostatic deposition of extra POMs.⁶⁵ Because the diazonium route yields strong Si-C bonds, thorough rinsing and sonication is feasible without degradation of the POM monolayer, which would not be possible in the case of an electrostatic deposition of the POMs. This is an important asset of the covalent immobilization of POMs. Formation of multilayers by radical attack on the already bound aromatics is a limitation of diazonium route, albeit strategies to control the layer growth have been devised, based on bulky substituents or the presence of radical scavengers.^{77, 78} Steric hindrance is probably provided here by the POM itself. Indeed, ellipsometry measurements confirm the formation of a monolayer with a mean thickness of 2.7 and 3.2 nm for the **K^W_{Sn}** and **K^{Mo}_{Sn}** derivatized monolayers respectively. This is in good agreement with a total thickness expected between 2.5 and 3.4 nm given that the diameter of the POM is ~1 nm, the length of the organic tether is around 1.4 nm and that the TBA cations should contribute for 0.5 nm to 1 nm, depending on the conformation of the butyl chains, and tacking into account the possible 30° bending of the layer. Ellipsometry measurements were reproducible on several samples and the homogeneity of the layer was further probed by AFM in tapping mode (Fig. 1), which gave smooth images with a root-mean-square (RMS) roughness of 0.15 nm and 0.19 nm for the **K^W_{Sn}** and **K^{Mo}_{Sn}** samples respectively, similar to the n-Si substrate with a RMS roughness measured at ~ 0.25 nm and in good accordance with our previous work (see ESI).⁶⁵

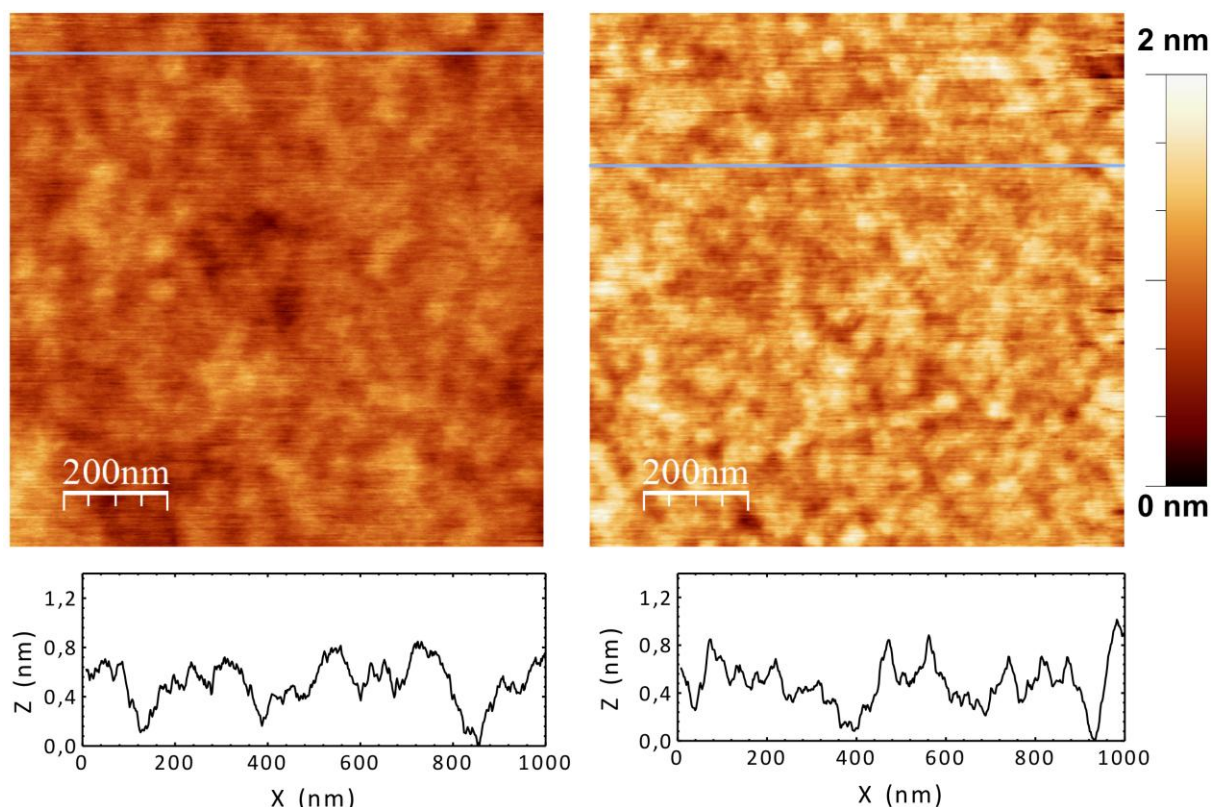


Fig. 1 Tapping-mode AFM images (up) (the scale bar corresponds to 100 nm) and height profile following the line in the AFM image (bottom) of KW_{Sn} (left) and KMo_{Sn} (right) modified n-Si(100) substrates.

Both samples were also studied by XPS and all constitutive elements of the POMs were detected: Sn, P, C, N, O and the metals, W for KW_{Sn} and Mo for KMo_{Sn} modified substrates. A part of the high resolution spectra are presented in Figure 2. For the KW_{Sn} compound, the spectrum for the 4f level of the element shows the typical spin orbit doublet at 36.1 ($\text{W}4f_{7/2}$) and 38.2 eV ($\text{W}4f_{5/2}$) corresponding to oxidized W(VI) atoms, as reported before.^{56, 65} This is confirmed by the presence of oxygen atoms bonded to metallic species, as attested by the contribution at 531.0 eV on the O1s deconvoluted peak (see - S11). The $\text{Sn}3d_{5/2}$ peak at 487.1 eV is in good agreement with oxidized Sn(IV) species as previously reported on tin oxide solids or tin derivative POM hybrid films^{79, 80} (the $\text{Sn}3d_{3/2}$ peak, not presented here for the sake of clarity, is usually spoiled by satellite peaks related to alkaline ions). The N1s and C1s photopeaks are quite similar to the ones of a diazonium terminated POM reference powder drop-casted on a silicon substrate (see Figure S12).

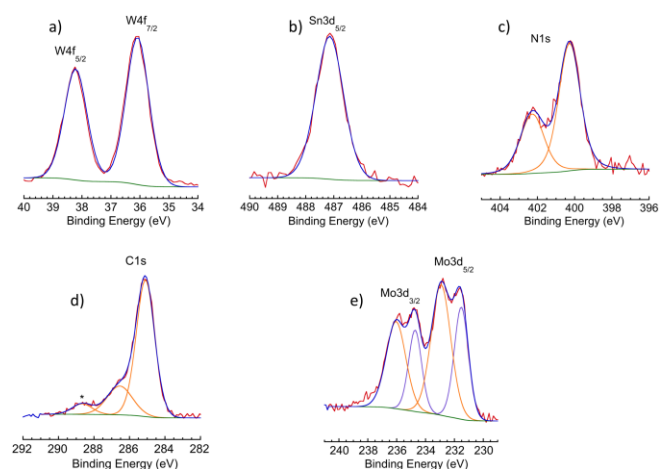


Fig. 2. High resolution X-ray photoelectron spectra for (a) W4f (b) $\text{Sn}3d_{5/2}$ (c) N1s (d) C1s in a KW_{Sn} monolayer and for (e) Mo3d in a KMo_{Sn} monolayer. The peak marked with an asterisk corresponds to a contamination by carbon atoms in an oxidized state.

On the N1s spectrum, one more contribution appears at 400.3 eV that is attributed to confined TBA counter-cations surrounding the grafted POMs. For the $\mathbf{K}^{\text{Mo}}_{\text{Sn}}$ sample, the P2p, Sn3d, O1s, N1s and C1s spectra exhibit the same features as for the $\mathbf{K}^{\text{W}}_{\text{Sn}}$ sample (see Figure S13). However, on the Mo3d spectrum, in addition to the expected Mo(VI) spin orbit doublet at 233.0 eV ($3d_{5/2}$) and 236.1 eV ($3d_{3/2}$), a new doublet at 231.5 ($3d_{5/2}$) and 234.7 eV ($3d_{3/2}$) was observed and attributed to Mo(V) atoms.⁸⁰ This means that around 40% of the grafted POMs are reduced. To understand the origin of this reduction, we compare the Mo3d spectra of a same batch of a $\mathbf{K}^{\text{Mo}}_{\text{Sn}}[\text{N}_2^+]$ powder, drop-casted on a Si/SiO₂ substrate, and used to covalently graft the $\mathbf{K}^{\text{Mo}}_{\text{Sn}}$ species on hydrogenated silicon (see Figure S14). We observe that the amount of Mo(V) is far lower (10%) in the powder reference than in the $\mathbf{K}^{\text{Mo}}_{\text{Sn}}$ layer. This means that the main contribution for the Mo reduction does come neither from the chemical synthesis, nor from the X-ray irradiation during XPS measurement. The Mo reduction may occur during the grafting on silicon. Indeed, the reduction potential of the polyoxomolybdate at -0.5 V vs SCE is close to the reduction potential of the diazonium group. Thus the hydrogenated silicon substrate, reducing agent for the diazonium function of the $\mathbf{K}^{\text{Mo}}_{\text{Sn}}[\text{N}_2^+]$ is also able to partially reduce the Mo(VI) atoms of the inorganic core.

Current-voltage (I-V) curves

Current-voltage (I-V) curves were measured by contacting the POM monolayer by a Hg drop acting as the top electrode in a glove box filled under a nitrogen flow. The drop was gently brought into contact with the sample surface thanks to a camera. The voltage V was applied on the Hg drop and the highly doped (degenerated, resistivity of $1\text{-}5 \times 10^{-3} \Omega\cdot\text{cm}$) n-type (100) silicon substrate is grounded through the ammeter to measure the current (Fig. 3a).

Figures 3-b and 3-c show 2D histograms of the 75 I-V curves measured on the SMM junctions with the mercury drop technique for the $\mathbf{K}^{\text{Mo}}_{\text{Sn}}$ and $\mathbf{K}^{\text{W}}_{\text{Sn}}$ derivatized monolayers, respectively. The 75 I-V traces were acquired at different location on the POM monolayer surface, with 1 to 5 traces per location, and they were not averaged. The two curves are quite similar with a low dispersion, inferior to one decade reflecting the homogeneity of the monolayer grafting on the substrate. However, we notice a slight difference between these two curves. The I-V curves obtained on the $\mathbf{K}^{\text{W}}_{\text{Sn}}$ monolayer are symmetric with respect to the voltage polarity, while the $\mathbf{K}^{\text{Mo}}_{\text{Sn}}$ monolayer present an absolute current value slightly higher at +1V than at -1V. However, the asymmetry is weak (rectification ratio $I_{+1\text{V}}/I_{-1\text{V}}$ is around 2.5, see Figure S15) and is not significant.^{19, 20}

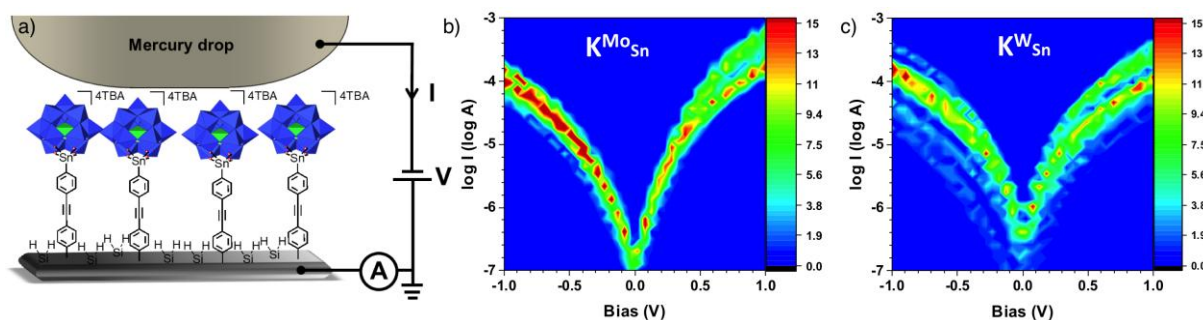


Fig. 3. a): Scheme of the molecular layer grafted on a Si substrate and the SMM junction Si- $\mathbf{K}^{\text{W}}_{\text{Sn}}$ /Hg. (b) 2D current histogram of 75 I-V curves measured for $\mathbf{K}^{\text{Mo}}_{\text{Sn}}$ and (c) $\mathbf{K}^{\text{W}}_{\text{Sn}}$ monolayers chemically grafted on a highly doped (degenerated) n-type <100> silicon substrates (resistivity of $\sim 1\text{-}5 \times 10^{-3} \Omega\cdot\text{cm}$). Voltages was applied on the mercury and Si substrate was grounded.

We analyzed the I-V curves using the Simmons model to extract the tunnel energy barrier of the monolayer placed between two electrodes^{81-84 85 86} For highly doped Si, there is no Schottky barrier related to the semiconductor and only tunneling through the molecular monolayer is operating, with a tunneling barrier height Φ_{POM} .

We extracted the barrier height and the effective mass of the monolayer directly by adjusting the I-V curve with the Simmon's equation⁸² (See Supporting Information for more details on this adjustment) since the other parameters are known: the thickness of the tunnel barrier corresponds to the monolayer thickness measured by ellipsometry and the surface contact area is estimated to $\sim 3 \times 10^{-4} \text{cm}^2$ for the mercury drop technique used (see details in ESI).

A representative I-V curve and the corresponding adjustment with the Simmon's equation are presented on Figure 4 for the $\mathbf{K}^{\text{Mo}}_{\text{Sn}}$ and $\mathbf{K}^{\text{W}}_{\text{Sn}}$ derivatized monolayers respectively. Histograms of the Φ_{POM} values extracted from these adjustments for the whole set of 75 I-V curves are shown on Figure 5a (histograms of the fitted effective mass are given in ESI, Figure S16). The histograms of Φ_{POM} are fitted by a Gaussian law, and there is a clear offset towards higher values for the $\mathbf{K}^{\text{W}}_{\text{Sn}}$ monolayers compared to the $\mathbf{K}^{\text{Mo}}_{\text{Sn}}$ ones, with mean values of $\Phi_{\text{POM}} = 1.8 \text{ eV}$ (standard deviation 0.26 eV) and $\Phi_{\text{POM}} = 1.6 \text{ eV}$ (standard deviation 0.35 eV), respectively. We note that these distributions include the raw current data distribution

(Fig. S15 in ESI) and the uncertainty of the measured thicknesses and contact area. From these average barrier height values for the both monolayers, we deduced the energy diagram of the junction (Fig. 5b), where the LUMO is localized at Φ_{POM} above the silicon Fermi energy (at 4.2 eV below the vacuum level). This value is an average between the LUMO of the POM unit itself and the one of the π -conjugated tether. However, since the tether is the same, the observed shift of Φ_{POM} is ascribed to the different chemical nature of the POM. These average values Φ_{POM} for both monolayers are, in the following, compared with the redox potentials determined by cyclic-voltammetry.

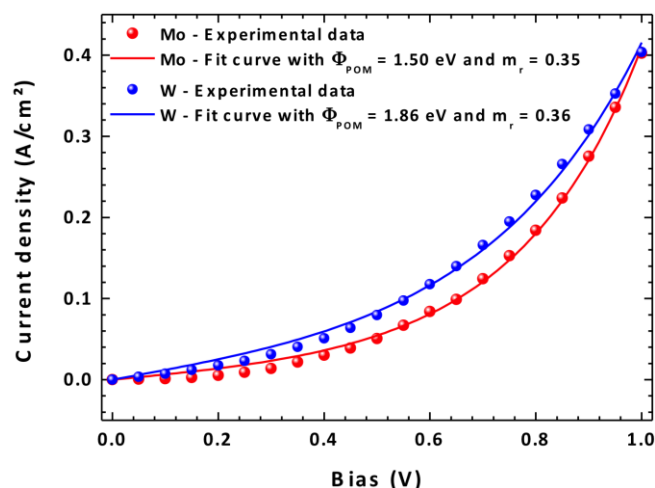


Fig. 4. Typical I-V adjustment between 0 and 1 V with the Simmons equation for the KMoSn (red curve) and KWSn (blue curve) derivatized monolayers. Adjustments of the Simmons equation with the experimental data for positive bias are good ($R > 0.98$). The other known parameters are: thickness 2.7 nm and 3.2 nm (W and Mo derivatives, respectively), contact area $3 \times 10^{-4} \text{ cm}^2$.

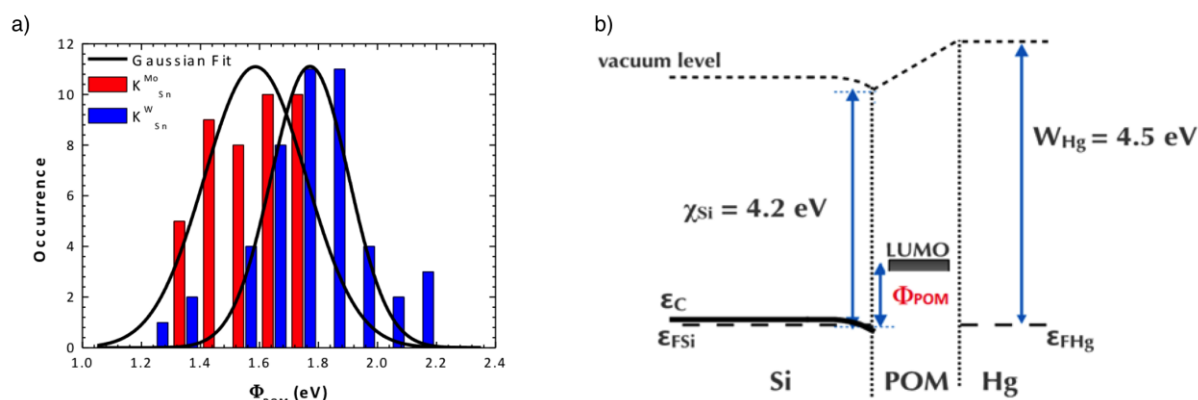


Fig. 5. (a) Barrier height (Φ_{POM}) histograms obtained by Simmons adjustments on the different I-V curves, and adjusted by a Gaussian distribution for the $\text{K}^{\text{Mo}}_{\text{Sn}}$ and $\text{K}^{\text{W}}_{\text{Sn}}$ derivatized monolayers. (b) Schematic energy diagram of the Si-POM/Hg drop as deduced from the I-V measurements.

Effect of the molecular structure of the POM

Significantly different mean values of the energy barrier Φ_{POM} have been inferred from the I-V measurements for the $\text{K}^{\text{W}}_{\text{Sn}}$ derivatized layers compared to the $\text{K}^{\text{Mo}}_{\text{Sn}}$ ones, which indicates that the POMs are indeed involved in the charge transport mechanism. Furthermore, in solution the reduction processes associated to the polyoxomolybdates are shifted to higher potentials compared to those of the analogous polyoxotungstates.⁸⁷⁻⁸⁹ The cyclic-voltammetry of $\text{K}^{\text{Mo}}_{\text{Sn}}[\text{I}]$ versus $\text{K}^{\text{W}}_{\text{Sn}}[\text{I}]$ and $\text{K}^{\text{Mo}}_{\text{Sn}}[\text{N}_3\text{Et}_2]$ versus $\text{K}^{\text{W}}_{\text{Sn}}[\text{N}_3\text{Et}_2]$ is presented in ESI as an illustration, together with the cyclic-voltammetry of the $\text{K}^{\text{W}}_{\text{Sn}}$ -derivatized layer and the relevant data gathered in Table S1. They show that (i) the redox mid-point potentials $E_{1/2}$ of molybdates are indeed about 500 mV higher than those of the corresponding tungstates, at about -0.5 V/SCE compared to -1.0 V/SCE; (ii) the nature of the tether has almost no influence on the $E_{1/2}$ values as we have already noticed on other POM hybrids;⁹⁰⁻⁹² (iii) the $E_{1/2}$ values for the POMs confined at the electrode are similar to those determined for the POMs in solution.^{58, 64, 65} An empirical linear correlation

between the Φ_{POM} and $E_{1/2}$ values has been proposed in the literature.^{20, 93} However, at variance with ferrocene, for example, the E potentials of POMs are dependent upon several parameters such as the solvent⁹⁴ and the counter-cations,^{95, 96} which might introduce some deviation. On a qualitative point of view, the matching between the relative Φ_{POM} values of $\text{K}^{\text{W}}_{\text{Sn}}$ and $\text{K}^{\text{Mo}}_{\text{Sn}}$ and their redox potential scale is however satisfying, the higher Φ_{POM} the more negative $E_{1/2}$. Note that in the case of the $\text{K}^{\text{Mo}}_{\text{Sn}}$ SMM, the value of Φ_{POM} is an average between the two oxidation states, Mo(V) and Mo(VI) observed by XPS.

Conclusions

Two diazonium-terminated POM hybrids $\text{TBA}_3[\text{PW}_{11}\text{O}_{39}\{\text{Sn}(\text{C}_6\text{H}_4)\text{C}\equiv\text{C}(\text{C}_6\text{H}_4)\text{N}_2\}]$ ($\text{K}^{\text{W}}_{\text{Sn}}[\text{N}_2^+]$) and $\text{TBA}_3[\text{PMo}_{11}\text{O}_{39}\{\text{Sn}(\text{C}_6\text{H}_4)\text{C}\equiv\text{C}(\text{C}_6\text{H}_4)\text{N}_2\}]$ ($\text{K}^{\text{Mo}}_{\text{Sn}}[\text{N}_2^+]$) differing only by the nature of their constitutive metals (W/Mo) have been prepared and covalently immobilized onto hydrogenated n-Si(100) by dediazonation to prepare $\text{K}^{\text{W}}_{\text{Sn}}$ - and $\text{K}^{\text{Mo}}_{\text{Sn}}$ -modified electrodes respectively. Solid-state molecular junctions have been closed by a Hg-top electrode and I-V curves have been recorded for the resulting SMM junctions. The I-V curves have been analyzed using the Simmons model to extract the tunneling barrier heights Φ_{POM} , which correspond to the position of the LUMOs relative to the silicon Fermi energy. The clear difference between the mean values of Φ_{POM} for the $\text{K}^{\text{W}}_{\text{Sn}}$ - and $\text{K}^{\text{Mo}}_{\text{Sn}}$ -modified electrodes, 1.8 eV and 1.6 eV respectively, follows the trend of the POM reduction potentials E as determined electrochemically in solution, with a higher energy barrier Φ_{POM} related to lower (more negative) redox potential E . We were thus able to detect the molecular signature of the POMs in the SMM junctions and this gives some insights into the energetics of the interface. Indeed, these results demonstrate a weak electronic coupling between the POMs and the Si substrate, making these electro-active molecules prone for molecular memory applications.

Experimental section

Chemicals and solvents were supplied from Aldrich or Acros and used as received, except triethylamine and acetonitrile that were distilled from CaH_2 . $\text{TBA}_4[\text{PW}_{11}\text{O}_{39}\{\text{Sn}(\text{C}_6\text{H}_4)\text{I}\}]$ ($\text{K}^{\text{W}}_{\text{Sn}}[\text{I}]$)⁹⁷ and $\text{TBA}_3[\text{PMo}_{11}\text{O}_{39}\{\text{Sn}(\text{C}_6\text{H}_4)\text{C}\equiv\text{C}(\text{C}_6\text{H}_4)\text{N}_2\}]$ ($\text{K}^{\text{Mo}}_{\text{Sn}}[\text{N}_2^+]$)⁹⁸ were prepared as previously reported (TBA stands for the tetrabutylammonium cation).

NMR spectra were recorded on a Bruker AvanceIII Nanobay 400 MHz spectrometer equipped with a BBFO probehead. ^1H chemical shifts are quoted as parts per million (ppm) relative to tetramethylsilane using the solvent signals as secondary standard (s: singlet, d: doublet, t: triplet, sex: sextet, m: multiplet) and coupling constants (J) are quoted in Hertz (Hz). ^{31}P chemical shifts are quoted relative to 85% H_3PO_4 . IR spectrum of the powder was recorded from a KBr pellet on a Jasco FT/IR 4100 spectrometer. 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 ($10 \text{ pmol}\cdot\mu\text{L}^{-1}$) were infused into the ESI source by using a syringe pump at a flow rate of $180 \mu\text{L}\cdot\text{h}^{-1}$. Mass spectra were acquired in the Orbitrap analyzer with a theoretical mass resolving power (R_p) of 100 000 at m/z 400, after ion accumulation to a target value of 10^5 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). Elemental analyses were performed at the Institut de Chimie des Substances Naturelles, Gif sur Yvette, France.

Synthesis of $\text{TBA}_4[\text{PW}_{11}\text{O}_{39}\{\text{Sn}(\text{C}_6\text{H}_4)\text{C}\equiv\text{C}(\text{C}_6\text{H}_4)\text{N}_3(\text{C}_2\text{H}_5)_2\}] \text{K}^{\text{W}}_{\text{Sn}}[\text{N}_3\text{Et}_2]$

$\text{K}^{\text{W}}_{\text{Sn}}[\text{I}]$ (199 mg, 0.050 mmol), 3,3-diethyl-1-(4-ethynylphenyl)triaz-1-ene ((33.5 mg, 0.166 mmol)), bis-(triphenylphosphine)palladium(II) dichloride (5.4 mg, 0.008 mmol) and copper iodide (1.4 mg, 0.007 mmol) were dissolved in dried and purged DMF (5 mL). The mixture was purged 5 min and freshly distilled triethylamine (150 μL , 1.133 mmol) was added. After stirring 24 hrs at room temperature, diethylether (40 mL) was added to precipitate the desired product. The recovered precipitate was dissolved in a minimum of TBABr solution (173.1 mg, 0.54 mmol in 10 mL acetonitrile). Finally, the pure product was obtained after precipitation by absolute ethanol (30 mL) and diethyl ether (2x30 mL) as beige powder (114.2 mg 55 %).

^1H NMR (400 MHz, CD_3CN): δ (ppm) 7,71 (d, $^3J_{\text{H-H}}=8,12\text{Hz}$, $^3J_{\text{Sn-H}}=96,42\text{Hz}$, 2H, Ar-H), 7,61 (d, $^3J_{\text{H-H}}=8,12\text{Hz}$, $^4J_{\text{Sn-H}}=32,72\text{Hz}$, 2H, Ar-H), 7,52 (d, $^3J_{\text{H-H}}=8,68\text{Hz}$, 2H, Ar-H), 7,38 (d, $^3J_{\text{H-H}}=8,68\text{Hz}$, 2H, Ar-H), 3,79 (q, $^3J_{\text{H-H}}=7,12\text{Hz}$, 4H, N- CH_2 - CH_3), 3,12 (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, $^3J_{\text{H-H}}=7,36\text{Hz}$, 32H, N- CH_2 - CH_2 - CH_2 - CH_3), 1,26 (m, 6H, N- CH_2 - CH_3), 0,98 (t, $^3J_{\text{H-H}}=7,36\text{Hz}$, 48H, N- CH_2 - CH_2 - CH_2 - CH_3); ^{31}P NMR (121 MHz, CD_3CN): δ (ppm) - 10,97 (s+d, $^2J_{\text{Sn-P}}=23,49\text{Hz}$); IR (KBr pellet, cm^{-1}): $\nu_{\text{C}\equiv\text{C}}=2962$ (m),

2931 (m), 2866 (m), 1482 (m), 1379 (w), 1334 (w), 1239 (w), 1071 (m), 968 (s), 884 (m), 806 (vs), 512 (w), 384 (m). HRMS (ESI-): m/z: [M]⁴⁺ calcd 768.07 found 768.07 ; [M+TBA]³⁻ calcd 1104.86 found 1104.86; [M+2TBA]²⁻ calcd 1778.48 found 1778.93 for C₁₈H₁₉SnN₃O₃₉PW₁₁. Anal. calcd for PSnW₁₁O₃₉C₈₂H₁₆₂N₇ (%): C 24.36, H 4.04, N 2.44; found : C 24.05, H 3.98, N 2.33.

Synthesis of TBA₃[PW₁₁O₃₉{Sn(C₆H₄)C≡C(C₆H₄)N₂}] K^W_{Sn}[N₂⁺]

Trifluoroacetic acid (TFA, 3.8 μL, 0.050 mmol) was slowly added to a solution of K^W_{Sn}[N₃Et₂] (40.9 mg, 0.010 mmol) in dried acetonitrile (2 mL) and the solution was stirred at room temperature for 10 minutes. A brown unidentified solid was filtered off the yellow solution and tetrabutylammonium hexafluorophosphate (TBAPF₆, 77.5 mg, 0.200 mmol) was added to the filtrate. The desired product was precipitated by dropping the filtrate into diethylether (20 mL). The yellow powder was dried at air for 30 minutes (31.1 mg, 79%). K^W_{Sn}[N₂⁺] is stored in a sealed flask at -30°C.

¹H NMR (400 MHz, CD₃CN, 298°K): δ(ppm): 8.51 (d, ³J_{H-H}=9.10Hz, 2H, Ar-H), 7.93 (d, ³J_{H-H}=9.10Hz, 2H, Ar-H), 7.75 (d, ³J_{H-H}=8.23Hz, 2H, Ar-H), 7.71 (d, ³J_{H-H}=8.23, 2H, Ar-H), 3.13 (m, 24H, N-CH₂-CH₂-CH₂-CH₃), 1.62 (m, 24H, N-CH₂-CH₂-CH₂-CH₃), 1.37 (sex, ³J_{H-H}=7.33Hz, 24H, N-CH₂-CH₂-CH₂-CH₃), 0.99 (t, ³J_{H-H}=7.33Hz, 36H, N-CH₂-CH₂-CH₂-CH₃); ³¹P NMR (121.5 MHz, CD₃CN, 298°K): δ(ppm) -10.96 (s+d, ²J_{Sn-P}=23,90Hz); IR (KBr pellet, cm⁻¹): ν̄ = 2960 (m), 2933 (m), 2872 (m), 2255 (w), 2208 (w), 1571 (m), 1482 (m), 1380 (w), 1070 (m), 963 (s), 886 (m), 795 (vs), 515 (w), 380 (m), 332 (w).

Surface covalent grafting

Pieces of silicon wafer (Highly phosphorous-doped n-Si(100) wafers purchased from Siltronix resistivity 1-5x10⁻³ Ω.cm) were sonicated for 5 min in dichloromethane and rinsed with absolute ethanol. They were then dipped two times in baths with the following compositions (i) 15 min piranha solution (mixture 2:1 of concentrated sulphuric acid and 30 wt% hydrogen peroxide solution) and (ii) 2 minutes in 5 % hydrofluoric acid (HF). After the last bath of HF, the hydrophobic dried substrate was directly dipped into a fresh 10⁻³ M solution of K^M_{Sn}[N₂⁺] (M = W or Mo) in distilled and degassed acetonitrile. The system was left under inert atmosphere at room temperature for about 1 hour (the dipping time was adjusted depending on the ellipsometry results), upon which the substrate was thoroughly rinsed with a flux of acetonitrile and sonicated for 1 minute in a bath of acetonitrile. Finally the substrate was dried under nitrogen flow and preserved from oxidation under inert atmosphere.

Caution: Piranha solution can be explosive in presence of organic compounds and HF is extremely toxic and corrosive. Manipulate with appropriate care.

Ellipsometry Ellipsometry measurements were performed on a 1*1 cm² sample and obtained using a monowavelength ellipsometer SENTECH SE 400 equipped with a He-Ne laser at λ = 632.8 nm. The incident angle was 70°. The values ns = 3.875 and ks = 0.018 were taken for the silicon wafer,⁹⁹ and ns = 1.48 and ks = 0 for the layer of POMs.¹⁰⁰ At least 6 measurements were performed on a same sample in different zones, to check the homogeneity of the layer. A mean value for the thickness was calculated when the standard deviation was lower than 0.2 nm.

AFM

The surface morphology of the POM monolayer was determined by imaging with a Dimension Icon atomic force microscope (AFM) from Bruker in tapping mode. Silicon cantilevers (Arrow™ NC from Nanoworld; k = 42 N.m⁻¹, radius < 10 nm) were used to acquire AFM images of 1 x 1 μm² at 1 Hz with a resolution of 512 x 512 pixels.

XPS

XPS analyses were performed using an Omicron Argus X-ray photoelectron spectrometer. The monochromated AlK_α radiation source (hν = 1486.6 eV) had a 300 W electron beam power. The emission of photoelectrons from the sample was analyzed at a takeoff angle of 90° under ultra-high vacuum conditions (≤ 10⁻¹⁰ Torr). Spectra were carried out with a 100 eV pass energy for the survey scan and 20 eV pass energy for the C1s, O1s, N1s regions. Binding energies were calibrated against the Si2p binding energy at 99.4 eV and element peak intensities were corrected by Scofield factors. The spectra were fitted using Casa XPS v.2.3.15 software (Casa Software Ltd., U.K.) and applying a Gaussian/Lorentzian ratio G/L equal to 70/30.

I-V measurements

We used a hanging mercury drop as top electrode, to contact electrically the POM monolayer grafted on a highly doped n-Si substrate. Calibrated mercury drops (99.9999%, purchased from Fluka) were generated by

a controlled growth mercury electrode system (CGME model from BASi) placed inside a glove box purged with a nitrogen flow. The mechanical contact between the sample and the hanging mercury drop was formed by moving up a precision lab-lift (supporting the sample) under the control of a digital video camera. The electrical contact area estimated by capacitance measurement on calibrated samples (Si/SiO₂ structure with a precisely known SiO₂ thickness of 10 nm and permittivity of 3.9 as measured by spectroscopic ellipsometry) is around 3 × 10⁻⁴ cm² (+/- 10%). The current-voltage measurements were done with an Agilent (Keysight) 4156C parameter analyzer. The voltage was always applied on the Hg top electrode, and the current measured at the grounded Si substrate.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

KDF thanks Sorbonne Universities and the program PER-SU for financial support.

References

- 1 M. T. Pope, *Heteropoly and Isopoly Oxometalates*, Springer, New-York, 1983.
- 2 M. Sadakane and E. Steckhan, *Chem. Rev.*, 1998, **98**, 219-237.
- 3 J. M. Maestre, X. Lopez, C. Bo, J. M. Poble and N. Casan-Pastor, *J. Am. Chem. Soc.*, 2001, **123**, 3749-3758.
- 4 J. M. Poble, X. Lopez and C. Bo, *Chem. Soc. Rev.*, 2003, **32**, 297-308.
- 5 M. T. Pope, in *Polyoxometalate Molecular Science*, Eds J. J. Borrás-Almenar, E. Coronado, A. Müller, M. T. Pope, Kluwer Academic Publishers, Dordrecht, pp. 3-31., 2003.
- 6 A. M. Douvas, E. Makarona, N. Glezos, P. Argitis, J. A. Mielczarski and E. Mielczarski, *ACS Nano*, 2008, **2**, 733-742.
- 7 N. Joo, S. Renaudineau, G. Delapierre, G. Bidan, L. M. Chamoreau, R. Thouvenot, P. Gouzerh and A. Proust, *Chem. Eur. J.*, 2010, **16**, 5043-5051.
- 8 B. L. Hu, C. Y. Wang, J. X. Wang, J. K. Gao, K. Wang, J. S. Wu, G. D. Zhang, W. Q. Cheng, B. Venkateswarlu, M. F. Wang, P. S. Lee and Q. C. Zhang, *Chem. Sci.*, 2014, **5**, 3404-3408.
- 9 X. L. Chen, Y. Zhou, V. A. L. Roy and S. T. Han, *Adv. Mater.*, 2018, **30**.
- 10 D. Velessiotis, A. M. Douvas, S. Athanasiou, B. Nilsson, G. Petersson, U. Sodervall, G. Alestig, P. Argitis and N. Glezos, *Microelectron. Eng.*, 2011, **88**, 2775-2777.
- 11 A. Balliou, A. M. Douvas, P. Normand, D. Tsikritzis, S. Kennou, P. Argitis and N. Glezos, *J. Appl. Phys.*, 2014, **116**, 143703.
- 12 L. Vila-Nadal, S. G. Mitchell, S. Markov, C. Busche, V. Georgiev, A. Asenov and L. Cronin, *Chem. Eur. J.*, 2013, **19**, 16502-16511.
- 13 C. Busche, L. Vila-Nadal, J. Yan, H. N. Miras, D. L. Long, V. P. Georgiev, A. Asenov, R. H. Pedersen, N. Gadegaard, M. M. Mirza, D. J. Paul, J. M. Poble and L. Cronin, *Nature*, 2014, **515**, 545-549.
- 14 J.-P. Bourgoïn, D. Vuillaume, M. Goffman and A. Filorama, *Molecular Electronics, in "Nanosciences, Nanotechnologies and Nanophysics"*, Eds C. Dupas, P. Houdy, M. Lahmani, Springer 2007, pp. 447-501., 2007.
- 15 C. C. Jia and X. F. Guo, *Chem. Soc. Rev.*, 2013, **42**, 5642-5660.
- 16 J. M. Beebe, B. Kim, J. W. Gadzuk, C. D. Frisbie and J. G. Kushmerick, *Phys. Rev. Lett.*, 2006, **97**, 4.
- 17 J. M. Beebe, B. Kim, C. D. Frisbie and J. G. Kushmerick, *ACS Nano*, 2008, **2**, 827-832.
- 18 S. Lenfant, C. Krzeminski, C. Delerue, G. Allan and D. Vuillaume, *Nano Lett.*, 2003, **3**, 741-746.
- 19 C. A. Nijhuis, W. F. Reus and G. M. Whitesides, *J. Am. Chem. Soc.*, 2010, **132**, 18386-18401.
- 20 W. F. Reus, M. M. Thuo, N. D. Shapiro, C. A. Nijhuis and G. M. Whitesides, *ACS Nano*, 2012, **6**, 4806-4822.
- 21 S. Y. Sayed, J. A. Fereiro, H. J. Yan, R. L. McCreery and A. J. Bergren, *P. Natl. Acad. Sci. USA*, 2012, **109**, 11498-11503.
- 22 S. Lenfant, D. Guerin, F. T. Van, C. Chevrot, S. Palacin, J. P. Bourgoïn, O. Bouloussa, F. Rondelez and D. Vuillaume, *J. Phys. Chem. B*, 2006, **110**, 13947-13958.
- 23 S. Rodriguez-Gonzalez, Z. Xie, O. Galangau, P. Selvanathan, L. Norel, C. Van Dyck, K. Costuas, C. D. Frisbie, S. Rigaut and J. Cornil, *J. Phys. Chem. Lett.*, 2018, **9**, 2394-2403.
- 24 H. J. Yoon, C. M. Bowers, M. Baghbanzadeh and G. M. Whitesides, *J. Am. Chem. Soc.*, 2014, **136**, 16-19.
- 25 M. Mas-Torrent, C. Rovira and J. Veciana, *Adv. Mater.*, 2013, **25**, 462-468.
- 26 C. T. Poon, D. Wu, W. H. Lam and V. W. W. Yam, *Angew. Chem. Int. Ed.*, 2015, **54**, 10569-10573.
- 27 H. Zhu, S. J. Pookpanratana, J. Y. Bonevich, S. N. Natoli, C. A. Hacker, T. Ren, J. S. Suehle, C. A. Richter and Q. L. Li, *ACS Appl. Mater. Interfaces*, 2015, **7**, 27306-27313.
- 28 H. Zhu and Q. L. Li, *Applied Sciences-Basel*, 2016, **6**.
- 29 J. E. Green, J. W. Choi, A. Boukai, Y. Bunimovich, E. Johnston-Halperin, E. Delonno, Y. Luo, B. A. Sheriff, K. Xu, Y. S. Shin, H. R. Tseng, J. F. Stoddart and J. R. Heath, *Nature*, 2007, **445**, 414-417.
- 30 Z. M. Liu, A. A. Yasseri, J. S. Lindsey and D. F. Bocian, *Science*, 2003, **302**, 1543-1545.
- 31 K. M. Roth, A. A. Yasseri, Z. M. Liu, R. B. Dabke, V. Malinovskii, K. H. Schweikart, L. H. Yu, H. Tiznado, F. Zaera, J. S. Lindsey, W. G. Kuhr and D. F. Bocian, *J. Am. Chem. Soc.*, 2003, **125**, 505-517.

- 32 J. S. Lindsey and D. F. Bocian, *Acc. Chem. Res.*, 2011, **44**, 638-650.
- 33 B. Fabre, *Acc. Chem. Res.*, 2010, **43**, 1509-1518.
- 34 D. A. Corley, T. He and J. M. Tour, *ACS Nano*, 2010, **4**, 1879-1888.
- 35 Y. Li, S. Calder, O. Yaffe, D. Cahen, H. Haick, L. Kronik and H. Zuilhof, *Langmuir*, 2012, **28**, 9920-9929.
- 36 W. J. I. DeBenedetti and Y. J. Chabal, *J. Vac. Sci. Technol. A*, 2013, **31**.
- 37 J. M. Buriak, *Chem. Mater.*, 2014, **26**, 763-772.
- 38 M. P. Stewart, F. Maya, D. V. Kosynkin, S. M. Dirk, J. J. Stapleton, C. L. McGuinness, D. L. Allara and J. M. Tour, *J. Am. Chem. Soc.*, 2004, **126**, 370-378.
- 39 M. Lu, W. A. Nolte, T. He, D. A. Corley and J. M. Tour, *Chem. Mater.*, 2009, **21**, 442-446.
- 40 C. H. deVilleneuve, J. Pinson, M. C. Bernard and P. Allongue, *J. Phys. Chem. B*, 1997, **101**, 2415-2420.
- 41 S. Q. Liu, H. Mohwald, D. Volkmer and D. G. Kurth, *Langmuir*, 2006, **22**, 1949-1951.
- 42 B. Fleury, M. Billon, F. Duclairoir, L. Dubois, A. Fanton and G. Bidan, *Thin Solid Films*, 2011, **519**, 3732-3738.
- 43 H. L. Li, S. P. Pang, S. Wu, X. L. Feng, K. Mullen and C. Bubeck, *J. Am. Chem. Soc.*, 2011, **133**, 9423-9429.
- 44 Y. C. Ji, L. J. Huang, J. Hu, C. Streb and Y. F. Song, *Energy Environ. Sci.*, 2015, **8**, 776-789.
- 45 S. M. Lauinger, J. M. Sumliner, Q. S. Yin, Z. H. Xu, G. J. Liang, E. N. Glass, T. Q. Lian and C. L. Hill, *Chem. Mater.*, 2015, **27**, 5886-5891.
- 46 Q. Liu and Z. Y. Tang, *Nano Today*, 2010, **5**, 267-281.
- 47 H. X. Yang, T. Song, L. Liu, A. Devadoss, F. Xia, H. Han, H. Park, W. Sigmund, K. Kwon and U. Paik, *J. Phys. Chem. C*, 2013, **117**, 17376-17381.
- 48 C. Musumeci, A. Luzio, C. P. Pradeep, H. N. Miras, M. H. Rosnes, Y. F. Song, D. L. Long, L. Cronin and B. Pignataro, *J. Phys. Chem. C*, 2011, **115**, 4446-4455.
- 49 M. Hutin, C. Yvon, J. Yan, A. Macdonell, D. L. Long and L. Cronin, *Crystengcomm*, 2013, **15**, 4422-4430.
- 50 L. C. Palilis, M. Vasilopoulou, A. M. Douvas, D. G. Georgiadou, S. Kennou, N. A. Stathopoulos, V. Constantoudis and P. Argitis, *Sol. Energy Mater. Sol. Cells*, 2013, **114**, 205-213.
- 51 A. Proust, B. Matt, R. Villanneau, G. Guillemot, P. Gouzerh and G. Izzet, *Chem. Soc. Rev.*, 2012, **41**, 7605-7622.
- 52 T. He, J. L. He, M. Lu, B. Chen, H. Pang, W. F. Reus, W. M. Nolte, D. P. Nackashi, P. D. Franzon and J. M. Tour, *J. Am. Chem. Soc.*, 2006, **128**, 14537-14541.
- 53 Y. C. Ji, J. Hu, L. J. Huang, W. Chen, C. Streb and Y. F. Song, *Chem. Eur. J.*, 2015, **21**, 6469-6474.
- 54 R. J. Errington, S. S. Petkar, B. R. Horrocks, A. Houlton, L. H. Lie and S. N. Patole, *Angew. Chem. Int. Ed.*, 2005, **44**, 1254-1257.
- 55 Y. F. Song, N. McMillan, D. L. Long, S. Kane, J. Malm, M. O. Riehle, C. P. Pradeep, N. Gadegaard and L. Cronin, *J. Am. Chem. Soc.*, 2009, **131**, 1340-1341.
- 56 D. Mercier, S. Boujday, C. Annabi, R. Villanneau, C. M. Pradier and A. Proust, *J. Phys. Chem. C*, 2012, **116**, 13217-13224.
- 57 D. Mercier, M. Ben Haddada, M. Huebner, D. Knopp, R. Niessner, M. Salmain, A. Proust and S. Boujday, *Sensors and Actuators B-Chemical*, 2015, **209**, 770-774.
- 58 M. Yaqub, J. J. Walsh, T. E. Keyes, A. Proust, C. Rinfray, G. Izzet, T. McCormac and R. J. Forster, *Langmuir*, 2014, **30**, 4509-4516.
- 59 A. Lombana, C. Rinfray, F. Volatron, G. Izzet, N. Battaglini, S. Alves, P. Decorse, P. Lang and A. Proust, *J. Phys. Chem. C*, 2016, **120**, 2837-2845.
- 60 J. Pinson and F. Podvorica, *Chem. Soc. Rev.*, 2005, **34**, 429-439.
- 61 M. D'amours and D. Belanger, *J. Phys. Chem. B*, 2003, **107**, 4811-4817.
- 62 D. E. Jiang, B. G. Sumpter and S. Dai, *J. Am. Chem. Soc.*, 2006, **128**, 6030-6031.
- 63 D. Belanger and J. Pinson, *Chem. Soc. Rev.*, 2011, **40**, 3995-4048.
- 64 C. Rinfray, G. Izzet, J. Pinson, S. G. Derouich, J. J. Ganem, C. Combellas, F. Kanoufi and A. Proust, *Chem. Eur. J.*, 2013, **19**, 13838-13846.
- 65 F. Volatron, J. M. Noel, C. Rinfray, P. Decorse, C. Combellas, F. Kanoufi and A. Proust, *J. Mater. Chem. C*, 2015, **3**, 6266-6275.
- 66 A. Cuentas-Gallegos, R. Martinez-Rosales, M. Baibarac, P. Gomez-Romero and M. E. Rincon, *Electrochem. Commun.*, 2007, **9**, 2088-2092.
- 67 S. H. H. Wang, Y. Nishimoto, S. Irle, T. Yokoyama, H. Yoshikawa, K. Awaga, *J. Am. Chem. Soc.*, 2012, **134**, 4918-4924.
- 68 M. Yang, B. G. Choi, S. C. Jung, Y. K. Han, Y. S. Huh and S. B. Lee, *Adv. Funct. Mater.*, 2014, **24**, 7301-7309.
- 69 J. Hu, H. L. Diao, W. J. Luo and Y. F. Song, *Chem. Eur. J.*, 2017, **23**, 8729-8735.
- 70 S. P. Cummings, J. Savchenko and T. Ren, *Coord. Chem. Rev.*, 2011, **255**, 1587-1602.
- 71 B. Fabre, *Chem. Rev.*, 2016, **116**, 4808-4849.
- 72 A. Vilan and D. Cahen, *Chem. Rev.*, 2017, **117**, 4624-4666.
- 73 G. Izzet, F. Volatron and A. Proust, *Chem. Rec.*, 2017, **17**, 250-266.
- 74 B. B. Xu, Y. G. Wei, C. L. Barnes and Z. H. Peng, *Angew. Chem. Int. Ed.*, 2001, **40**, 2290-2292.
- 75 J. Kang, B. B. Xu, Z. H. Peng, X. D. Zhu, Y. G. Wei and D. R. Powell, *Angew. Chem. Int. Ed.*, 2005, **44**, 6902-6905.
- 76 V. Duffort, R. Thouvenot, C. Afonso, G. Izzet and A. Proust, *Chem. Commun.*, 2009, 6062-6064.
- 77 Y. R. Leroux, H. Fei, J. M. Noel, C. Roux and P. Hapiot, *J. Am. Chem. Soc.*, 2010, **132**, 14039-14041.
- 78 T. Menanteau, E. Levillain and T. Breton, *Chem. Mater.*, 2013, **25**, 2905-2909.
- 79 Y. Okazaki, T. Tateishi and Y. Ito, *Materials Transactions Jim*, 1997, **38**, 78-84.
- 80 L. Huder, C. Rinfray, D. Rouchon, A. Benayad, M. Baraket, G. Izzet, F. Lipp-Bregolin, G. Lapertot, L. Dubois, A. Proust, L. Jansen and F. Duclairoir, *Langmuir*, 2016, **32**, 4774-4783.
- 81 J. G. Simmons, *J. Appl. Phys.*, 1963, **34**, 1793.
- 82 J. G. Simmons, *J. Appl. Phys.*, 1963, **34**, 2581.
- 83 J. G. Simmons, *J. Appl. Phys.*, 1964, **35**, 2655.

- 84 A. Salomon, T. Boecking, O. Seitz, T. Markus, F. Amy, C. Chan, W. Zhao, D. Cahen and A. Kahn, *Adv. Mater.*, 2007, **19**, 445-450.
- 85 H. B. Akkerman, R. C. G. Naber, B. Jongbloed, P. A. van Hal, P. W. M. Blom, D. M. de Leeuw and B. de Boer, *P. Natl. Acad. Sci. USA*, 2007, **104**, 11161-11166.
- 86 A. J. Bergren, R. L. McCreery, S. R. Stoyanov, S. Gusarov and A. Kovalenko, *J. Phys. Chem. C*, 2010, **114**, 15806-15815.
- 87 S. Himeno and M. Takamoto, *J. Electroanal. Chem.*, 2002, **528**, 170-174.
- 88 S. Himeno, M. Takamoto, R. Santo and A. Ichimura, *Bull. Chem. Soc. Jpn.*, 2005, **78**, 95-100.
- 89 J. I. Nambu, T. Ueda, S. X. Guo, J. F. Boas and A. M. Bond, *Dalton Trans.*, 2010, **39**, 7364-7373.
- 90 B. Matt, C. Coudret, C. Viala, D. Jouvenot, F. Loiseau, G. Izzet and A. Proust, *Inorg. Chem.*, 2011, **50**, 7761-7768.
- 91 B. Matt, X. Xiang, A. L. Kaledin, N. N. Han, J. Moussa, H. Amouri, S. Alves, C. L. Hill, T. Q. Lian, D. G. Musaev, G. Izzet and A. Proust, *Chem. Sci.*, 2013, **4**, 1737-1745.
- 92 C. Rinfraay, S. Renaudineau, G. Izzet and A. Proust, *Chem. Commun.*, 2014, **50**, 8575-8577.
- 93 X. Lefevre, F. Moggia, O. Segut, Y. P. Lin, Y. Ksari, G. Delafosse, K. Smaali, D. Guerin, V. Derycke, D. Vuillaume, S. Lenfant, L. Patrone and B. Jousset, *J. Phys. Chem. C*, 2015, **119**, 5703-5713.
- 94 B. Keita, D. Bouaziz and L. Nadj, *J. Electrochem. Soc.*, 1988, **135**, 87-91.
- 95 V. A. Grigoriev, C. L. Hill and I. A. Weinstock, *J. Am. Chem. Soc.*, 2000, **122**, 3544-3545.
- 96 B. Chen and R. Neumann, *PCCP*, 2016, **18**, 22487-22493.
- 97 B. Matt, J. Moussa, L. M. Chamoreau, C. Afonso, A. Proust, H. Amouri and G. Izzet, *Organometallics*, 2012, **31**, 35-38.
- 98 C. Rinfraay, V. Brasiliense, G. Izzet, F. Volatron, S. Alves, C. Combellas, F. Kanoufi and A. Proust, *Inorg. Chem.*, 2016, **55**, 6929-6937.
- 99 B. V. Zeghbroke, *Principle of conductor devices*, <http://ecee.colorado.edu/~bart/book/contents.htm>.
- 100 S. G. Derouich, C. Rinfraay, G. Izzet, J. Pinson, J. J. Gallet, F. Kanoufi, A. Proust and C. Combellas, *Langmuir*, 2014, **30**, 2287-2296.