

Ni^{II}₃₆-Containing 54-Tungsto-6-Silicate: Synthesis, Structure, Magnetic and Electrochemical Studies

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Abstract: The 36-Ni^{II}-containing 54-tungsto-6-silicate, $[Ni_{36}(OH)_{18}(H_2O)_{36}(SiW_9O_{34})_6]^{6-}$ (Ni₃₆) was synthesized by a simple one-pot reaction of the Ni_2 -pivalate complex [$Ni_2(\mu$ - OH_2)(O_2CCMe_3)₄(HO_2CCMe_3)₄] with the trilacunary [SiW₉O₃₄]¹⁰⁻ polyanion precursor in water and structurally characterized by a multitude of physicochemical techniques including single-crystal XRD, FTIR, TGA, elemental analysis, magnetic and electrochemical studies. Polyanion Ni₃₆ comprises six equivalent {Ni^{II}₆SiW₉} units which are linked by Ni–O–W bridges forming a macrocyclic assembly. Magnetic studies demonstrate that the {Ni₆} building blocks in Ni₃₆ remain magnetically intact while forming a hexagonal ring with antiferromagnetic exchange interactions between adjacent {Ni₆} units. Electrochemical studies indicate that the first reduction is reversible and associated with the $W^{VI/V}$ couple, whereas the second reduction is irreversible attributed to the Ni^{II/0} couple.

Polyoxometalates (POMs) are discrete, anionic metal-oxo clusters with a large variety of structures, compositions and physicochemical properties. While plenary POMs such as $[XW_{12}O_{40}]^{n-}$ (X = P, Si, Ge) are not reactive, lacunary derivatives such as the monolacunary $[XW_{11}O_{39}]^{n-}$ or trilacunary $[XW_9O_{34}]^{n-}$ (prepared by controlled base hydrolysis of the plenary precursor) are highly reactive towards oxophilic electrophiles such as d and f block metal ions.^[1] For decades the synthesis of high-nuclearity transition metal ion-containing POMs has received much attention, mainly aiming at novel structures with associated interesting magnetic, electrochemical, catalytic and biomedical properties.^[2] From a magnetism point of view, highnuclearity magnetic POMs may possess large uniaxial anisotropy (D) and high-spin ground states (S), possibly resulting in single-molecule magnetic behaviour and relevance for highdensity magnetic data storage and nanotechnology.^[3] Some selected examples of high-nuclearity, magnetic d-metal ion- $[Mn_{40}P_{32}W_{224}O_{888}]^{144-[4c]}$ containing POMs^[4] are $[Mn_{19}(OH)_{12}(SiW_{10}O_{37})_6]^{34-[4e]}$

 $[Fe_{48}(OH)_{76}(H_2O)_{16}(H_2P_2W_{12}O_{48})_8]^{28-}$,

 $[Fe_{28}P_8W_{48}O_{248}H_{56}]^{28-,[4g]},$

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$$\label{eq:constraint} \begin{split} & [Co_{16}(OH)_{12}(PO_4)_4(PW_9O_{34})_4]^{28-} \overset{[4d]}{,} & \text{and} & [Cu_{20}CI-(OH)_{24}(H_2O)_{12}(P_8W_{48}O_{184})]^{25-} \overset{[4h]}{,} \end{split}$$

With respect to the synthesis of Ni^{II}-containing POMs, several species with different nickel-nuclearities ranging from 2 to 40 have been reported.^[5,6] Most of these species have a sandwich-type structure, with a recent member being the heptanuclear tungstosilicate [Ni₇(OH)₆(H₂O)₄(SiW₈O₃₁)₂]¹²⁻, which showed ferromagnetic coupling between the hydroxo-bridged centres.^[5a] 2013 14-nickel-containing Ni In the $[Ni_{14}(OH)_6(H_2O)_{10}(HPO_4)_4(P_2W_{15}O_{56})_4]^{34-}$ was reported, $^{[5d]}$ and in 2015 the 25-nickel-containing $[Ni_{25}(OH)_{18}(H_2O)_2(CO_3)_2(PO_4)_6(SiW_9O_{34})_6]^{50-}.^{[5b]}$ Most of the known examples involve bridging secondary ligands such as HPO₄²⁻, PO_4^{3-} , BO_3^{3-} or CO_3^{2-} .^[5] Some POMs contain { Ni_6XW_9 } (X = P, Si, Ge) units, which are connected to adjacent units via Ni-O=W bonds, resulting in different topologies.^[6a-e] The hydroxobridged {Ni₆} subunit represents a key fragment of the brucitetype sheet structure, which has also been observed in $[Mn_{19}(OH)_{12}(SiW_{10}O_{37})_6]^{34-}$ ({Mn_{19}}).^[4e] Interestingly, Long et al. used a $Zr_6O_4(OH)_4(bpydc)_6$ "lacunary" MOF to capture $\{M_{19}\}$ brucite-type sheet structures, very similar to the {Mn19} sheets, but with halide bridges rather than OH. This resembles the situation in MX₂ solid-state structures in general for X=Cl, Br, I, and OH. In general, the occupancy is around 13-17, out of the maximum calculated 19. This suggests that using lacunary POMs as capping groups is a more robust strategy than trying to fill "holes" in MOF structures, as there are no defects in the former.^[6b]

Here we report on the synthesis of a 36 Ni^{II}-containing POM comprising exclusively trilacunary 9-tungstosilicate ions and hydroxo ligands. Reaction of the dinuclear coordination com- $[Ni_2(\mu-OH_2)(O_2CCMe_3)_4(HO_2CCMe_3)_4]$ plex (Ni₂-Piv) (Piv = $HO_2CCMe_3)^{[7]}$ with the trilacunnary 9-tungstosilicate $[SiW_9O_{34}]^{10-}$ in water under hydrothermal conditions resulted in the formation of $[Ni_{36}(OH)_{18}(H_2O)_{36}(SiW_9O_{34})_6]^{6-}$ (Ni₃₆), which was hydrated sodium isolated as а salt, $Na_{6}[Ni_{36}(OH)_{18}(H_{2}O)_{36}(SiW_{9}O_{34})_{6}] \cdot 105H_{2}O$ (Na-Ni₃₆). The title compound was characterized by a multitude of physicochemical techniques including single-crystal XRD, IR, TGA, elemental analysis, magnetic and electrochemical studies. Single-crystal XRD revealed that Na-Ni₃₆ crystallized in the triclinic system with space group $P\overline{1}$. The crystallographic parameters are shown in the Supporting Information (Table S1). The polyanion Ni₃₆ consists of six {Ni₆SiW₉} subunits, each comprising a trilacunary $\{B-\alpha$ -SiW₉\} Keggin unit with six incorporated Ni^{II} ions coordinated octahedrally and bridged to each other by hydroxo ligands, and arranged in a coplanar fashion (Figure 1). The six {Ni₆SiW₉} subunits are connected to each other via Ni–O=W bonds, leading to a cyclic assembly with the {Ni₆SiW₉} subunits alternatingly pointing up and down, resulting in polyanion Ni₃₆ with S_6 point group symmetry. Interestingly, a side view of Ni₃₆ reveals six, planar {Ni₆} assemblies arranged in a coplanar cyclic fashion and sandwiched by three $\{B-\alpha-SiW_9\}$ units from above and below, respectively. Bond valence sum (BVS) calculations confirm that all the 36 nickel ions in Ni_{36} have a +2 oxidation state (Table S2), and that the bridging oxygens are monoprotonated and hence hydroxo groups (Table S3).^[8] A close inspec-

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Figure 1. Structural representation of Ni_{36} in top view (top), side view (middle), and one of the six hydroxo-bridged { Ni_6 } subunits (bottom). Color code: nickel (green), oxygen (red). The WO₆ octahedra and Si heteroatoms are shown as transparent octahedra and balls, respectively, to highlight the six { Ni_6 } subunits within Ni_{36} .

tion of the {Ni₆SiW₉} subunits in Ni₃₆ indicates that each planar Ni₆-hydroxo cluster comprises four interconnected Ni₃O₄ cubane units. Such Ni₃O₄ or Ni₄O₄ motifs are known in POM chemistry as well as in coordination complexes.^[9]

The magnetic properties of **Na-Ni**₃₆ have been measured in powdered sample employing a SQUID magnetometer, in the temperature range of 2 to 300 K and fields ranging from 0 to 7 T. The χT at room temperature is 53.1 cm³ mol⁻¹ K, which is in the expected range for six uncoupled Ni₆ units.^[10] Upon cooling, the χT profile increases, reaching a maximum χT value of 81.1 cm³ at ca. 40 K, before decreasing to 13.2 cm³mol⁻¹ K at 2 K. The upsurge indicates the existence of ferromagnetic interactions, as observed within the composing Ni₆ units. The M(B) plot shows no saturation of the magnetization, with an almost linear profile with increasing field. Expectations were that the magnetism of **Ni**₃₆ is equal to that of its six uncoupled independent Ni₆ subunits. Both, χT versus *T* as well as *M* versus *B*, see Figure 2, suggest that there is a small exchange interaction through O–W–O bridges that couple adjacent Ni₆



Figure 2. Magnetic susceptibility at B = 0.1 T and magnetization at T = 2.0 K (inset) – experimental data (symbols), ALPS QMC simulation (red curves), simulations of 6 independent Ni₆ subunits (black curves).

triangles. We model this interaction by a single exchange integral J_3 (blue in Figure 3), while leaving J_1 and J_2 as previously derived from the magnetism of the Ni₆ subunits.^[11]

The magnetism of discrete molecular objects as large as Ni_{36} cannot be modelled by exact diagonalization due to a prohibitive size of the related Hilbert space.^[12] However, the magnetic interactions of Ni_{36} are non-frustrating since J_1 and J_2 are ferromagnetic and only J_3 is antiferromagnetic; therefore, quantum Monte Carlo (QMC) calculations can be used. We simulated the magnetic observables using the program *dirloop_se* of the ALPS package.^[13] As a result, we found that a weak antiferromagnetic exchange $J_3 = -1$ cm⁻¹ connects neighboring subunits. The agreement with the experimental data is very good. Small deviations may result from not considered Ni anisotropies, possible modifications of J_1 or J_2 , or an exchange pattern that differs from Figure 3. The tiny wiggles of the red curves are due to the fact that the QMC calculations are not yet fully converged, even after several days of simulation.



Figure 3. Schematic representation of exchange interactions in Ni₃₆: J_1 (red) and J_2 (green) as in Liu,^[10] J_3 (blue) denotes the exchange between adjacent {Ni₆} triangles. A "-2J"-Hamiltonian was employed in the calculations.

The electrochemistry of Na-Ni₃₆ was carried out in aqueous solutions in a pH 7.8 acetate medium, which is close to the pH used for the synthesis. For this purpose, the stability of polyanion Ni₃₆ was assessed by cyclic voltammetry (CV). Only the tungsten centers are expected to give rise to an electrochemical response, with the Ni^{II} ions being silent at our experimental conditions.^[14] The solid Na-Ni₃₆ was first immobilized on the surface of the basal plane of the pyrolytic graphite disk (PGB) and the electrochemical responses studied in pH 7.8 acetate medium. Figure 4A and B features the CVs of Ni₃₆ obtained at 20 mV.s⁻¹, at two different reverse potentials in a pH 7.8 acetate medium. The CV of Ni₃₆ exhibits a quasireversible reduction wave at -0.980 V versus SCE (step I), follow by the second irreversible wave at -1.232 versus SCE (step II), which is very close to the solvent discharge. As shown in Figure 4B (dark-blue and dark-red curves), nearly the same cyclic voltammograms were measured after the first and the second scan without polishing the electrode showing the chemically reversible W-reduction at -0.980 V versus SCE for the first step. In contrast, the second reduction (step II) corresponds to an irreversible multielectron transfer (Figure S3, blue curve). It must be noted that at the reverse sweep, one additional anodic peak at -0.502 V versus SCE appeared (peak *). It corresponds to the anodic dissolution peak of Ni^o and indicates that during the second irreversible reduction at

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Figure 4. Cyclic voltammograms of $Na-Ni_{36}$ immobilized at a PGB electrode (d = 2 mm) in a 1 M pH 7.8 LiOAc-HAcO solution. Scan rate: 20 mV s⁻¹.

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-1.232 versus SCE, reduction of $Ni^{\rm II}$ to $Ni^{\rm 0}$ occurred, leading to decomposition of Ni_{36} , but only during the second reduction.

Comparable measurements on a 0.50 mM solution of Ni_{36} were obtained using a glassy carbon electrode (GC) in acetate medium at pH 7.8, as shown in Figure S4 (see Supporting Information). After polishing the GC electrode, similar CV curves were always obtained, even after 1 day. Moreover, no change of the UV-visible spectrum was observed (Figure S6). These results indicate that Ni_{36} is stable in acetate medium at pH 7.8.

In summary, we have synthesized and structurally characterized the 36 Ni^{II}-containing discrete, cyclic polyanion Ni₃₆, which not only contains a record number of Ni^{II} ions, but the magnetic core is constructed without the need for any secondary bridging capping groups such as phosphate, borate or carbonate. The title polyanion Ni₃₆ is prepared by reaction of the dinuclear coordination complex Ni₂-Piv with the trilacunary {SiW₉} Keggin polyanion precursor in water, and heating in a stainless-steel autoclave under autogenous pressure at 140°C for three days. The structure of Ni₃₆ can be rationalized as a cyclic assembly of six identical {Ni₆SiW₉} units, which are oriented in alternating up/down orientations, resulting in a structure with S_6 symmetry. The magnetic properties of Ni_{36} result from the combined action of ferromagnetic exchange interactions within the {Ni₆} units, as well as antiferromagnetic exchange interactions between neighboring {Ni₆} units. For low temperatures and fields this qualitatively corresponds to a hexagonal ring of weakly antiferromagnetically interacting {Ni₆} units, each $\{Ni_6\}$ unit possessing an effective spin S=6. The electrochemical properties indicate a good stability of Ni₃₆ toward the first reduction step, which is almost reversible, whereas the polyanion decomposes at the second reduction step, due to the reduction of Ni^{II} to Ni⁰.

Experimental Section

Synthesis of $Na_6[Ni_{36}(H_2O)_{36}(OH)_{18}(SiW_9O_{34})_6] \cdot 105H_2O$ (Na-Ni₃₆): The coordination complex $[Ni_2(\mu-OH_2)(O_2CCMe_3)_4(HO_2CCMe_3)_4]$ (Ni₂-**Piv**) (Piv = HO_2CCMe_3)^[7] (0.255 g, 0.272 mmol) was dissolved in H_2O (40 mL) and then $Na_{10}[A-\alpha-SiW_9O_{34}]\cdot 24H_2O$ (0.200 g, 0.068 mmol)^[15] was added with constant stirring for 10 min at room temperature. The resultant reaction mixture was transferred into a Teflon-lined stainless-steel autoclave and was heated under autogenous pressure at 140°C for three days. After overnight cooling a precipitate had formed at the bottom of the Teflon container, which was removed by centrifugation and/or filtration. The solution was kept in an open vial at room temperature to allow for slow evaporation. After about one month a green, crystalline product started to appear. Evaporation was allowed to continue until most of the solvent had disappeared. Then the solid product was collected by filtration and air dried. Yield 15 mg (7%, based on W). IR (2% KBr pellet): $\tilde{\nu} = 3412$ (br), 1619(s), 1379(sh), 947(s), 878(s), 833 (sh), 790(s), 687(s), 541(w), 513(w), 478(sh). Elemental analysis (%) for Na₆[Ni₃₆(H₂O)₃₆(OH)₁₈(SiW₉O₃₄)₆] ·105H₂O (Na-Ni₃₆), calcd: W 53.79, Ni 11.45, Na 0.75, Si 0.91; found W 51.35, Ni 12.02, Na 0.78, Si 0.77. Elemental analysis was performed at CREALINS (Villeurbanne, France), except for Na, which was determined by atomic absorption at Jacobs University.

Deposition Number 2001196 (Na-Ni₃₆) contains the supplementary crystallographic data for this paper. These data are provided free of

charge by the joint Cambridge Crystallographic Data Centre and Fachinformationszentrum Karlsruhe Access Structures service www.ccdc.cam.ac.uk/structures.

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Conflict of Interest

The authors declare no conflict of interest.

Keywords: Electrochemistry · magnetic properties · nickel · polyoxometalates · structure elucidation

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