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Dithiolene complexes as metallo-ligands: a crown-ether approach†

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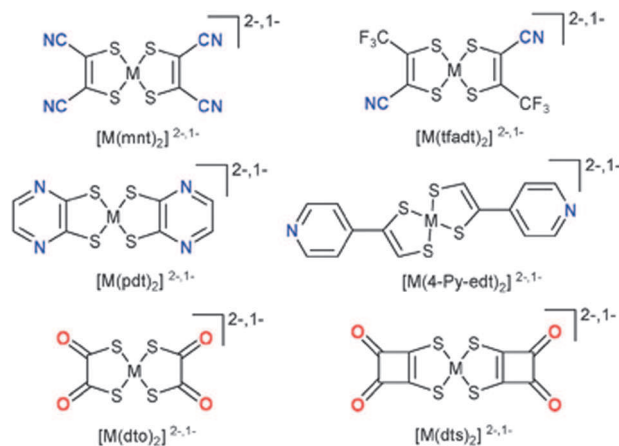
A nickel dithiolene complex substituted with crown ether cyclic moieties incorporating four O atoms, abbreviated as $[\text{Ni}(\text{S}_2\text{O}_4)_2]^{1-0}$, is isolated in its radical anionic (as Na^+ salt) and neutral forms. The Na^+ cation is six-coordinated with short $\text{Na} \cdots \text{O}$ distances (2.46–2.52 Å), involving two crown ether moieties of two different complexes. The oxidized neutral complex $[\text{Ni}(\text{S}_2\text{O}_4)_2]^0$ was also isolated and structurally characterized. In the absence of alkaline cations during the synthesis, a mixed salt associating $[\text{Ni}(\text{S}_2\text{O}_4)_2]^{1-}$ with Ni^{2+} was isolated, and formulated as $(\text{Ni}_2\text{H}_2\text{O})[\text{Ni}(\text{S}_2\text{O}_4)_2]_2$, with the Ni^{2+} cation weakly bonded to two crown ether moieties. The salt exhibits an unprecedented solid state association with extremely short $\text{S} \cdots \text{S}$ intermolecular contacts [3.332(2) Å], leading to a pairing of the radical $[\text{Ni}(\text{S}_2\text{O}_4)_2]^-$ into antiferromagnetic uniform spin chains.

Introduction

Paramagnetic metal dithiolene complexes¹ are currently extensively investigated for their conducting and magnetic properties,² with the most common magnetic species found among the $S = \frac{1}{2} d^9$ $[\text{Cu}(\text{dithiolene})_2]^{2-}$ copper complexes, the formally d^7 , $S = \frac{1}{2}$ $[\text{M}(\text{dithiolene})_2]^{1-}$ nickel, palladium or platinum anionic complexes,^{2,3} and the $[\text{Au}(\text{dithiolene})_2]^0$ gold neutral complexes.^{4,5} The electronic properties of these “simple” salts are essentially controlled by the overlapping interactions between radical species. Weak interactions favour the observation of localized magnetic states (dimers, spin chains, spin ladders, ...) while stronger interactions might allow the formation of conduction bands^{2d} and associated metallic behavior of the conductivity.

Paramagnetic dithiolene complexes can also be considered as elementary building blocks for more elaborated architectures where they would play the role of a *metallo-ligand* toward other metallic centers,^{6,7} provided that they bear substituents adapted for further outer coordination, essentially through the lone pairs of nitrogen (nitrile and pyridine functions) or oxygen (ether and carbonyl functions) atoms. For example, derivatives of nitrile containing complexes^{3,8} such as $[\text{M}(\text{mnt})_2]^{2-,1-}$ or $[\text{M}(\text{tfadt})_2]^{2-,1-}$ were found to successfully coordinate other metallic centers through nitrogen atoms,

such as the $S = 2$ Mn^{III} porphyrin, $[\text{Mn}(\text{TPP})]^+$,⁹ the $S = 1$ $[\text{Ni}(\text{cyclam})]^{2+}$ complex¹⁰ and polymetallic complexes such as $[\text{Mn}_4(\text{hmp})_6(\text{CH}_3\text{CN})_2]^{4+}$ (hmp: 2-hydroxymethylpyridinate).^{11,12} Other examples of nitrogen coordination can be also found in fused pyrazine¹³ or in 4-pyridyl derivatives.¹⁴ Complexes with oxygen-based coordinating functions are essentially limited to the chelating dithiooxalate (dto)⁶ and dithiosquarate (dts) ligands. Because of the strong electron-withdrawing nature of the carbonyl groups, both ligands only stabilize the most reduced dianionic forms of the dithiolene complexes, for example the diamagnetic $[\text{Ni}(\text{dto})_2]^{2-}$ or the paramagnetic d^9 $S = \frac{1}{2}$ $[\text{Cu}(\text{dto})_2]^{2-}$. The latter was indeed engaged in bimetallic polymeric chains with ferrimagnetic interactions upon Mn^{II} coordination.¹⁵ Other examples involved the tris(dithiolene) complexes¹⁶ such as $[\text{Cr}(\text{dto})]^{3-}$ or the polymeric $\text{N}(n\text{-C}_3\text{H}_7)_4[\text{Co}^{\text{II}}\text{Fe}^{\text{III}}(\text{dto})_3]$.¹⁷



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Looking for alternative substitution patterns which would work with the monoanionic, formally d^7 , $S = \frac{1}{2}$ $[\text{Ni}(\text{dithiolene})_2]^-$ complexes while keeping oxygen atoms for secondary coordination, we envisioned crown ether derivatives as they would provide a powerful coordination function, particularly toward oxophilic cationic centers. Their structure is shown in Scheme 1 with the corresponding abbreviations used in the following.

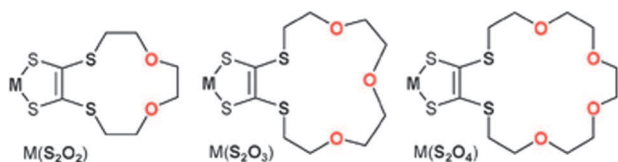
Dithiolene complexes bearing such a crown ether functionality were first described in 1990 by Green,¹⁸ aiming at coupling the rich redox chemistry of dithiolene complexes with the cation binding ability of the crown ether group, as demonstrated in the heteroleptic $\text{Cp}_2\text{M}(\text{S}_2\text{O}_3)$, $\text{M} = \text{Mo}, \text{W}$ complexes, and the square planar diamagnetic $[\text{Cu}(\text{S}_2\text{O}_3)_2]^-$ species. This attractive approach was also extended to the corresponding tetrathiafulvalene derivatives,^{19,20} showing also notable electrochemical anodic shifts of the oxidation potentials upon alkaline cation coordination.²¹ However, the coordination ability of these crown ether derivatives has been strictly restricted up to now to alkaline cations: other, eventually paramagnetic, metal cations were never considered. This is all the more unfortunate that these dithiolene complexes can be isolated into various paramagnetic states, with a spin density largely delocalized on the dithiolate ligands while the ether functionalities of the crown ether substituents would be particularly adapted to the coordination of oxophilic metal centres.

We describe here our first results along this line with the synthesis of the crown-ether substituted paramagnetic complex abbreviated as $[\text{Ni}(\text{S}_2\text{O}_4)_2]^{1-}$, isolated as its Na^+ salt and structurally characterized. In the absence of alkaline cations during the synthesis, a mixed salt associating $[\text{Ni}(\text{S}_2\text{O}_4)_2]^-$ units with Ni^{2+} was isolated, showing an unprecedented solid state association with short S...S intermolecular contacts, leading to a pairing of the radical $[\text{Ni}(\text{S}_2\text{O}_4)_2]^-$ into antiferromagnetic uniform spin chains.

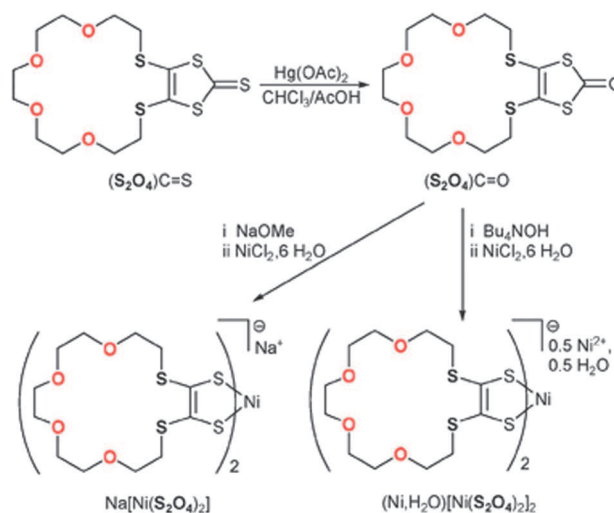
Results and discussion

The preparation of the nickel complex (Scheme 2) relies on the base opening of the dithiocarbonate $(\text{S}_2\text{O}_4)\text{C}=\text{O}$, obtained itself from the oxymercuration of known trithiocarbonate^{19b} $(\text{S}_2\text{O}_4)\text{C}=\text{S}$ in 60% yield. The dithiocarbonate was characterized by single crystal X-ray diffraction (Fig. 1).

When NaOMe is used as a base to produce the free dithiolate from the dithiocarbonate $(\text{S}_2\text{O}_4)=\text{O}$, followed by addition of half an equivalent $\text{NiCl}_2 \cdot 6\text{H}_2\text{O}$, the initially formed $[\text{Ni}(\text{S}_2\text{O}_4)_2]^{2-}$ oxidizes in air to afford the monoanionic complex, isolated as its sodium salt, that is, $\text{Na}[\text{Ni}(\text{S}_2\text{O}_4)_2]$. On the other hand, using (*n*-Bu)₄NOH as a base instead of NaOMe afforded the Ni^{2+} salt, formulated as $[\text{Ni}(\text{H}_2\text{O})_{0.5}[\text{Ni}(\text{S}_2\text{O}_4)_2]$. The oxidized neutral complex $[\text{Ni}(\text{S}_2\text{O}_4)_2]^0$ was obtained from iodine oxidation of the latter monoanionic species.



Scheme 1 Crown-ether dithiolene complexes and their acronyms.



Scheme 2 Synthesis of nickel complexes.

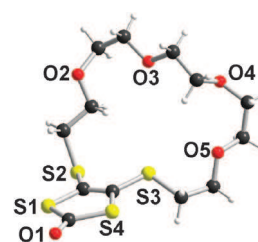


Fig. 1 X-Ray crystal structure of the dithiocarbonate $(\text{S}_2\text{O}_4)\text{C}=\text{O}$.

Cyclic voltammetry experiments performed on $\text{Na}[\text{Ni}(\text{S}_2\text{O}_4)_2]$ (Fig. 2) show two reversible redox waves associated with the $[\text{Ni}(\text{S}_2\text{O}_4)_2]^{-2/-1}$ and $[\text{Ni}(\text{S}_2\text{O}_4)_2]^{-1/0}$ redox processes, at $E_{1/2} = -0.64$ and $+0.13$ V (vs. SCE) respectively. A third redox process observed at $E_{\text{ox}} = +0.78$ V and associated to the oxidation of the neutral complex is irreversible. Note that the rather low oxidation potential of the monoanionic $[\text{Ni}(\text{S}_2\text{O}_4)_2]^{1-}$ ($+0.13$ V) gives an explanation for its experimentally observed easy oxidation to the neutral complex.

$\text{Na}[\text{Ni}(\text{S}_2\text{O}_4)_2]$ crystallizes in the monoclinic system, space group $C2/c$ with the Ni atom on an inversion centre while the Na ion lies on a two-fold axis. As shown in Fig. 3, the Na^+ cation is coordinated by six oxygen atoms from two

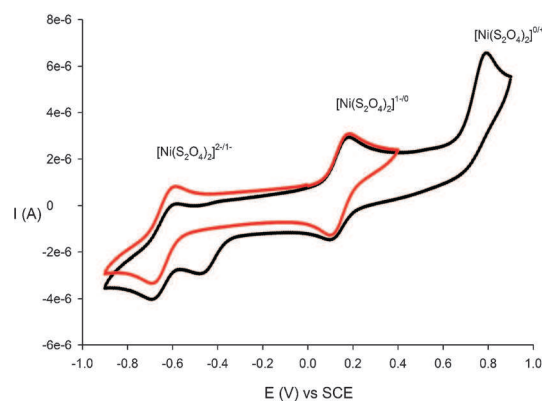


Fig. 2 Cyclic voltammogram of $\text{Na}[\text{Ni}(\text{S}_2\text{O}_4)_2]$, in DMF with 0.2 M (*n*-Bu₄N)PF₆, at 100 mV s^{-1} scan rate.

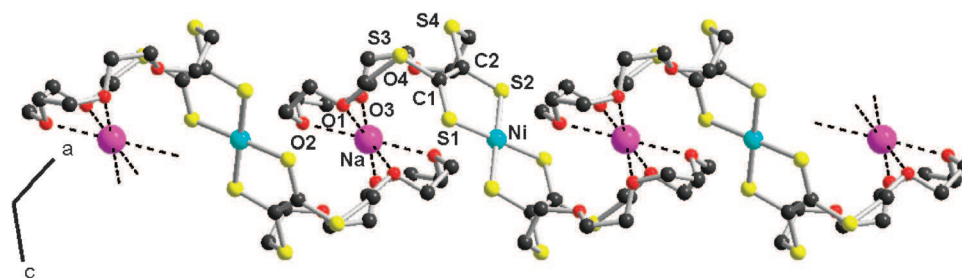


Fig. 3 View of the $\text{Na}[\text{Ni}(\text{S}_2\text{O}_4)_2]$ complex, showing the Na^+ coordination pattern.

neighbouring dithiolene complexes, with $\text{Na}\cdots\text{O}$ distances of 2.463(4), 2.523(6) and 2.4651(4) Å with O1, O2 and O3 respectively, giving rise to coordination chains running along $a + c$, and demonstrating that the crown ether functions can effectively act as secondary ligands in these dithiolene complexes. The $\text{Na}\cdots\text{O}$ distances are particularly short when compared with those observed for example in $[(18\text{-crown-6})\text{Na}^+]$ derivatives where they are usually found around 2.50–2.90 Å. An interesting comparison can also be made with the only structurally reported analog incorporating two sulfur atoms, shown in Fig. 4, where the eight-coordinated Na^+ cation is sandwiched between two dithiatetraoxa crown molecules.²² In this salt, the $\text{Na}\cdots\text{O}$ distances are even longer, between 2.85 and 3.11 Å.

Intramolecular bond distances within the metallacycle in $[\text{Ni}(\text{S}_2\text{O}_4)_2]^{1-}$ (Table 1) are characteristic of radical anion nickel dithiolene complexes such as the analogous thioalkyl substituted $[\text{Ni}(\text{ddd}t)_2]^-$,²⁵ $[\text{Ni}(\text{pdd}t)_2]^-$,²³ or $[\text{Ni}(\text{F}_2\text{pdd}t)_2]^-$ complexes.^{24,27}

Because of its limited solubility, EPR spectra of the sodium salt in solution could be obtained only from DMF solutions with added CH_2Cl_2 . At room temperature, a single Lorentzian line is observed with $g = 2.055$ and a 9.1 G linewidth. In frozen solution (Fig. 5), a rhombic pattern is found with $g_{\text{max}} = 2.102$, $g_{\text{int}} = 2.040$ and $g_{\text{min}} = 2.001$, comparable to those described for $[\text{Ni}(\text{mnt})_2]^-$ (2.14, 2.04, 1.99), $[\text{Ni}(\text{dmit})_2]^-$ (2.105, 2.041, 2.001)²⁸ or $[\text{Ni}(\text{ddd}t)_2]^-$ [2.119(1), 2.057(1), 2.022(1)].²⁹ The powder spectrum of $\text{Na}[\text{Ni}(\text{S}_2\text{O}_4)_2]$ shows an axial symmetry with $g_{\perp} = 2.071$ and $g_{\parallel} = 2.044$. In the solid state, the radical complexes are fully isolated from each other and the shortest intermolecular S \cdots S distances exceed 4.88 Å. This is also confirmed by the temperature dependence of the magnetic susceptibility, which follows a Curie law in the whole temperature range.

As mentioned above, the $[\text{Ni}(\text{S}_2\text{O}_4)_2]^{1-}$ radical anion is oxidation sensitive and many attempts to replace the Na^+ cation with other metallic cations very often afforded the neutral oxidized complex $[\text{Ni}(\text{S}_2\text{O}_4)_2]^0$. The latter was found to crystallize in the triclinic system, space group $P\bar{1}$ with the

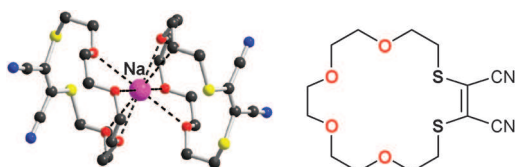
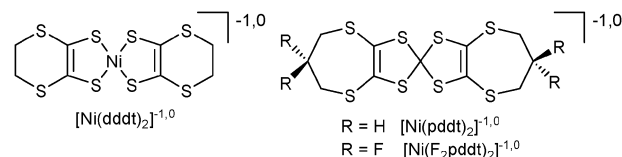


Fig. 4 The structurally characterized Na^+ salt of a dithia-tetraoxa crown.²²

Table 1 Intramolecular bond distances within the $[\text{Ni}(\text{S}_2\text{O}_4)_2]^{n-}$ and analogous reference compounds known in both oxidation states such as $[\text{Ni}(\text{ddd}t)_2]^{n-}$, $[\text{Ni}(\text{pdd}t)_2]^{n-}$ and $[\text{Ni}(\text{F}_2\text{pdd}t)_2]^{n-}$ ($n = 1, 0$)



	Charge	Ni-S	S-C	C=C	Ref.
$\text{Na}[\text{Ni}(\text{S}_2\text{O}_4)_2]$	-1	2.142(4) 2.155(2)	1.726(6) 1.728(5)	1.370(6)	This work
$(\text{Ni}, \text{H}_2\text{O})_{0.5}[\text{Ni}(\text{S}_2\text{O}_4)_2]$	-1	2.140(1)	1.729(4)	1.367(5)	This work
$\text{Et}_4\text{N}[\text{Ni}(\text{ddd}t)_2]$	-1	2.147(2)	1.735(8)	1.339(11)	29
$\text{Et}_4\text{N}[\text{Ni}(\text{pdd}t)_2]$	-1	2.142(10)	1.725(20)	1.356(3)	23
$\text{Bu}_4\text{N}[\text{Ni}(\text{F}_2\text{pdd}t)_2]$	-1	2.150(1)	1.735(4)	1.346(5)	24
$[\text{Ni}(\text{S}_2\text{O}_4)_2]^0$	0	2.134(1) 2.136(1)	1.708(2) 1.712(2)	1.399(3)	This work
$[\text{Ni}(\text{ddd}t)_2]^0$	0	2.121(5)	1.69(2)	1.37(2)	25
$[\text{Ni}(\text{pdd}t)_2]^0$	0	2.133(1)	1.712(3)	1.401(4)	26
$[\text{Ni}(\text{F}_2\text{pdd}t)_2]^0$	0	2.133(1)	1.698(3)	1.376(4)	27

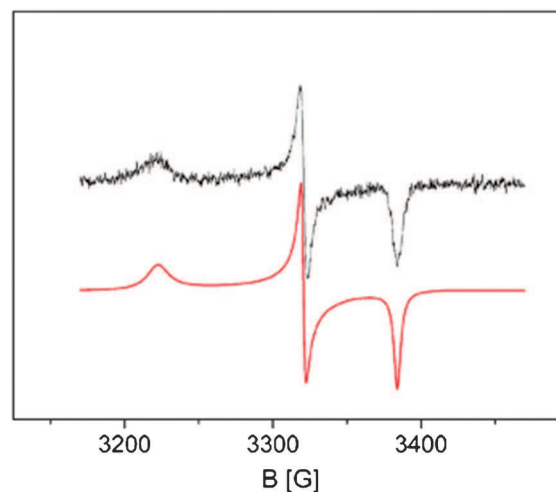


Fig. 5 Frozen solution EPR spectrum of $\text{Na}[\text{Ni}(\text{S}_2\text{O}_4)_2]$ (in DMF/ CH_2Cl_2) with fit in red line (see text).

nickel atom on the inversion centre (Fig. 6). Intramolecular bond distance within the NiS_2C_2 metallacycle (Table 1) confirms the oxidation state of the complex, with a notable shortening of the C-S and lengthening of the C=C bonds when compared with the $[\text{Ni}(\text{S}_2\text{O}_4)_2]^{1-}$ radical anion, associated with the partial oxidation of the two dithiolate ligands.

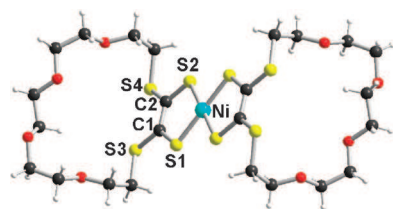


Fig. 6 View of the neutral oxidized complex $[\text{Ni}(\text{S}_2\text{O}_4)_2]^0$.

When the dithiolene complex is prepared in the absence of Na^+ cations, using Bu_4NOH rather than NaOMe , the $[\text{Ni}(\text{S}_2\text{O}_4)_2]^{-1}$ radical complex was isolated as Ni^{2+} salt, together with a water molecule. The salt, formulated as $(\text{Ni}, \text{H}_2\text{O})_{0.5}[\text{Ni}(\text{S}_2\text{O}_4)_2]$, crystallizes in the monoclinic system, space group $C2/c$. The dithiolene complex is located on an inversion centre while the Ni^{2+} cation and water molecule are disordered on the same position, on the two-fold axis (Fig. 7). Intramolecular bond distances within the NiS_2C_2 metallacycle (Table 1) are identical to those observed for the sodium salt, confirming the actual formulation. The very peculiar solid state organization parallels that which was observed with the Na^+ salt, where the sodium site would be alternatively occupied by a Ni^{2+} or a water molecule, allowing for lattice electro-neutrality. At variance however with the structure of $\text{Na}[\text{Ni}(\text{S}_2\text{O}_4)_2]$, the coordination of the $\text{Ni}^{2+}/\text{H}_2\text{O}$ centers by the crown ether oxygen atoms is much weaker, with $\text{Ni}\cdots\text{O}$ distances of 2.977(3), 2.885(4) and 2.898(3) Å with O1, O2 and O3 respectively. These distances are larger than those observed above in $\text{Na}[\text{Ni}(\text{S}_2\text{O}_4)_2]$ with the Na^+ cation and also than those reported in the Ni^{2+} crown ether complexes such as $[\text{Ni}(18\text{-crown-6})(\text{EtOH})_3](\text{PF}_6)_2$ where three out of the six available O atoms coordinate Ni^{2+} with $\text{Ni}\cdots\text{O}$ distance around 2.03–2.11 Å,³⁰ or $[\text{Ni}(15\text{-crown-5})(\text{H}_2\text{O})_2](\text{NO}_3)_2$ and $[\text{Ni}(15\text{-crown-5})(\text{H}_2\text{O})_2](\text{HSO}_4)_2$ with $\text{Ni}^{2+}\cdots\text{O}$ distances 2.06–2.30 Å.³¹ This tends to indicate that the Ni^{2+} coordination is not the only driving force directing the supramolecular arrangement of the structure.

From a magnetic point of view, $(\text{Ni}, \text{H}_2\text{O})[\text{Ni}(\text{S}_2\text{O}_4)_2]$ can be described as chains of alternative Ni^{2+} cations in an octahedral environment with an anticipated $S = 1$ spin state, and $[\text{Ni}(\text{S}_2\text{O}_4)_2]^{-1}$ radical anion complexes with $S = \frac{1}{2}$, also taking into consideration that the Ni^{2+} occupancy is only 50%. Temperature dependence of the magnetic susceptibility for a trimolecular $(\text{Ni}^{2+})[\text{Ni}(\text{S}_2\text{O}_4)_2]$ formulation, reported here as the $\chi \times T$ product, is given in Fig. 8. We note that the room

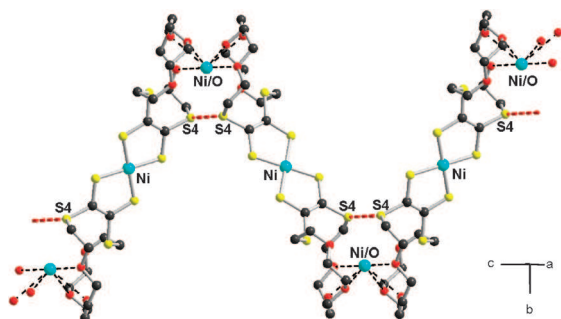


Fig. 7 Solid state organization within $(\text{Ni}, \text{H}_2\text{O})_{0.5}[\text{Ni}(\text{S}_2\text{O}_4)_2]$. Hydrogen atoms were omitted for clarity.

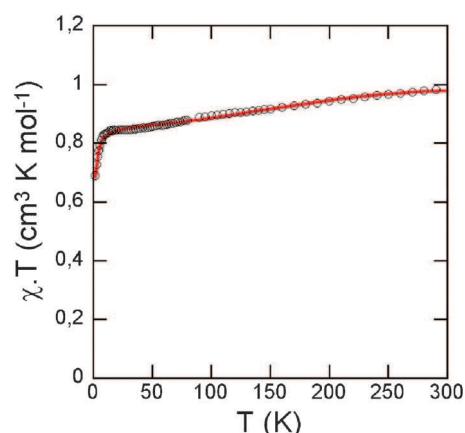


Fig. 8 Temperature dependence of the $\chi \times T$ product for $(\text{Ni}^{2+}, \text{H}_2\text{O})[\text{Ni}(\text{S}_2\text{O}_4)_2]^{-1}$. The solid line is the fit for the sum of the $S = 1$ Ni^{2+} species and uniform spin chain of antiferromagnetically coupled dithiolene complexes (see text).

temperature value of $\chi \times T$ does not exceed $1.0 \text{ cm}^3 \text{ K mol}^{-1}$, while a total value of $1.75 = 1$ (for the Ni^{2+}) + 2×0.375 (for the dithiolene complexes) would have been expected, considering a $g = 2$ value for both magnetic species, in the absence of any antiferromagnetic interactions.

Closer inspection of the crystal structure gives a rationale for this unexpected behaviour. As shown in Fig. 7, we note the presence of an extremely short intermolecular $\text{S}\cdots\text{S}$ contact between the radical dithiolene complexes with $\text{S}4\cdots\text{S}4^i$ at $3.332(2)$ Å ($i: 2 - x, y, 1.5 - z$). This intermolecular distance should be compared to the sulfur \cdots sulfur van der Waals contact distance at 3.70 Å. Furthermore, while the spin density in most radical nickel bis(dithiolene) complexes is essentially delocalized on the $\text{Ni}(\text{S}_2\text{C}_2)_2$ core, complexes with thioether ($-\text{SR}$) substituents are well known to allow a sizeable fraction of the spin density to delocalize on the outer sulfur atoms such as atom S4 in $[\text{Ni}(\text{S}_2\text{O}_4)_2]^{-1}$.^{2a} The overlap associated with the $\text{S}4\cdots\text{S}4^i$ intermolecular interaction is therefore at the origin of a strong antiferromagnetic interaction between the radical dithiolene complexes, which would then form a uniform chain of spins running along the c axis, independently from the Ni^{2+} cations. Accordingly, a tentative fit of the $\chi \times T$ product was performed (see Fig. 8), with the susceptibility χ as a sum of the Curie–Weiss contribution of the $S = 1$ Ni^{2+} species, $\chi_{\text{CW}} = 1/(T - \theta)$, with the contribution of a uniform spin chain for the dithiolene complexes noted χ_{BF} , associated with the following Hamiltonian $H = -J \sum_{i=1}^{n-1} S_i S_{i+1}$.^{32,33} It gives $\chi = \chi_0 + x(\chi_{\text{CW}} + 2\chi_{\text{BF}})$ with $\chi_0 = -0.8 \times 10^{-4} \text{ cm}^3 \text{ K mol}^{-1}$, $x = 0.86$, $\theta = -0.51(1)$ K and $J/k = -340(10)$ K ($J = -236 \text{ cm}^{-1}$).

In conclusion, we have shown here that dithiolene complexes substituted with crown ether moieties, investigated up to now only with alkaline cations in solution, can be used as metallo-ligands toward 3d magnetic metal cations for the elaboration of more complex magnetic systems. The obtained salt of $[\text{Ni}(\text{S}_2\text{O}_4)_2]^{-1}$ and Ni^{2+} illustrates however that this attractive approach is always in competition with direct antiferromagnetic interactions between the dithiolene complexes. This work is currently being extended to trivalent metal cations such as rare-earth cations as well as to smaller crown-ether moieties.

Experimental

All manipulations were performed under argon by use of Schlenk techniques. Solvents (MeOH, DMF and Et₂O) were distilled prior to use and degassed with argon.

Synthesis of the dithiocarbonate (S₂O₄)C=S

A 100 mL round bottom flask containing acetic acid (20 mL) and chloroform (7 mL) was charged with (S₂O₄)C=S^{19b} (0.470 g, 1.17 mmol) and Hg(OAc)₂ (0.820 g, 2.57 mmol). The reaction mixture was refluxed for 2 h. After cooling to room temperature, the reaction mixture was filtered through Celite® and the filtrate evaporated on the rotary evaporator. The residue was taken up in chloroform, filtered and the filtrate was concentrated, leaving an orange oil, purified by column chromatography (Silica, eluent: CH₂Cl₂/acetone 5 : 1 v/v, R_f = 0.88). After concentrating the second yellow band eluted from the column, the product crystallized in the form of yellow needles (0.276 g, 61%). Mp 71–72 °C. ¹H NMR (300 MHz, CDCl₃, TMS) δ 3.00 (t, J = 6 Hz, 4H), 3.61 (s, 12H), 3.71 (t, J = 6 Hz, 4H). ¹³C NMR (300 MHz, CDCl₃, TMS) δ 36.20, 69.59, 70.59, 70.78, 70.81, 127.43, 189.85. Elem. Anal. Calcd for C₁₃H₂₀S₄O₅. C, 40.60; H, 5.24%. Found: C, 40.39; H, 5.22%.

Synthesis of Na[Ni(S₂O₄)₂]

A Schlenk tube was charged with MeOH (5 mL) and sodium (0.010 g) for the *in situ* preparation on NaOMe. After the complete dissolution of Na, (S₂O₄)=O (0.085 g, 0.22 mmol) was added and the solution stirred for 1 h. Addition of NiCl₂·6H₂O (0.026 g, 0.11 mmol) to the yellow solution yields a brown precipitate. The reaction was left for stirring for another

hour. It was then filtered and the brown solid was dissolved in DMF, filtered and set to crystallize by Et₂O vapor diffusion. X-Ray diffraction quality single crystals were obtained after one week (0.07 g, 70%). Elem. Anal. Calcd. for C₂₄H₄₀NaNiO₈S₈: C, 36.27; H, 5.07%. Found: C, 36.30; H, 5.07%. UV-vis-NIR (DMF): λ_{max} = 1010 nm (ε = 8 437 L mol⁻¹ cm⁻¹).

Synthesis of (Ni,H₂O)[Ni(S₂O₄)₂]₂

A Schlenk tube was charged with MeOH (15 mL), (S₂O₄)=O (0.1 g, 0.25 mmol) and Bu₄NOH 1 M in MeOH (0.55 mL, 0.55 mmol). After the complete dissolution of (S₂O₄)=O the solution was stirred for 1 h. Addition of NiCl₂·6H₂O (0.060 g, 0.26 mmol) to the yellow solution yields a brown precipitate. The reaction was left for stirring for another hour. It was then filtered and the brown solid was dissolved in DMF, filtered and set to crystallize by Et₂O vapor diffusion. X-Ray diffraction quality single crystals were obtained after one week (0.056 g, 56%). Elem. Anal. Calcd for C₄₈H₈₂Ni₃O₁₇S₁₆: C, 35.58; H, 5.10%. Found: C, 35.29; H, 4.98%. UV-vis-NIR (DMF): λ_{max} = 1011 nm (ε = 9900 L mol⁻¹ cm⁻¹).

Synthesis of [Ni(S₂O₄)₂]₂

A Schlenk tube was charged with DMSO (3 mL) and (Ni,H₂O)[Ni(S₂O₄)₂]₂ (40.0 mg, 0.05 mmol). I₂ (20.2 mg, 0.08 mmol) in DMSO (1.0 mL) is added to the solution. A precipitate appears and the suspension was stirred for 15 minutes. EtOH was then added to complete the precipitation and the solid, once filtered and dried by suction, was redissolved in CH₂Cl₂, which, upon diffusion of Et₂O vapours, afforded needle shaped crystals (27 mg 70%). Elem. Anal. Calcd. for C₂₄H₄₀NiO₈S₈:

Table 2 Crystallographic data

Compound	(S ₂ O ₄)C=O	Na[Ni(S ₂ O ₄) ₂]	[Ni(S ₂ O ₄) ₂] ⁰	(Ni,H ₂ O)[Ni(S ₂ O ₄) ₂] ₂
Formula	C ₁₃ H ₂₀ O ₅ S ₄	C ₂₄ H ₄₀ NaNiO ₈ S ₈	C ₂₄ H ₄₀ NiO ₈ S ₈	C ₂₄ H ₄₀ Ni _{1.5} O _{8.5} S ₈
FW/g mol ⁻¹	384.53	794.74	771.75	809.14
Crystal color	Colorless	Black	Black	Black
Crystal size/mm	0.25 × 0.11 × 0.10	0.13 × 0.07 × 0.06	0.21 × 0.12 × 0.06	0.30 × 0.10 × 0.08
Crystal system	Monoclinic	Monoclinic	Triclinic	Monoclinic
Space group	P2 ₁ /c	C2/c	P $\bar{1}$	C2/c
T/K	293(2)	150(2)	150(2)	293(2)
a/Å	8.2045(16)	23.6588(14)	9.1918(4)	13.4074(4)
b/Å	11.888(2)	10.1920(6)	9.5486(4)	14.0031(4)
c/Å	17.949(4)	17.5265(12)	10.6697(4)	19.1875(6)
α/°	90.0	90.0	67.4590(10)	90.0
β/°	99.79(3)	127.742(2)	74.151(2)	104.5930(10)
γ/°	90.0	90.0	84.463(2)	90.0
V/Å ³	1725.1(6)	3342.0(4)	832.01(6)	3486.15(18)
Z	4	4	1	4
D _{calc} /g cm ⁻³	1.481	1.580	1.540	1.542
μ/mm ⁻¹	0.569	1.137	1.128	1.341
Total reffs.	16 204	12 062	10 148	15 378
Abs. corr.	Multi-scan	Multi-scan	Multi-scan	Multi-scan
T _{min} , T _{max}	0.928, 0.945	0.909, 0.934	0.850, 0.935	0.878, 0.917
Unique reffs.	3937	3800	3777	3991
R _{int}	0.0541	0.0563	0.0317	0.0268
Unique reffs (I > 2σ(I))	2726	2562	3498	3469
Refined param.	199	192	187	192
R ₁ (I > 2σ(I))	0.0357	0.0417	0.0233	0.0528
wR ₂ (all data)	0.0916	0.1568	0.0809	0.1811
Goodness-of-fit	1.088	1.044	1.206	1.061
Res. dens./e Å ⁻³	0.299, -0.284	0.523, -0.617	0.365, -0.384	1.352, -0.849

$$R_1 = \frac{\sum |F_o| - |F_c|}{\sum |F_o|}; wR_2 = \frac{[\sum w(F_o^2 - F_c^2)^2 / \sum w F_o^4]^{1/2}}$$

C, 37.35; H, 5.22%. Found C, 37.46; H, 5.76%. UV/Vis/NIR (DMF) $\lambda = 1015$ nm ($\epsilon = 10970$ L mol⁻¹ cm⁻¹).

X-Ray diffraction studies

Single crystals were mounted on the top of a thin glass fiber. Data were collected on a Nonius KappaCCD diffractometer at room temperature with graphite-monochromated Mo-K α radiation ($\lambda = 0.71073$ Å). Structures were solved by direct methods (SHELXS-97) and refined (SHELXL-97) by full-matrix least-squares methods,³⁴ as implemented in the WinGX software package.³⁵ Absorption corrections were applied. Hydrogen atoms were introduced at calculated positions (riding model), included in structure factor calculations, and not refined. Crystallographic data are summarized in Table 2.

Magnetic properties

The magnetic susceptibility measurements were obtained from a Quantum Design SQUID magnetometer MPMS-XL. This magnetometer works between 1.8 and 400 K for dc applied fields ranging from -5 to 5 T. Measurements were performed on polycrystalline samples of Na[Ni(S₂O₄)₂] (5.7 mg) and (Ni₂H₂O)[Ni(S₂O₄)₂] (14 mg). The magnetic data were corrected for the sample holder and the diamagnetic contributions. EPR spectra were obtained either on a X-band Bruker EMX-8/2.7 spectrometer equipped with a liquid nitrogen cooling system. Simulations were performed with Bruker WinEPR Symphonia.

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