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Molecular and electronic structure of the dithiooxalato radical ligand stabilised by rare earth coordination†

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Heterometallic rare earth transition metal compounds of dithiooxalate (dto)²⁻, [Ni^{II}{(dto)Ln^{III}Tp₂}₂] (Ln = Y (1), Gd (2); Tp = hydrotris(pyrazol-1-yl)borate) were synthesised. The Lewis acidic rare earth ions are bound to the dioxolene and chemical reduction of 1 and 2 with cobaltocene yielded [CoCp₂]⁺[Ni^{II}{(dto)Ln^{III}Tp₂}₂]¹⁻ Ln = Y (3), Gd (4). The reduction is ligand-based and 3 and 4 are the first examples of both molecular and electronic structural characterisation of the dithiooxalato radical (dto)^{3•-}.

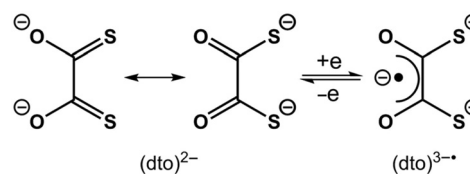
A century of coordination chemistry of the dithiooxalate ligand, (dto)²⁻, has revealed a unique set of properties that sets it apart from other 1,2-dithiolate ligands, *i.e.* dithiolenes.^{1,2} The utility of this ditopic ligand stems from the asymmetry of the two binding sites: mid-to-late transition metal ions bind preferentially to the soft sulfur-donor dithiolate, whereas Lewis acidic metal ions bind preferentially to the hard oxygen-donor α -diketonate.² This has led to the assembly of numerous mixed-metal oligomers and coordination polymers by simply combining hard and soft metal ions with a dithiooxalate salt in a one-pot reaction.³⁻⁶ The majority of these studies have focussed on the structural topology of the metal ions, and the magnetic properties of the constituent metal ions linked by this ligand.

The electron transfer chemistry, inherent to bis(dithiolene) compounds, does not exist in monometallic bis(dithiooxalate) species such as [Ni(dto)₂]²⁻. This stems from the resonance stabilisation within the (dto)²⁻ ligand, between the dithiolate and diolate forms (Scheme 1). Prior to this work the only example of reversible one electron transfer chemistry using (dto)²⁻ was [M{(dto)SnX₄}₂]²⁻ (M = Ni, Pd, Pt; X = F, Cl).^{5,6} Akin to bis(dithiolene) compounds,⁷ the redox chemistry was shown to be ligand-based, with consecutive electrons added to a π^*

orbital stabilised by coordination of the Lewis acidic Sn, generating the α -diketonate radical chelate (dto)^{3•-}.⁸ However, attempts to isolate and characterise the one electron reduced compounds, were unsuccessful.

Herein we report the synthesis and characterisation of two heterometallic compounds [Ni^{II}{(dto)Y^{III}Tp₂}₂] (1) and [Ni^{II}{(dto)Gd^{III}Tp₂}₂] (2) (Tp = hydrotris(pyrazol-1-yl)borate), in which the α -diketonate site is occupied by Lewis acidic rare earth ions. Compounds 1 and 2 both display consecutive one-electron reduction waves in their cyclic voltammograms, at reduction potentials that are significantly more negative than [M{(dto)SnX₄}₂]²⁻. The chemical reduction of 1 and 2 with cobaltocene yielded [CoCp₂]⁺[Ni^{II}{(dto)Y^{III}Tp₂}₂]¹⁻ (3) and [CoCp₂]⁺[Ni^{II}{(dto)Gd^{III}Tp₂}₂]¹⁻ (4). The locus of the reduction is confirmed as ligand-based, thus 3 and 4 are the first structurally characterised compounds of the elusive dithiooxalato radical ligand (dto)^{3•-}.

Compounds 1 and 2 were synthesised by simultaneous addition of two equivalents of aqueous rare earth chloride and four equivalents of KTp, to a stirred aqueous solution of K₂[Ni(dto)₂] at room temperature (Scheme 2). Compounds 1 and 2 precipitated immediately from aqueous solution and were isolated as violet powders in good yields (65–70%). We attribute the increase in yield when compared to related 3d–4f species to the modular approach of using K₂[Ni(dto)₂] as a synthon.⁴ The simultaneous addition of metal and capping ligand to the reaction mixture limits the formation of [LnTp₂X] by-products and circumvents the need for several purification



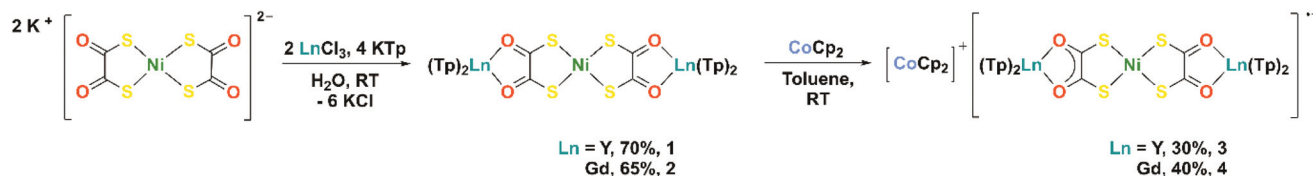
Scheme 1 Resonance stabilisation and one electron redox chemistry of the (dto)²⁻ ligand.

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† Electronic supplementary information (ESI) available: Experimental details, IR spectra, electronic spectra, cyclic voltammogram of 2, crystal structure data, computational data. CCDC 1886429–1886432 and 1889410. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c9dt00688e





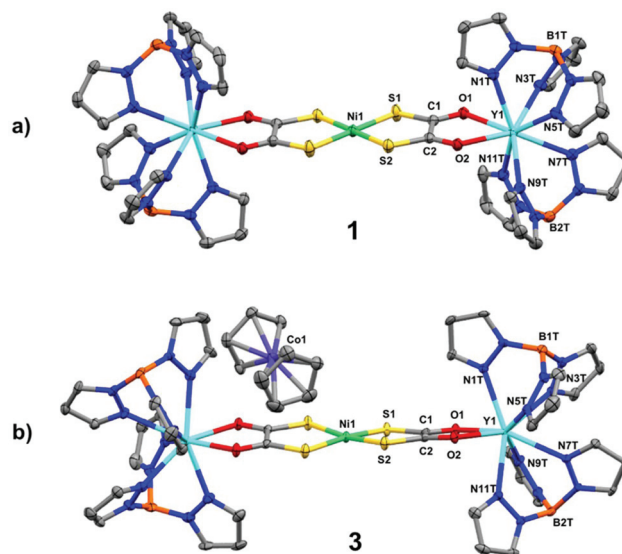
Scheme 2 Synthesis of compounds 1–4.

steps.^{9,10} The ¹H NMR spectrum of **1** in C₆D₆ at room temperature displays three resonances at δ = 7.34, 7.09, and 5.68 ppm, in the expected 1 : 1 : 1 ratio for magnetically equivalent pyrazolyl rings coordinated to a rare earth ion.^{9,11} The BH is observed as a very broad resonance at δ = 4.7 ppm. We also synthesised [PPh₄]₂[Ni{(dto)SnCl₄}]₂ **5** to benchmark our experimental data on **1–4**. The ATR IR spectra of **1**, **2** and **5** are consistent with dioxolene binding. The C–O stretching vibrations in **1**, **2** (1523–1503 cm⁻¹) and **5** (1493–1479 cm⁻¹) are observed at lower frequency than ~1600 cm⁻¹ observed in [PPh₄]₂[Ni(dto)]₂.¹²

Based on the first reduction potential of **1** and **2**, determined by electrochemistry (see below), chemical reduction of **1** and **2** with one equivalent of cobaltocene in toluene at room temperature yielded **3** and **4**, respectively (Scheme 2). Compounds **3** and **4** were isolated as teal coloured powders in moderate yields (30–40%). We attribute the successful isolation of the (dto)^{3•-} radical anion in **3** and **4** to the combined effect of the Lewis acidic rare earth metal and stabilising Tp co-ligands. The purity of **1–5** were confirmed by elemental analysis and all complexes were fully characterised (see ESI† for full experimental details and spectra).

The solid-state molecular structures of compounds **1** and **2** were determined by single crystal X-ray diffraction and are shown in Fig. 1a and Fig. S1.† The square planar NiS₄ unit has Ni–S bond distances and S–Ni–S angles (Table 1) consistent with those reported for monometallic [Ni(dto)]₂²⁻,¹⁴ and **5** (Fig. S3†).⁶ In **1** and **2** the bridging [Ni(dto)]₂²⁻ unit is capped by two eight-coordinate Y or Gd ions whose distorted square antiprismatic geometry is completed by two facially binding Tp ligands. Both complexes are present as *meso* compounds, where one rare earth centre has Λ and the other has Δ absolute configuration because of the crystallographic inversion symmetry at Ni.¹⁵ The Ln–O and Ln–N distances fall within the normally observed range. The intraligand S–C, C–C and C–O bond distances (Table 1) do not vary significantly between **1**, **2** and **5**, but do demonstrate the resonance stabilisation (Scheme 1) that gives (dto)²⁻ its distinctive chemistry in comparison to other 1,2-dithiolenes.⁷

Compounds **3** and **4** are the first examples of structurally characterised dithiooxalato (dto)^{3•-} ligands in a coordination compound (Fig. 1b and Fig. S2†). Their overall geometry differs from **1** and **2** only by a modest tetrahedralisation of the NiS₄ core of 10.3° in **3** and 9.9° in **4** (Table 1) and the absence of a crystallographic inversion centre at Ni. The increased charge on the (dto)^{3•-} ligands significantly shortens the Y–O and Gd–O bonds (Table 1). Reduction of (dto)²⁻ to (dto)^{3•-}

Fig. 1 Crystal structures of (a) **1** and (b) **3**. Thermal ellipsoids drawn at 50% probability and hydrogen atoms omitted for clarity.Table 1 Selected bond distances and angles^a

	1	2	3	4	5
M'	Y	Gd	Y	Gd	Sn
Avg. Ni–S	2.1851(9)	2.187(1)	2.168(1)	2.164(1)	2.1803(6)
Avg. M'–O	2.390(2)	2.435(2)	2.326(3)	2.370(3)	2.202(2)
Avg. S–C	1.696(3)	1.699(4)	1.711(4)	1.714(5)	1.681(3)
C1–C2	1.525(5)	1.537(5)	1.468(5) ^c	1.466(5) ^c	1.526(3)
Avg. C–O	1.244(4)	1.241(4)	1.275(5)	1.271(5)	1.248(3)
Avg. M'–N	2.470(3)	2.508(3)	2.497(3)	2.520(4)	
Ni...M'	6.105	6.156	6.071 ^c	6.099 ^c	5.825
M'...M'	12.210	12.313	12.114	12.177	11.650
S–Ni–S	92.78(3)	92.72(4)	92.22(4) ^c	92.21(5) ^c	93.07(2)
O–M'–O	66.23(7)	65.30(8)	68.84(7) ^c	68.0(1) ^c	74.41(6)
α ^b	0	0	10.3	9.9	0

^a Distances in angstrom; angles in degrees. ^b Dihedral angle between mean NiS₂ planes. ^c Average value.

leads to a significant shortening of the C–C bonds and concomitant lengthening of the C–O bonds from those found in **1** and **2** (Table 1). The C–O bond distances in **3** and **4** are consistent with those found in semiquinone ligands^{17,18} (quinoidal distortion).⁷ This is consistent with the dithiooxalato radical (dto)^{3•-}, with the unpaired electron delocalised over both dto ligands (Fig. 3).



the complex distributes +1.04 spins at each end of the bridging $\{\text{Ni}(\text{dto})_2\}$ unit with -0.03 spins on the Ni ion from a minor polarisation of the S–C bonds. Notably there is no spin density on the Y(III) ions in accordance with the absence of hyperfine structure in the EPR spectrum. The identical spin distribution is found in the computational one-electron reduced 5, $[\text{Ni}\{\text{dto}\text{SnCl}_4\}_2]^{3-}$ (Fig. S14†). As the analogous chemical reduction of 5 was unsuccessful, the electronic structure was calculated on an optimised geometry which exhibited the same intraligand bond distances and angles as seen in 3 and 4 (Fig. S13†).

The electronic structures of 2 and its one-electron reduced product 4 have been calculated using the broken symmetry (BS) method in order to account for the spin coupling between the terminal Gd(III) $S = 7/2$ ions and $(\text{dto})^{3*-}$ radical. It has been shown that DFT can reliably estimate the spin coupling between lanthanide ions and organic radicals.¹⁹ A BS(7,7) calculation for 2 gave isoenergetic $M_S = 0$ state for antiferromagnetically coupled Gd(III) ions, and $M_S = 7$ for ferromagnetically coupled Gd(III) ions. This implies that two paramagnetic centres are uncoupled, as expected for the 12.31 Å separation with negligible isotropic exchange coupling ($J \approx 0$). Reduction of 2 to 4 introduces a third spin centre into the system, and so BS(14,1), BS(8,7) and BS(7,6) calculations that account for all spin coupled permutations were performed on the crystallographic coordinates of 4 (Table S3†). The results revealed the BS(14,1) solution as the most stable, which is the parallel alignment of the 7 spins on each Gd(III) ions with the opposed alignment of the single spin on the $(\text{dto})^{3*-}$ ligand, *i.e.* $M_S = 13/2$ (Fig. 3b). This solution is marginally more stable than the uncoupled scenario with the Gd(III)-radical exchange interaction estimated at $J = -1.3 \text{ cm}^{-1}$. Although small in magnitude, it does fall in the range observed for many other Gd(III)-radical systems.¹⁹ Compound 4 is most closely related to $[\text{Tp}_2\text{Gd}^{\text{III}}(\text{dtbsq})]$ (dtbsq = 3,5-di-*tert*-butylsemiquinonato), with $J = -5.7 \text{ cm}^{-1}$.¹⁷ The smaller exchange interaction estimated for 4 lies in the fact that the $(\text{dto})^{3*-}$ unpaired electron is delocalised over two dioxolene units in the complex. This reduces the relative spin concentration at each of the four oxygen atoms in $\{\text{Ni}(\text{dto})_2\}^{*-}$ compared to the two oxygen atoms of *o*-benzosemiquinones.²⁰ In spite of this, reduction of the bridging $\{\text{Ni}(\text{dto})_2\}$ unit in 4 enforces a ferromagnetic alignment of the terminal Gd(III) ions, which maximises the total spin ground state of this system, albeit at very low temperatures. This spin distribution in the bridging metallodithiooxalate unit can be modified by changing the metal at its core, which will not only modulate the covalency but also the geometry and therein the alignment of the magnetic anisotropy of the terminal ions, which we will continue to investigate using this system.

In conclusion, heterometallic rare earth transition metal complexes of the $(\text{dto})^{2-}$ ligand, display ligand-based quasi-reversible electron transfer chemistry. Chemical reduction of these complexes led to the first compounds in which both molecular and electronic structure characterisation of the dithiooxalato radical $(\text{dto})^{3*-}$ was possible.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

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References

- H. O. Jones and H. S. Tasker, *J. Chem. Soc., Trans.*, 1909, **95**, 1904.
- W. Dietzsch, P. Strauch and E. Hoyer, *Coord. Chem. Rev.*, 1992, **121**, 43.
- (a) P. D. W. Boyd, J. Hope, C. L. Raston and A. H. White, *Aust. J. Chem.*, 1990, **43**, 601; (b) J. M. Bradley, S. G. Carling, D. Visser, P. Day, D. Hautot and G. J. Long, *Inorg. Chem.*, 2003, **42**, 986; (c) D. Coucouvanis and D. Piltingsrud, *J. Am. Chem. Soc.*, 1973, **95**, 5556; (d) M. Leitheiser and D. Coucouvanis, *Inorg. Chem.*, 1977, **16**, 1611; (e) F. J. Hollander and D. Coucouvanis, *Inorg. Chem.*, 1973, **13**, 2381; (f) C. Frasse, J. C. Trombe, A. Gleizes and J. Galy, *C. R. Acad. Sci., Ser. II: Mec., Phys., Chim., Sci. Terre Univers*, 1985, **200**, 403; (g) A. Gleizes, F. Maury, P. Cassoux and J. Galy, *Z. Kristallogr.*, 1981, **155**, 293; (h) J. C. Trombe, A. Gleizes and J. Galy, *C. R. Acad. Sci., Ser. II: Mec., Phys., Chim., Sci. Terre Univers*, 1982, **294**, 1369; (i) A. Gleizes and M. Verdagner, *J. Am. Chem. Soc.*, 1984, **106**, 3727.
- (a) M. Siebold, S. Eidner, A. Kelling, M. U. Kumke, U. Schilde and P. Strauch, *Z. Anorg. Allg. Chem.*, 2006, **632**, 1963; (b) M. Siebold, A. Kelling, U. Schilde and P. Strauch, *Z. Naturforsch., B: J. Chem. Sci.*, 2005, **60**, 1149; (c) M. Siebold, M. Korabik, U. Schilde, J. Mrozinski and P. Strauch, *Chem. Pap.*, 2008, **62**, 487; (d) G.-F. Xu, P. Gamez, J. Tang, R. Clérac, Y.-N. Guo and Y. Guo, *Inorg. Chem.*, 2012, **51**, 5693.
- D. Coucouvanis, *J. Am. Chem. Soc.*, 1970, **92**, 707.
- D. Coucouvanis, N. C. Baenziger and S. M. Johnson, *J. Am. Chem. Soc.*, 1973, **95**, 3875.
- (a) K. Ray, T. Petrenko, K. Wieghardt and F. Neese, *Dalton Trans.*, 2007, 1552; (b) S. Sproules and K. Wieghardt, *Coord. Chem. Rev.*, 2011, **255**, 837.
- G. A. Bowmaker, P. D. W. Boyd and G. K. Campbell, *Inorg. Chem.*, 1982, **21**, 3565.
- T. Sanada, T. Suzuki and S. Kaizaki, *J. Chem. Soc., Dalton Trans.*, 1998, 959.
- T. Sanada, T. Suzuki, T. Yoshida and S. Kaizaki, *Inorg. Chem.*, 1998, **37**, 4712.
- D. P. Long, A. Chandrasekaran, R. O. Day and P. A. Bianconi, *Inorg. Chem.*, 2000, **39**, 4476.



- 12 F. Junnosuke and N. Kazuo, *Bull. Chem. Soc. Jpn.*, 1964, **37**, 528.
- 13 A. R. Latham, V. C. Hascall and H. B. Gray, *Inorg. Chem.*, 1965, **4**, 788.
- 14 P. Román, A. Luque, J. M. Gutiérrez-Zorilla and J. I. Beitia, *Z. Kristallogr.*, 1992, **198**, 213.
- 15 N. G. Connelly, in *Nomenclature of inorganic chemistry. IUPAC recommendations 2005*, Royal Society of Chemistry, 2005.
- 16 (a) D. R. Eaton, *Inorg. Chem.*, 1964, **3**, 1268; (b) B. S. Prabhananda, C. C. Felix, B. Kalyanaraman and R. C. Sealy, *J. Magn. Reson.*, 1988, **76**, 264.
- 17 A. Caneschi, A. Dei, D. Gatteschi, L. Sorace and K. Vostrikova, *Angew. Chem., Int. Ed.*, 2000, **39**, 246.
- 18 (a) A. Caneschi, A. Dei, D. Gatteschi, S. Poussereau and L. Sorace, *Dalton Trans.*, 2004, 1048; (b) J. R. Hickson, S. J. Horsewill, J. McGuire, C. Wilson, S. Sproules and J. H. Farnaby, *Chem. Commun.*, 2018, **54**, 11284; (c) G. H. Spikes, S. Sproules, E. Bill, T. Weyhermüller and K. Wieghardt, *Inorg. Chem.*, 2008, **48**, 10935.
- 19 (a) T. Gupta, T. Rajeshkumar and G. Rajaraman, *Phys. Chem. Chem. Phys.*, 2014, **16**, 14568; (b) T. Rajeshkumar and G. Rajaraman, *Chem. Commun.*, 2012, **48**, 7856.
- 20 R. R. Kapre, E. Bothe, T. Weyhermüller, S. DeBeer George, N. Muresan and K. Wieghardt, *Inorg. Chem.*, 2007, **46**, 7827.

