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Caught! Crystal trapping of a side-on peroxo bound to Cr(IV)[†]

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A Cr(IV)η²-peroxo complex crystallizes from 33% aqueous H₂O₂. The complex is a likely intermediate in catalytic disproportionation of H₂O₂ proposed to occur through a single metal site mechanism in solution – and solid state.

The study of metal-dioxygen, peroxo and oxo adducts has attracted considerable attention as such metal-activated oxygen species have emerged as the common precursors and active oxidants in a wide range of biological and non-biological oxidation systems.^{1–5} The majority of first row transition metals are represented as co-factors in enzymes and proteins. However, in spite of its rich redox chemistry, reasonable availability and oxophilicity, chromium is by large absent in the natural world. Instead, it is generally considered a cell toxin capable of causing damage to DNA and RNA, through both oxidative mechanisms and adduct formation.^{6,7} Several molecular Cr(v)-oxo complexes have been reported in the literature over the last 40 years,^{8–13} whereas only a handful Cr-dioxygen complexes have been identified. In all cases these compounds are derived from the reaction of Cr(II) precursors with O₂. An ion proposed to be a Cr(III)-superoxo complex [CrO₂(bipy)₂]²⁺ (bipy = 2,2'-bipyridine) was obtained by reacting coordinatively unsaturated [Cr(bipy)₂]²⁺ with O₂ in the gas phase.^{14,15} Subsequently Theopold *et al.*¹⁶ structurally characterized a “side-on” Cr(III)-superoxo adduct generated from the reaction of [Cr^{II}(Tp^{tBu,Me})NC₆H₅]⁺ (Tp^{tBu,Me} = hydrotris(3-*tert*-butyl-5-methylpyrazolyl)borate) with O₂ in diethylether. Recently Nam *et al.* have reported both an “end-on” Cr(III)-superoxo¹¹

and a “side-on” Cr(IV)-peroxo¹⁷ adduct supported by macrocyclic tetramethylcyclams of different ring sizes.

Molecular Cr-dioxygen species are relevant with respect to the biotoxicity of chromium, and are useful as structural mimics of more elusive metal-dioxygen adducts of the later transition metals. We have previously used amino-acid derived pyridine ligands, such as tpena[−] (*N,N,N'*-tris(2-pyridylmethyl)ethylenediamine-*N'*-acetate), to generate reactive iron(IV)-oxo,^{18,19} manganese(IV)-oxo complexes and putative O₂ adducts,^{20,21} and a metal-oxidant (iodosylbenzene) adduct.²² The instability of such high-valent iron and manganese complexes makes their structural characterization especially difficult. The isolation of more stable structural analogues based on earlier first row transition metals is a potentially fruitful strategy for gleaning structural information on such intermediates.¹⁸ Here we describe the structural trapping of a catalytically competent Cr(IV)-peroxo adduct formed from the reaction of a Cr(III) precursor with H₂O₂.

[Cr^{III}(tpena)]²⁺ (**1**) is prepared in aqueous solution from the reaction of chromium(III)-nitrate with the sodium salt of the ligand, followed by crystallization as a diperchlorate salt. The X-ray crystal structure shows that tpena[−] coordinates through all six donor atoms. The Cr(III) atom displays a particularly irregular octahedral geometry, Fig. S1 (ESI[†]), most notable being an extremely obtuse N1–Cr–N5 angle of 115.40(4)°. When **1** is dissolved, UV-Vis spectra (pH 4–8) show that aqueous solutions contain an equilibrium between **1**, its “pseudo hydrate”, [Cr^{III}(tpenaH)OH]²⁺ (**2**) and the blue congener base [Cr^{III}(tpena)OH]⁺ (**3**), Scheme 1. Complex **3** is the base peak in ESI-MS spectra and this speciation is supported by cyclic voltammetry, *vide infra*. Addition of base drives the equilibrium towards **3**. When 100 eq. of H₂O₂ are added to solutions of **1**, a colour change from red to violet occurs over 30 min accompanied by the evolution of dioxygen²³ consistent with modest catalase activity. The solutions are EPR silent.

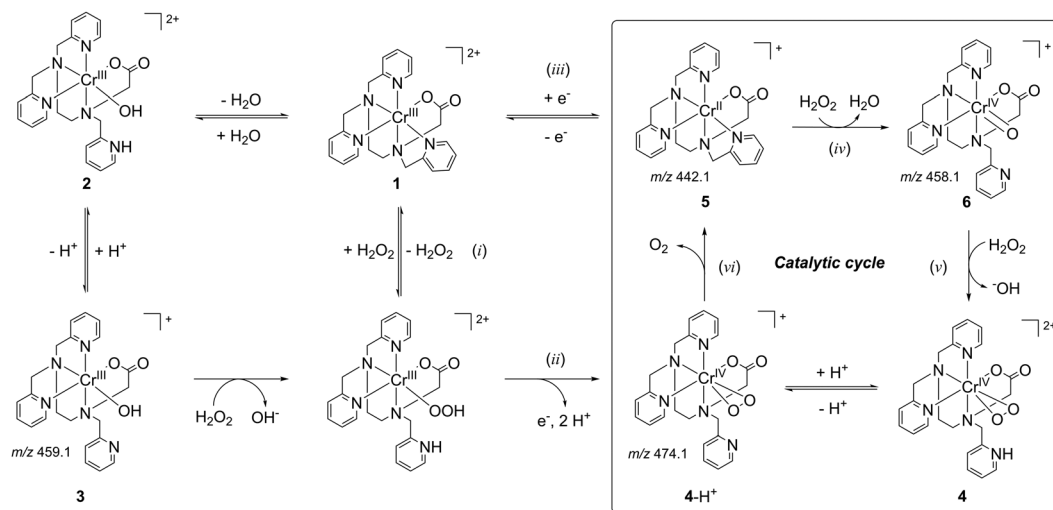
This observation, together with UV-Vis spectroscopy (λ, 539 nm; ε, 150 M^{−1} cm^{−1}), Fig. 1, and electrospray ionization (ESI) mass spectra – which show the dominant presence of a Cr(tpena)-dioxygen adduct [CrO₂(tpena)]⁺ at *m/z* 474.1 (calcd 474.1),

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[†] Electronic supplementary information (ESI) available: Experimental details of syntheses, crystal structure determination, spectroscopic techniques and supporting CV, EPR, ESI-MS, GC-TCD, rR and time-resolved UV-Vis spectra. CCDC 1031521 and 1031522. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c4cc08785b





Scheme 1 Interrelationship between *tpena*[−] complexes of Cr(II), Cr(III) and Cr(IV), including a mechanistic proposal for the catalytic decomposition of H₂O₂.

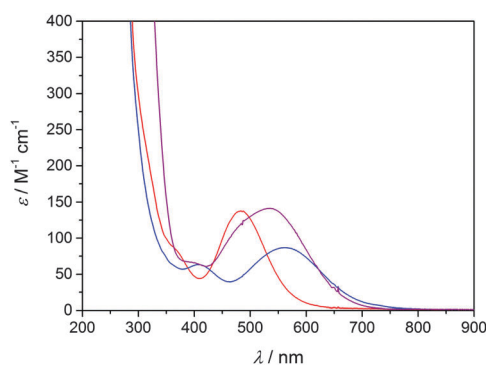


Fig. 1 Aqueous UV-Vis spectra of 4 mM **1** (red line), 3.7 mM **4** (violet line, generated *in situ* from **1** and 200 eq. H₂O₂) and 3.6 mM **3** (blue line, generated *in situ* from **1** and 50 eq. NaOH).

suggests either of the isomeric species: a triplet Cr(III)–superoxo system consisting of a Cr(III) center anti-ferromagnetically coupled to a O₂^{•−} radical, or a triplet Cr(IV)–peroxo system. A Cr(III)–hydroperoxo adduct was ruled out, since this formulation would give rise to a simple 3/2 system presumably with a spectrum similar to that obtained for **1**, Fig. S5 (ESI[†]). The presence of a band at 878 cm^{−1} in the resonance Raman spectrum of the purple solutions, Fig. 2(a) and Fig. S8(b) (ESI[†]), (full spectrum) identifies the species unambiguously as a peroxo, rather than a superoxo complex.²⁴ Consistently the IR spectrum of an EPR silent solid precipitated with diethyl ether–dioxane shows a strong sharp absorption at 871 cm^{−1} not seen in IR spectrum of **1**, Fig. 2(b). The differences in the ligand vibrations at 1256, 1373 and 1669 cm^{−1} suggest that a pyridine arm of the ligand has become uncoordinated and protonated as seen in the salts of [V^{IV}O(*tpenaH*)]²⁺ and [Fe^{III}₂O(*tpenaH*)₂]⁴⁺,^{18,22} and the formulation [Cr^{IV}O₂(*tpenaH*)]²⁺ (**4**) could be proposed.

Single crystals of 4(ClO₄)₂(H₂O)₂(H₂O) were obtained only on a couple of occasions by placing the largest crystals of 1(ClO₄)₂(C₄H₈O₂)_{0.5} available into aqueous 33% H₂O₂. Over the course of a day at 4 °C the starting material was replaced

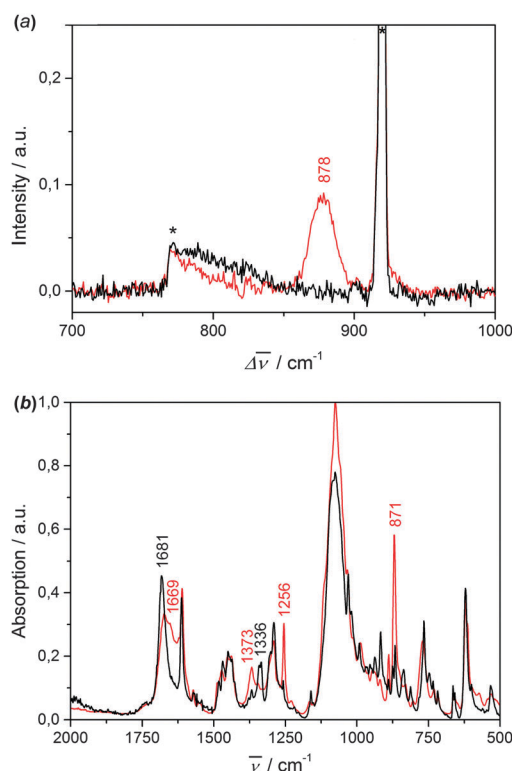


Fig. 2 (a) Frozen solution rRaman spectra of 9 mM **1** (black) and **4** (red) recorded at 77 K in CH₃CN (λ_{ex} = 514.5 nm, power = 65 mW). Asterisks denote solvent signals. (b) Solid state IR spectra of **1**(ClO₄)₂ (black) and **4**(ClO₄)₂ (red).

by plates of the purple peroxo complex. Bubbles of O₂ were observed to slowly emerge from both the solution and the surface of the crystals. The X-ray crystal structure, Fig. 3(a), shows that the peroxide is coordinated to a seven-coordinate Cr-centre in a side-on fashion, and charge balance dictates that the cation is a Cr(IV) species. As anticipated one methylpyridyl arm is uncoordinated and protonated. The distance between

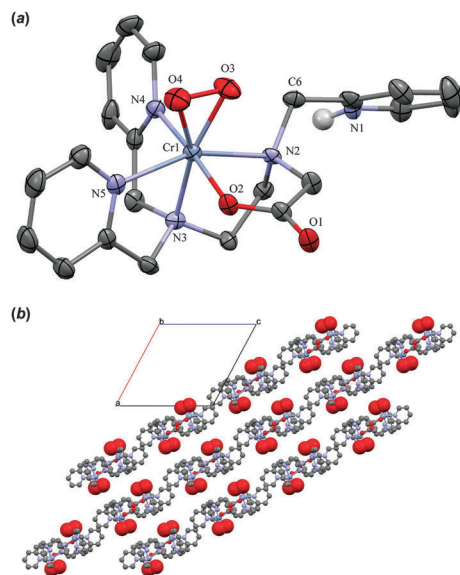


Fig. 3 (a) ORTEP plot of the cation in $4(\text{ClO}_4)_2(\text{H}_2\text{O}_2)_3(\text{H}_2\text{O})$ showing 50% probability ellipsoids. (b) Ball-and-stick illustration of the crystal lattice defined by the chains of H-bonded **4** with the peroxide ligand as space filled spheres after *in silico* removal of H_2O_2 , H_2O and ClO_4^- . Viewed down the *b*-axis.

the O atoms of the O_2 moiety is 1.383(8) Å. The average Cr–L distances for **1** and **4** respectively are 2.040(1) Å and 2.014(6) Å supporting a higher oxidation for the chromium ion in **4**. The +4 oxidation state is confirmed by a Bond Valence Sum (BVS) analysis that yields a BVS = 4.001.²⁵ A hydrogen-bonding motif is present in the crystal, with the cations linked together by the protonated dangling pyridyl arm of one molecule and the non-coordinated carbonyl oxygen of a neighbour ($\text{PyH}^+ \cdots \text{OOC}$, 1.904 Å), forming herringbone chains. The hydrogen-bonded chains stack regularly to form sheets parallel to the *b*-axis, forming the basis for a network of pores running along the *c*-axis. The pores are occupied by three co-crystallized H_2O_2 molecules per cation. The perchlorate ions H-bond predominantly to the substrate guest H_2O_2 . The unusually large number of H_2O_2 molecules in the crystal lattice, and their disorder, suggest that these might be rather mobile guests. Thus we conclude that a “crystal trapping” of the reactive **4** is made possible because it is surrounded by substrate that is taken up in the solid state, selectively, from solution. A calculation of the pore dimensions after *in silico* removal of the H_2O_2 molecules reveals about 16% pseudo void space. These voids lie predominantly between internal lamellar surfaces with the coordinated peroxo ligands lining the surface of these sheets, Fig. 3(b). With such a spatial arrangement, it seems reasonable to assume that guest H_2O_2 molecules can creep between the sheets, while product O_2 is able to leave the crystals the same way by diffusion.

Deprotonated **4** (m/z 474.1, 4-H^+) is present in ESI mass spectra obtained on both working solutions containing excess H_2O_2 , and in the spectra of isolated $4(\text{ClO}_4)_2(\text{H}_2\text{O}_2)_4$ redissolved in acetonitrile. These spectra furnish some hints about the mechanism of the catalytic H_2O_2 disproportionation, and the decomposition of **4** in the absence of substrate H_2O_2 , respectively. The dominant ion in both spectra is $[\text{Cr}^{\text{II}}(\text{tpena})]^+$ (**5**; m/z 442.1, calcd 442.1). While **5** is

accessible electrochemically (*vide infra*), it was present, only occasionally, as a minor ion in the spectra of **1** (complex **3** dominates the spectra of **1**, Fig. S2, ESI†). These observations suggest that **5** is not predominantly formed *via* an ionization induced one-electron reduction of **1**. Collision Induced Dissociation (CID) experiments on the 474.1 ion (4-H^+) result in the loss of the mass equivalent of two oxygen atoms to directly generate **5**. That no stepwise O atom loss occurs, strongly indicates that the O–O bond remains intact during this process¹⁴ and that this is the dominant route for generation of the observed Cr(II) species **5**. Ions assigned to $[\text{Cr}^{\text{IV}}\text{O}(\text{tpena})]^+$ (m/z 458.1, calcd 458.1) and $[\text{Cr}^{\text{VO}}(\text{tpena})]^{2+}$ (m/z 229.1, calcd 229.1), and its ion pair, $[\text{Cr}^{\text{VO}}(\text{tpena})]\text{ClO}_4^+$ (m/z 557.1 calcd 557.1) and other high-valent O containing species appear only in the spectra of isolated $4(\text{ClO}_4)_2$. These species might result from the inter- and intra-molecular O atom transfer from **4** to an unknown substrate or the ligand.²⁶ It is noteworthy that a greater number of ligand decomposition products are observed in Fig. 4(b). On the basis of the electronic and geometrical structure of **4**, the gas phase experiments through which various Cr(II), Cr(IV)–peroxo, Cr(IV)–oxo, and Cr(V)–oxo species are observed, we propose in Scheme 1, a catalytic acid–base type cycle for the relatively slow disproportionation of H_2O_2 . The cycle is similar to mechanisms proposed for heme catalases.²⁷ Indeed the possible involvement of a second coordination sphere basic pyridine to aid proton transfers is reminiscent of the role of the proximal histidine in peroxidases. The species

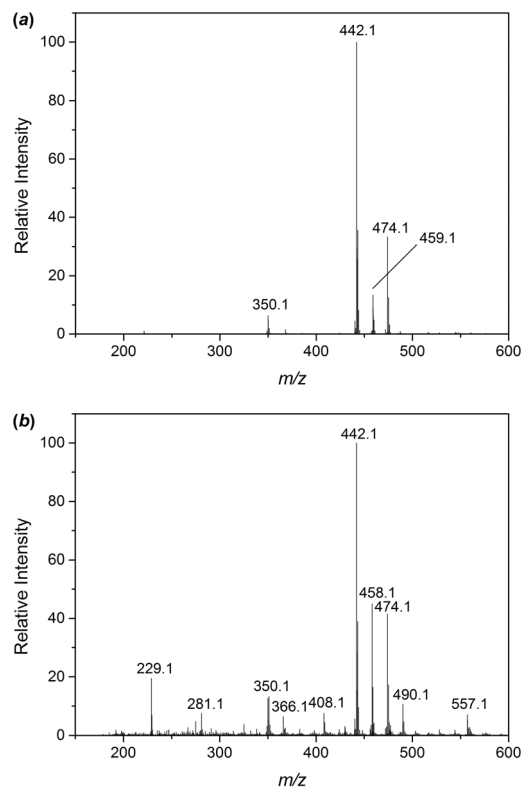


Fig. 4 (a) ESI mass spectrum of solution of **4** generated *in situ* from the reaction of **1** with 200 eq. H_2O_2 in H_2O . (b) ESI mass spectrum of $4(\text{ClO}_4)_2$ dissolved in CH_3CN . Important assignments not in the text: m/z 490.1 $[\text{Cr}^{\text{VI}}(\text{O})(\text{O}_2)(\text{tpena})]^+$ (or $[\text{Cr}^{\text{IV}}(\text{O}_2)(\text{tpenaO})]^+$), 366.1 $[\text{Cr}^{\text{III}}(\text{tpenaO}-\text{CHC}_5\text{H}_4\text{N})]^+$, 350.1 $[\text{Cr}^{\text{III}}(\text{tpena}-\text{CH}_2\text{C}_5\text{H}_4\text{N})]^+$.



labelled by a number have been detected, and a structural analogue for **6**, $[\text{V}^{\text{IV}}\text{O}(\text{tpenaH})]^+$ is known.¹⁸ The direct observation of the O_2 release process step (vi) by CID suggests that the pro-catalyst **1** is not directly involved in the catalytic cycle, but requires reductive activation, by direct (i) + (ii) or indirect (iii) involvement by H_2O_2 , before it can enter the cycle. Time-resolved UV-Vis spectroscopy indicates that the direct (i) + (ii) mechanism is the dominant activation pathway, since the time trace for the appearance of the λ_{max} at 553 nm due to the chromophore of **4** can be modelled satisfactorily with a 2-step mechanism (see ESI†, Fig. S7). The cyclic voltammogram of $\mathbf{1}(\text{ClO}_4)_2$ reveals the presence of **1** and **3** in neutral aqueous solutions, with reversible $\text{Cr}^{2+/3+}$ redox couples at $E_{1/2} = -885$ mV and -1099 mV (vs. Ag/Ag^+), Fig. S3 (ESI†), supporting the plausibility of the involvement of **5** in the catalytic cycle. The $\text{Cr}(\text{IV})$ -oxo species, **6**, formed by a O atom transfer from H_2O_2 to **5** (iv) is expected to be a strong base which can deprotonate H_2O_2 and assist the substitution step (v),²⁸ perhaps with involvement of the dangling pyridine.

In summary, we have isolated a rare example of a side-on $\text{Cr}(\text{IV})$ -peroxo adduct. This species is a catalytically competent species in solutions, and within crystals, for catalytic H_2O_2 dismutation. Its preparation involves the reaction of an unusually labile $\text{Cr}(\text{III})$ precursor and H_2O_2 . This is an interesting contrast to the syntheses of the handful of other known $\text{Cr}-\text{O}_2$ (superoxo, peroxo) adducts which typically involves the reaction of $\text{Cr}(\text{II})$ complexes with O_2 – precisely the reverse of step (vi) in Scheme 1.

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