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## Synthesis of a nanoscale Cu(II)<sub>31</sub>-oxo-carboxylate cluster, and effect of Cu-oxo cluster size on visible-light absorption†

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**Cu<sub>16</sub>-Cu<sub>31</sub> Cu(II)-oxo-carboxylate clusters are reported, including those with condensed 1.5 nm Cu-O cores supported exclusively by O-donor ligands. A size-colour correlation is observed due to a red-shift of the charge transfer absorption band on increasing size; hence these clusters sit between small molecules and (black) CuO nanostructures.**

Metal-oxo clusters are fascinating molecular analogues of metal oxide materials, with recent discoveries described as ‘molecular nanoparticles’.<sup>1</sup> They have well-defined and tuneable structures making them excellent options as photocatalysts,<sup>2,3</sup> molecules for energy conversion or storage,<sup>4</sup> molecular magnets,<sup>5</sup> or soluble precursors to semiconducting oxide materials.<sup>6,7</sup> Metal-oxo clusters may absorb light in a similar manner to their bulk oxide counterparts, however, as most are smaller than the exciton Bohr radius of the parent material, quantum confinement theory predicts a size-dependent blue-shift of their absorption onset relative to the bulk oxide.<sup>8</sup> This allows tuneability of optical properties which may be further optimised by adjusting surface interactions for efficient photochemical applications.

Cupric oxide (CuO) is an important semiconducting oxide material with applications in electrocatalysis (e.g. water splitting<sup>9,10</sup> or biomass valorisation<sup>11</sup>), heterogeneous catalysis,<sup>12</sup> and in photovoltaic cells.<sup>13</sup> Cupric oxide is a black *p*-type (strongly correlated) semiconductor with a small band gap (reported values range from ~1 – 2.1 eV)<sup>13–16</sup> attributed to {O + Cu(3d<sub>x<sup>2</sup>-y<sup>2</sup>)} → Cu(3d<sub>x<sup>2</sup>-y<sup>2</sup>)} transitions with significant oxygen to metal charge transfer character.<sup>15,16</sup> CuO has a moderately large exciton Bohr radius (~6 nm).<sup>17</sup> Therefore it is logical that quantum size effects should bring the absorption onset of small Cu(II)-oxo clusters into</sub></sub>

the mid-visible region (e.g. 2–2.8 eV), potentially providing appropriate photovoltage for challenging photocatalytic processes driven by sunlight, such as water oxidation.

Despite potential uses in photocatalysis or as precursors for the deposit of ultrasmall copper or copper-oxide-based particles, Cu(II)-oxo clusters with a condensed Cu(II) + O/OH core remain remarkably rare,<sup>5,18–26</sup> contrasting with the rich chemistry known for other metal-oxo clusters. Some interesting Cu-oxo structures have been reported with internal templates,<sup>20,21,25</sup> multidentate ligands,<sup>5,19,21,26</sup> or with mixed-metal systems;<sup>27</sup> however, in these structures the incorporation of halides, heteroatom donors, or heterometals, will inherently change the electronic properties and alter the relationship with CuO. Pioneering work by Christou *et al.* reported the structure of Cu<sub>16</sub>-oxo clusters [Cu<sub>16</sub>O<sub>4</sub>(OH)<sub>4</sub>(OR)<sub>8</sub>(O<sub>2</sub>C<sup>t</sup>Bu)<sub>12</sub>(ROH)], **1**, R = <sup>n</sup>Pr and **2**, R = <sup>n</sup>Bu, as rare examples of Cu-oxo clusters with exclusively O-donor ligands.<sup>18</sup> The use of multidentate carboxylate ligands has also led to the discovery of the Cu-hydroxo nanodisc [Cu<sub>15</sub>(dhs)<sub>6</sub>(OH)<sub>6</sub>(H<sub>2</sub>O)<sub>10</sub>](H<sub>2</sub>O)<sub>20</sub> (H<sub>4</sub>dhs = 2,3-dihydroxysuccinic acid),<sup>22</sup> and the 1D metal-organic chain [(CO<sub>3</sub>)@Cu<sub>20</sub>(suc)<sub>6</sub>(OH)<sub>26</sub>(H<sub>2</sub>O)<sub>12</sub>·18H<sub>2</sub>O]<sub>n</sub> (suc = succinate) comprising of Cu cages with the (8-capped) Keggin structure encapsulating a carbonate anion.<sup>28</sup>

In previous reports,<sup>18,26</sup> Cu-oxo clusters are synthesised from Cu(NO<sub>3</sub>)<sub>2</sub> and ligands in alcohol solutions, using a base to activate the Cu salt into more reactive species. We hypothesised that the base initiates Cu(II)-OR formation which is then susceptible to hydrolysis and condensation reactions to build Cu-O-Cu connectivity, leading to clusters, but which may retain some surface alkoxides (Scheme 1). Therefore, the alcohol solvent plays an important role in the assembly of clusters, affecting solubility and also providing a steric influence on the cluster surface during growth. Furthermore, as water is necessary for hydrolysis and cluster growth, the amount of water in solution is also a critical factor, with uncontrolled hydrolysis likely leading to extended CuO materials.

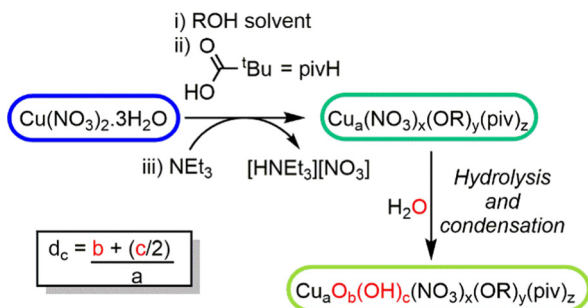
Using this approach with pivalic acid (<sup>t</sup>BuCO<sub>2</sub>H) as a supporting ligand and sterically unencumbered primary alcohols

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Scheme 1 Proposed reaction scheme for the growth of Cu-oxo clusters via hydrolysis and condensation of a 'pre-hydrolysis intermediate'.

( $R = \text{Et}$ ,  ${}^n\text{Pr}$ ,  ${}^n\text{Bu}$ ) leads to previously reported green  $\text{Cu}_{16}$  clusters **1** and **2** (Fig. S1–S3, ESI<sup>†</sup>),<sup>18</sup> or similarly structured ethoxide derivative **3**,  $[\text{Cu}_{16}\text{O}_4(\text{OH})_4(\text{OEt})_8(\text{O}_2\text{C}^t\text{Bu})_{12}(\text{H}_2\text{O})]$  which contains a symmetrically bound water molecule in its central pocket (Fig. 1 and Fig. S4, S5, ESI<sup>†</sup>). These structures all crystallise directly from alcohol solutions (or more rapidly from pentane, see ESI<sup>†</sup>), and have a degree of condensation ( $d_c$ ), defined as  $(\{n(\text{O}^{2-}) + 1/2n(\text{OH}^-)\}/n(\text{Cu}^{2+}))$ , of 0.375, some way from that of  $\text{CuO}$  ( $d_c = 1$ ), indicating that their structures are mostly 'surface'.

By switching to bulkier secondary alcohols ( $R = {}^i\text{Pr}$  or  $\text{Cyp}$  ( $\text{C}_5\text{H}_{11}$ )) grass-green crystals of larger clusters form after two weeks (in powdered form, or when dissolved, these compounds appear yellow/green). Isostructural forms were identified by X-ray crystallography, supported by bond-valence sum calculations, as  $[\text{Cu}_{31}\text{O}_{12}(\text{OH})_{18}(\text{O}_2\text{C}^t\text{Bu})_{18}(\text{NO}_3)_2(\text{ROH})_6 \cdot n(\text{ROH})]$  ( $R = {}^i\text{Pr}$ ,  $n = 6$ , **4**;  $R = \text{Cyp}$ ,  $n = 12$ , **5**) (Fig. 2 and Fig. S6–S8, ESI<sup>†</sup>). This structure has a pseudo-spherical Cu–O core (with 12  $\mu_4\text{-O}^{2-}$ , 12  $\mu_3\text{-OH}^-$  and 6  $\mu_2\text{-OH}^-$ ) with a maximum (Cu–O) core diameter of 1.5 nm and can thus be described as a 'molecular nanoparticle'. However, the Cu–O core is not a clear representation of the repeating structure in  $\text{CuO}$ . In  $\text{CuO}$  the copper centres coordinate to four oxygens in a close to square planar geometry (Cu–O bond lengths = 1.91–2.02 Å),<sup>29</sup> with longer distances (2.78 Å) to axially positioned oxygens. Similar

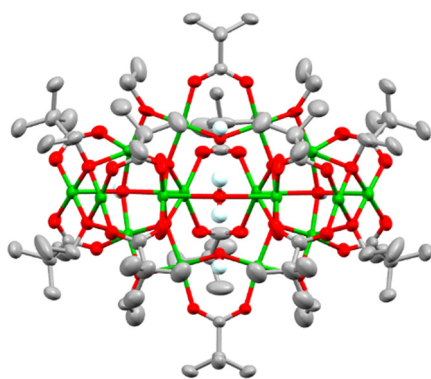


Fig. 1 Solid-state structure of ethoxide capped  $\text{Cu}_{16}$ -oxo cluster **3**. Ellipsoids displayed at 50% probability, hydrogen atoms (except OH or  $\text{H}_2\text{O}$ ) omitted for clarity. Cu = green, O = red, C = grey, H = very pale blue.

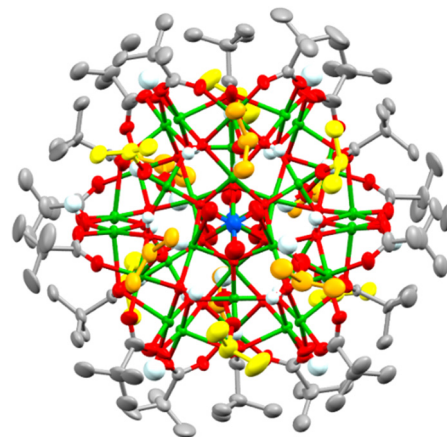


Fig. 2 Solid-state structure of  $\text{Cu}_{31}$ -oxo cluster **4**. Cu = green, O = red, C = grey, N = blue, H = very pale blue (non-coordinated/H-bonded  ${}^i\text{PrOH}$  Cs = yellow, coordinated  ${}^i\text{PrOH}$  Cs = orange). Ellipsoids displayed at 50% and hydrogen atoms (except OH) omitted.

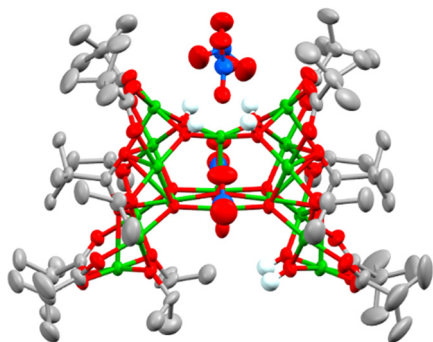
coordination modes are found in **4** and **5** with comparable Cu–O bond lengths to the four nearest oxygens (range 1.88–2.28 Å) and most sites having a fifth oxygen occupying an apical site (range 2.29–2.89 Å, Supporting note 1, Fig S9, S10 and Table S1, ESI<sup>†</sup>). In contrast, the central Cu atom in **4** or **5** has a tilted octahedral geometry (Cu–O bond lengths: 2.05 to 2.22 Å). The cluster lies on a pseudo- $\text{C}_3$  symmetry axis passing through the centrosymmetric octahedral Cu and the N atoms of the two nitrates (Fig. 2) with overall pseudo- $\text{D}_{3d}$  symmetry. The  $d_c$  of **4** and **5** is 0.68, showing a much more condensed core and thus greater resemblance to  $\text{CuO}@$ carboxylate nanoparticles than the smaller clusters **1–3**. The surface of **4** and **5**, which can be considered a model for  $\text{CuO}@$ carboxylate nanoparticles,<sup>30</sup> is capped by 18 anionic ( $\mu_2$  or  $\mu_3$ ) pivalate ligands, 18 hydroxides, six neutral ROH ligands and two loosely coordinated  $\mu_2$ -nitrate anions (Cu– $\text{ONO}_2$  bond lengths 2.6–2.8 Å), which have disordered coordination geometries in the crystal structure and are each surrounded by the three coordinated and a further three non-coordinated ROH ligands (which likely engage in H-bonding interactions to the cluster surface, Fig. 2, Fig. S6 and S8, ESI<sup>†</sup>). Crucially these clusters have no remaining alkoxide groups (confirmed by bond valence sum calculations, Table S3, ESI<sup>†</sup>) which may provide enhanced stability to further hydrolysis and condensation reactions. The greater steric bulk of secondary alcohols appears to support condensation and enlargement of the core compared to less bulky primary alcohols. The synthesis of **4** and **5** is very condition-dependent, with brown precipitates containing  $\text{CuO}@$ pivalate nanoparticles ( $\sim 8\text{--}15$  nm by Scherrer analysis) and  $\text{Cu}_2(\text{OH})_3(\text{NO}_3)$  (Fig S11 and S12, ESI<sup>†</sup>) alongside solutions containing the paddlewheel  $[\text{Cu}_2(\text{O}_2\text{C}^t\text{Bu})_4(\text{NEt}_3)_2]$  able to form as by-products if the reaction is conducted at higher concentrations; left open to ambient atmosphere; or allowed to sit for longer times. The by-products likely form due to uncontrolled hydrolysis. It appears essential to maintain good solubility of intermediates



to allow **4/5** to self-assemble without the formation of insoluble by-products.

Based on our observations and hypothesised reaction mechanism (Scheme 1), moisture plays a crucial role in cluster synthesis. However, the copper source  $\text{Cu}(\text{NO}_3)_2 \cdot 3\text{H}_2\text{O}$  contains three equivalents of water per copper atom, thus always provides an excess of water for hydrolysis and condensation reactions. Therefore, selective cluster formation is likely defined by crystallisation of the growing clusters from the reaction medium (for example when **3** is dissolved in dichloromethane or isolated as a powder it is susceptible to further hydrolysis when exposed to moisture). To test these concepts, the synthesis of clusters using anhydrous precursors was explored.  $\text{Cu}(\text{OEt})_2$  was reacted with 1 equiv. of pivalic acid in dry toluene at  $70^\circ\text{C}$  to produce previously reported loop-shaped  $[\text{Cu}_6(\text{OEt})_6(\text{O}_2\text{C}^t\text{Bu})_6]$ , **6**,<sup>18,31</sup> (Fig. S13, ESI†). This cluster is a 'pre-hydrolysis' structure as it contains no oxo or hydroxide groups ( $d_c = 0$ ), and could potentially form at the beginning of the reaction between  $\text{Cu}(\text{NO}_3)_2 \cdot 3\text{H}_2\text{O}$ , pivalic acid and  $\text{NEt}_3$  in ethanol, *i.e.* before any hydrolysis (Scheme 1).<sup>32</sup> The reaction of **6** with 2.7 equivalents of water in EtOH induces a slow colour change from blue-green to green, and a precipitate of **3** forms after two weeks (confirmed by diffraction after recrystallisation), demonstrating that water promotes cluster growth.<sup>32</sup>

Considering that water plays a crucial role in cluster formation, the reaction conditions to produce **4** were repeated, but, after 30 minutes of reaction (under air), the solution was degassed and placed under  $\text{N}_2$  over activated  $3\text{ \AA}$  molecular sieves. The flask was kept under air-free conditions and after one month, crystals of a new structure, **7**,  $[\text{Cu}_{17}\text{O}_5(\text{OH})_6(\text{O}^t\text{Pr})_6(\text{O}_2\text{C}^t\text{Bu})_{10}(\text{NO}_3)_2] \cdot ([\text{NEt}_3\text{H}][\text{NO}_3])_2$ , reproducibly form. **7** has a  $d_c$  of 0.47 ( $> 1-3$  but  $< 4-5$ ) indicating that this structure has formed under partial hydrolysis conditions, *i.e.* before **4** forms (Fig. 3 and Fig. S14–S16, ESI†). This also indicates that hydrolysis and cluster growth is a relatively slow process as it can be arrested by removing moisture from the reaction solution after 30 minutes of reaction. The structure of **7** maintains some  $\text{Cu}-\text{O}^t\text{Pr}$  functionality, suggesting that it may be susceptible to further reaction with water. **7** is a C-shaped cluster with an intriguing pocket which could act as a sterically

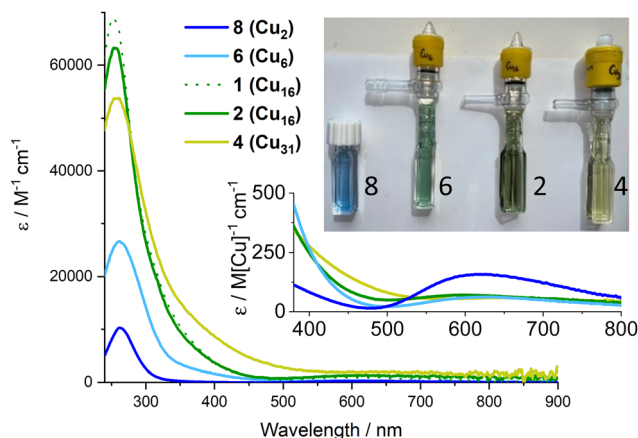


**Fig. 3** Solid-state structure of **7**. Ellipsoids displayed at 50% probability, hydrogen atoms (except OH groups) and  $[\text{NEt}_3\text{H}]^+$  cations omitted for clarity. Cu = green, O = red, C = grey, N = blue, H = very pale blue. The base of the structure (within the pocket) has disordered  $\text{O}^t\text{Pr}$  and OH groups.

encumbered reactive site. **7** contains 17 5-coordinate Cu centres which are supported by eight  $\mu_2$ -bridging pivalate anions and two unusually coordinated pivalates that each coordinate to seven Cu centres (Fig. S17, ESI†). Two chelating nitrates ( $\text{Cu}-\text{O} = 2.02-2.65\text{ \AA}$ ) and four bridging isopropoxide groups coordinate around the edges of the cluster, whilst a combination of two isopropoxides and two hydroxides occupy the internal cleft of the C-shape in a disordered manner (Fig. S18, ESI†). Two further nitrate anions are connected *via* H-bonding to four  $\text{Cu}-\text{OH}$  sites which sit in a square on the top of the cluster.

The relatively asymmetric shapes and range of different Cu carboxylate coordination geometries in these clusters highlights the flexibility of the  $\text{Cu}(\text{II})$  centre and the unpredictability of the structures that it can form. This implies that many further possible cluster structures can be obtained by variation of the supporting ligands and degree of condensation.

The colours of these Cu-oxo clusters suggests a relationship with size (Fig. 4 and Fig. S19, S20, ESI†), as the compounds change from blue through green to yellow/green with increasing size, eventually reaching dark brown/black  $\text{CuO}$  nanoparticles (Fig. S21, ESI†). The UV/visible absorption spectra of **1-4**, **6**, the small  $\text{Cu}_2$  'paddlewheel'  $[\text{Cu}_2(\text{O}_2\text{C}^t\text{Bu})_4(\text{HOR})_2]$  ( $R = ^t\text{Pr}$  or H), **8** (Fig. S22, ESI†), and  $\text{Cu}(\text{NO}_3)_2$  were recorded (Fig. 4 and Fig. S23–S31, ESI†). All display Beer–Lambert behaviour. The well-known paddlewheel structure (with two carboxylates per Cu) is blue in solution (Fig. 4) and has a high energy carboxylate (O) to metal ( $d_{x^2-y^2}$ ) charge transfer (LMCT) band in the UV region<sup>33,34</sup> (**8**:  $\lambda_{\text{max}} = 262\text{ nm}$ ;  $\epsilon \sim 10\,300\text{ M}^{-1}\text{ cm}^{-1}$ ) with a shoulder ("dimer band") arising from the bridging carboxylate coordination mode at slightly lower energy, and weaker d–d transitions (to  $d_{x^2-y^2}$ ) in the visible region ( $\lambda_{\text{max}} = 615\text{ nm}$ ;  $\epsilon \sim 300\text{ M}^{-1}\text{ cm}^{-1}$ ) which cause the blue colour. Compound **6** is blue/green in solution and shows a similar LMCT peak  $\lambda_{\text{max}} = 261\text{ nm}$ , and a shoulder which now extends into the visible region, possibly from added contributions of alkoxide to Cu LMCT,<sup>8</sup> and a weaker d–d absorption per Cu (Fig. 4). Upon growth of the cluster to (green)  $\text{Cu}_{16}$  (**2**) and (yellow/green)  $\text{Cu}_{31}$  (**4**), the



**Fig. 4** UV/vis spectra of **2**, **4**, **6** & **8** reported as molar extinction coefficients. Spectra recorded at  $0.1-0.4\text{ mM}[\text{Cu}]$  in  $\text{CH}_2\text{Cl}_2$  (or pentane for **2**), inset shows magnified visible region as 'per copper' normalised extinction coefficients, spectra recorded at  $1.7-3.3\text{ mM}[\text{Cu}]$ . Photos of solutions (with  $[\text{Cu}] = \sim 0.24\text{ mM}$ ).





