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## Subsurface Oxygen in Oxide-Derived Copper Electrocatalysts for Carbon Dioxide Reduction

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# Subsurface Oxygen in Oxide-Derived Copper Electrocatalysts for Carbon Dioxide Reduction

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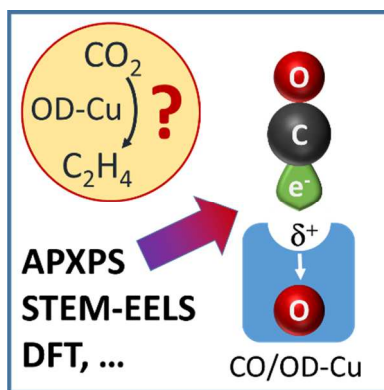
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## ABSTRACT

Copper electrocatalysts derived from an oxide have shown extraordinary electrochemical properties for the carbon dioxide reduction reaction (CO<sub>2</sub>RR). Using *in situ* Ambient Pressure X-ray Photoelectron Spectroscopy (APXPS) and quasi *in situ* Electron Energy Loss Spectroscopy (EELS) in a Transmission Electron Microscope (TEM), we show that there is a substantial amount of residual oxygen in nanostructured, oxide-derived copper electrocatalysts, but no residual copper oxide. Based on these findings in combination with Density Functional Theory (DFT) simulations, we propose that residual subsurface oxygen changes the electronic structure of the catalyst and creates sites with higher carbon monoxide binding energy. If such sites are stable under the strongly reducing conditions found in CO<sub>2</sub>RR, these findings would explain the high efficiencies of oxide-derived copper in reducing carbon dioxide to multi-carbon compounds such as ethylene.



**KEYWORDS** In Situ, Ambient-Pressure X-Ray Photoelectron Spectroscopy, Scanning Transmission Electron Microscopy, Electron Energy Loss Spectroscopy, Density Functional Theory

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3 A promising path towards a sustainable energy future with net-neutral CO<sub>2</sub> emissions is the  
4 combination of renewable electricity production with the electroreduction of carbon dioxide  
5 (CO<sub>2</sub>RR), that needs a suitable electrocatalyst.<sup>1,2</sup> Among all pure metals, copper is the only one  
6 that gives a rich gamut of single- and multi-carbon products,<sup>3-5</sup> which can be explained by an  
7 optimal binding energy of CO to the surface.<sup>6,7</sup> However, the explanation becomes more  
8 complicated when dealing with nanostructured copper materials derived from oxidized  
9 precursors, which show both higher activity and greatly enhanced selectivity towards multi-  
10 carbon products.<sup>8-16</sup> It has been hypothesized that this could be related for instance to an increase  
11 of local pH,<sup>17-19</sup> grain boundaries,<sup>20</sup> undercoordinated sites,<sup>21</sup> or residual oxides.<sup>22-24</sup>

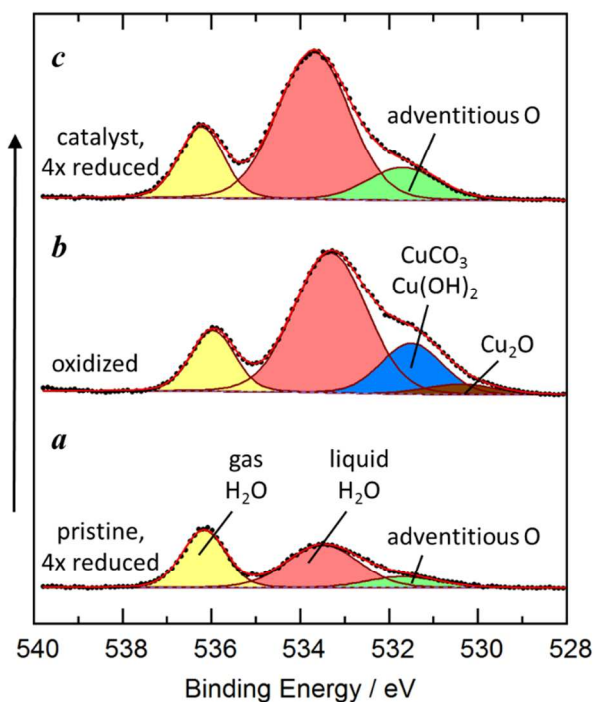
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25 Several groups have shown the existence of sites with higher CO binding energy in oxide-  
26 derived copper.<sup>23,25</sup> However, if the increased binding energy is caused by a simple shift of the *d*-  
27 band in the catalyst (such as if we had nickel instead of copper), scaling relations would actually  
28 predict a higher overpotential for CO-CO-coupling on Cu(100).<sup>7</sup> Furthermore, an experimental  
29 study of different transition metals also suggests that an increase in CO binding energy actually  
30 lowers CO<sub>2</sub>RR activity.<sup>5</sup> In contrast, if the increased binding energy has a different origin, such  
31 an understanding could become a future strategy to overcome scaling relations.

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41 A polycrystalline copper foil was exposed to electrochemical oxidation-reduction cycles that  
42 increased the overall CO<sub>2</sub>RR activity of the catalyst and improved the product yield towards  
43 more ethylene vs. methane (Figure S1). This change in activity and selectivity comes together  
44 with a pronounced change in the O 1s APXPS (Figure 1). The reported photoelectron spectra  
45 were measured *in situ*, i.e. in the chemical environment of a few nm thick layer of electrolyte (4  
46 mM KCl in 0.1 M KHCO<sub>3</sub>), immediately after pulling the sample out of the beaker (Figure  
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55 S2).<sup>26-28</sup>

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Figure 1a shows a spectrum of the pristine copper foil after four electrochemical reduction cycles, to remove surface oxides. The spectrum consists of a gas phase water peak at 536.2 eV apparent binding energy (yellow), a liquid water peak at 533.7 eV (red) and a third, adventitious oxygen peak at 531.7 eV (green). The intensity of the third (green) peak does not decrease upon further reduction as shown in Figure S3. Its origin is discussed below. Shifts induced by an electrochemical double-layer can be excluded due to the sufficiently high concentration of the electrolyte used.<sup>29</sup> The corresponding Cu  $2p_{3/2}$  spectrum (Figure 2a) shows metallic copper. However, small contributions from Cu(I) cannot be excluded as Cu(I) compounds are typically shifted by only ca. 100 meV to lower binding energies with respect to metallic copper.<sup>30</sup>

After oxidation, the O  $1s$  spectrum (Figure 1b) gains two components that can be assigned to  $\text{CuCO}_3/\text{Cu}(\text{OH})_2$  and  $\text{Cu}_2\text{O}$ , respectively.<sup>30,31</sup> Possible underlying intensity of the aforementioned adventitious oxygen peak at 531.7 eV cannot be resolved due to overlap with these components. The corresponding Cu  $2p_{3/2}$  spectrum (Figure 2b) shows additional intensity at 934.6 eV which is in accordance with  $\text{CuCO}_3/\text{Cu}(\text{OH})_2$ , whereas the  $\text{Cu}_2\text{O}$  fraction cannot be resolved due to the small shift of Cu(I) species. Scanning the Cl  $2p$  spectral region did not show any significant intensity (Figure S4), proving the absence of CuCl compounds in spite of the presence of 4 mM chloride in the electrolyte (which is below the detection limit), which is in accordance with an earlier XAS study on a similar system.<sup>9</sup>

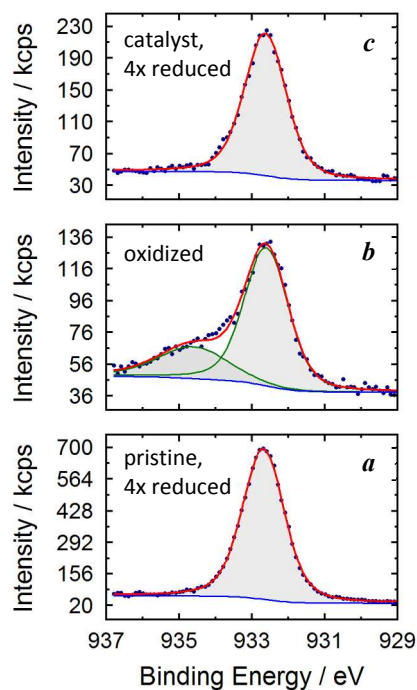


**Figure 1.** In situ O 1s APXPS spectra measured with an incident photon energy of 3996 eV. (a) Pristine sample. (b) Oxidation of the pristine sample leads to additional intensity at binding energies below 533 eV due to the formation of oxidized compounds such as CuCO<sub>3</sub>, Cu(OH)<sub>2</sub> and Cu<sub>2</sub>O. Furthermore, the liquid water peak increased indicating a thicker water overlayer. (c) The initially oxidized and then reduced sample contains significantly more adventitious oxygen (green) than the pristine sample before oxidation. Intensities not normalized.

After reduction, the O 1s spectrum (Figure 1c) shows a substantially increased adventitious component at 531.7 eV (green), compared to the initial spectrum before oxidation. Furthermore, the oxidized and reduced samples give rise to an increased amount of liquid water on the sample, visible both by an increased liquid water peak in the O 1s spectrum and by a stronger attenuation of the Cu 2p line. This behavior was reproducible and not an effect of the background pressure of water vapor which was slightly lower for the oxidized and reduced samples, compared to the

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4 pristine one. The fact that the oxidized and reduced samples show a similar moist content (which  
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6 is significantly higher than for the pristine sample) points away from surface polarity being the  
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8 main cause of this effect. It is likely that mainly nano-structuring of the sample surface (Figure  
9  
10 S5) causes the increased hydrophilicity.  
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13 For quantification of the relative increase of adventitious oxygen, attenuation by the water  
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15 overlayer has to be considered. Figure S6 describes different normalization schemes which are  
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17 based on different assumptions. As a result, the increase of adventitious oxygen after oxidation  
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19 and four subsequent reduction cycles can be estimated to be in the order of a factor 10.  
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48 **Figure 2.** *In situ* Cu  $2p_{3/2}$  APXPS spectra. Spectra of the reduced sample (*a*, *c*) do not contain  
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50 any Cu(II). The oxidized spectrum (*b*) shows an additional Cu(II) component such as malachite  
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52 ( $\text{CuCO}_3/\text{Cu}(\text{OH})_2$ ).  
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3 We also observed K  $2p$  intensity and additional C  $1s$  intensity at 289 eV (Figure S7)  
4 sporadically associated with samples, which roughly scaled with each other. This indicates that  
5 the electrolyte contributes to the adventitious (green) oxygen component in the O  $1s$  spectrum  
6 (Figure 1), especially since the O  $1s$  binding energy of carbonate is centered at about 531.7 eV.<sup>30</sup>  
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8 However, whereas a strong correlation between adventitious oxygen and sample history is  
9 present (*i.e.* whether it was pre-oxidized or not), the correlation between K  $2p$  and adventitious O  
10  $1s$  intensity is rather weak, giving rise to the conclusion that the adventitious oxygen signal is not  
11 solely caused by electrolyte. Furthermore, potential dependent XPS using the dip-and-pull  
12 method<sup>26,27</sup> show that potassium is electronically not connected to the Cu working electrode  
13 (observable by a potential-dependent shift in apparent binding energy), whereas the best fit  
14 indicates that the adventitious oxygen component is mostly pinned to the Cu working electrode  
15 (Figure S8). This supports the hypothesis that an additional species, which is electronically  
16 connected to the working electrode, contributes to the adventitious (green) component in the O  
17  $1s$  spectrum (Figure 1). The appearance of electrolyte in the spectra is further discussed in the  
18 Supporting information.

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38 Recently, the existence of residual  $\text{Cu}_2\text{O}$  in copper catalysts for  $\text{CO}_2\text{RR}$  was discussed.<sup>23,24</sup>  
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40 However, the additional (green) peak at 531.7 eV in the O  $1s$  spectrum cannot be caused by  
41  $\text{Cu}_2\text{O}$  which would have been observed at 530.4 eV (compare to Figure 1*b*). Furthermore, Cu(II)  
42 components can be excluded by the Cu  $2p_{3/2}$  spectrum (Figure 2*c*) that shows no peak at high  
43 binding energies. Other imaginable Cu(I) components, such as the hypothetical Cu(OH) or  
44  $\text{Cu}_2\text{CO}_3$ , are generally not stable,<sup>32</sup> but cannot entirely be excluded.

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53 It is a reasonable hypothesis that subsurface oxygen contributes to the additional (green)  
54 spectral component in Figure 1*c*, overlapping with the aforementioned electrolyte contribution.  
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56 Also a contribution from specifically adsorbed  $\text{HCO}_3^-$ , supported by subsurface oxygen, is  
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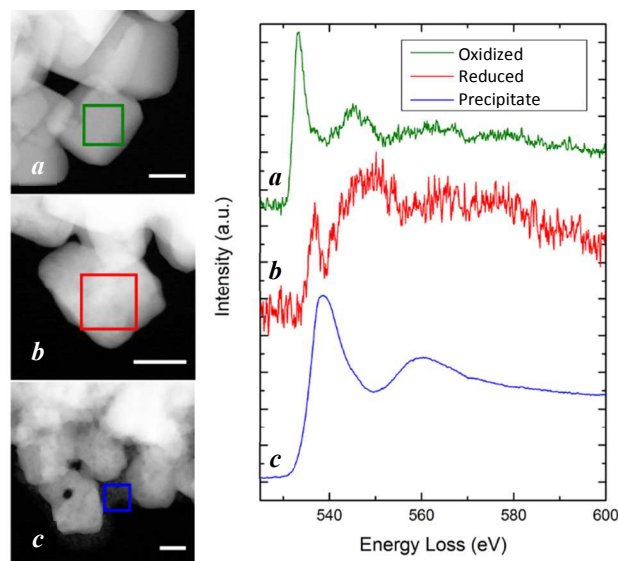
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imaginable. The inelastic mean free path of O 1s photoelectrons at a kinetic energy of 3470 eV in metallic copper is 3.9 nm.<sup>33</sup> Since the attenuation of photoelectrons follows an exponential decay with increasing depth, the majority of the observed signal stems from the top few nanometers of the catalyst. From a theoretical point of view, Korzhavyi and Sandström have calculated the trapping barriers of oxygen and OH<sup>-</sup> in a monovacancy of Cu as 0.95 and 1.23 eV, respectively, and excluded water formation in this site, which points to the fact that oxygen could indeed be stable in certain subsurface sites in an aqueous environment and in the presence of hydrogen.<sup>34</sup>

Bluhm and co-workers have found an XPS spectroscopic signature of subsurface oxygen earlier in a copper catalyst for methanol oxidation.<sup>35</sup> In their study, peak assignments are tentative due to the challenging interpretation of data with many components and limited signal-to-noise ratio at high photon energies. Our Density Functional Theory (DFT) calculations suggest a relative difference of XPS binding energy between subsurface and surface oxygen of 0.8-1.7 eV (Table S2). Together with the known XPS binding energy of surface oxygen of ca. 530 eV,<sup>36</sup> it is consistent with the hypothesis that subsurface oxygen contributes to the additional spectral component at 531.7 eV. This is also in accordance with the silver-oxygen system that shows subsurface oxygen shifted towards higher XPS binding energy compared to surface oxygen.<sup>37</sup>

This conclusion is further backed by quasi *in situ* oxygen K-edge Electron Energy Loss Spectra (EELS) of a copper nanoparticle catalyst in a Transmission Electron Microscope (TEM) (Figure 3). In this experiment, electrochemical modifications and electrolyte removal by rinsing in water were performed in a protective nitrogen atmosphere outside the microscope, followed by sample transfer into the microscope using a vacuum transfer holder. Due to this experimental scheme, the sample was never in contact with air during the experiment. Figure 3a shows the oxidized sample with an EELS signature resembling that of Cu<sub>2</sub>O.<sup>38-40</sup> Experimental details were

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3 slightly different between EELS and APXPS (nanoparticles versus foil and lower pH in EELS),  
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5 resulting in Cu<sub>2</sub>O being the dominant species in EELS. After reduction, two chemically different  
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7 oxygen species were found: one inside the catalyst, assignable to subsurface oxygen (*b*) and one  
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9 in a precipitate on the surface of the catalyst (*c*). The precipitate contains K, C and O (Figure S9)  
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11 and could be related to K<sub>2</sub>CO<sub>3</sub> precipitation; more details can be found in the Supporting  
12  
13 information, section ‘Appearance of electrolyte in the APXPS spectra’. Both spectra (*b*) and (*c*)  
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15 in Figure 3 exhibit a different line shape and a shift of +3.2 and +3.5 eV with respect to spectrum  
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17 (*a*), respectively. This gives clear evidence for the changed chemical environment of subsurface  
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19 oxygen with respect to Cu<sub>2</sub>O. The energy shift of the oxygen K edge to higher energy losses in  
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21 the reduced sample (*b*) compared to the oxidized one (*a*) is consistent with that observed in the O  
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23 1s APXPS (Figure 1), where adventitious oxygen (green) was found at ca. 1.3 eV higher binding  
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25 energy than Cu<sub>2</sub>O (brown). The different values for the energy shift between EELS and APXPS  
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27 are caused by the non-metallic nature of the oxidized sample, that does not allow metallic  
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29 screening in the APXPS of Cu<sub>2</sub>O.<sup>41</sup> However, the similar shifts in Figs. 3*b* and (*c*) support the  
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31 modelling of adventitious oxygen in APXPS by one peak (Figure 1, green component). Copper  
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33 L-edge EELS spectra of the oxidized and reduced catalyst confirm Cu<sub>2</sub>O and metallic copper,  
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35 respectively (Figure S11).  
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**Figure 3.** Scanning TEM annular dark field (ADF) images (left) and oxygen K EEL spectra (right) obtained in a quasi *in situ* TEM experiment mimicking the APXPS experiment. *a*) Oxidized sample: spectroscopic EELS signature of  $\text{Cu}_2\text{O}$ ; *b,c*) reduced sample, *b*) EELS from a particle and *c*) EELS from the precipitate. Scale bars in the TEM images correspond to 30 nm. Squares in the TEM images depict the scanned areas for EELS. Edge shifts match qualitatively with APXPS binding energy shifts.

Recently, Lee *et al.*<sup>23</sup> and Mistry *et al.*<sup>24</sup> proposed residual  $\text{Cu}_2\text{O}$  after reduction that would explain the enhanced selectivity of oxide-derived Cu. Their *in situ* XANES data are consistent with a similar study by Eilert *et al.*<sup>9</sup> and an *in situ* Raman spectroscopy study by Ren *et al.*,<sup>11</sup> that showed a pure metallic phase after holding the sample at reductive potentials, but their *ex situ* TEM work that occurred after long exposure to air and *post mortem* sample preparation showed the presence of an oxide phase (a reference sample reduced with  $\text{H}_2$  indicated that the oxide phase was not solely caused by exposure to air). However, in the present study our APXPS experiment was carried out *in situ* and our TEM experiment in a quasi *in situ* environment

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3 without exposure to air and shows that there are no residual oxides but a pure metallic phase with  
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5 a small amount of subsurface oxygen.  
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8 It is worth noting the difference between residual copper oxides such as  $\text{Cu}_2\text{O}$  or  $\text{Cu}(\text{OH})_2$  and  
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10 trapped subsurface oxygen or  $\text{OH}^-$ . Since the catalyst is formed by reducing an oxidized Cu  
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12 sample, it is supposed that subsurface oxygen is still in a negative oxidation state, which is  
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14 compensated by polarization of the conduction electrons in the surrounding, mostly metallic  
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16 copper atoms. This leads to substantially different properties and crystal structure compared to  
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18 the oxide phase, which contains Cu in an oxidation state +I or +II.  
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22 CO bonding on late transition metals such as copper is governed by an interplay between  $\pi$ -  
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24 bonding and  $\sigma$ -repulsion.<sup>42-44</sup> Both depend on the  $d$ -band position, however, the  $\sigma$ -repulsion is  
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26 also sensitive to the occupancy of the Cu  $sp$ -band. Thus an electron acceptor which withdraws  
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28 charge from the  $sp$ -band can reduce the repulsion as shown by Xin *et al.* for the case 2O-  
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30 CO/Ru(0001).<sup>45</sup> In particular the free energy at elongated distances in comparison to the  
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32 equilibrium position is affected.<sup>45</sup>  
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36 As an example, the influence of subsurface oxygen on the binding energy of CO on a Cu(100)  
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38 surface was investigated by DFT. The CO binding energy was calculated for clean Cu(100) and  
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40 for Cu(100) with an oxygen atom placed in interstitial sites in different layers below the surface  
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42 (Table S3). Significant CO binding energy enhancement is observed for oxygen atom  
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44 implantations down to the 3<sup>rd</sup> subsurface layer.  
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48 Hence, we find that the interaction of the subsurface oxygen atom with the metal causes an  
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50 increased CO binding energy (Figure S12). A larger CO binding energy could lead to a higher  
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52 CO coverage of the catalyst, which kinetically favors C-C-coupling over hydrogenation. There  
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54 can be additional effects in the C-C coupling at longer CO-metal distances when electrolyte and  
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56 water are present since the oxygen-induced changes minimize the  $\sigma$ -repulsion from the more  
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3 spatially extended *sp*-band wave functions.<sup>45</sup> This could solve the apparent contradiction  
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5 between studies by Ma *et al.*<sup>7</sup> and Kuhl *et al.*<sup>5</sup> on the one side, which propose (on the basis of *d*-  
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7 band scaling relations) no improved catalytic properties for catalysts with higher CO binding  
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9 energy compared to clean Cu, and studies by Lee *et al.*<sup>23</sup> and Verdaguer-Casadevall *et al.*,<sup>25</sup>  
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11 which show sites with an increased CO binding energy in oxide-derived Cu. The question is,  
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13 however, whether subsurface oxygen could be stable close to the surface under such reducing  
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15 conditions and if so it would be expected to require some unique sites created in the  
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17 nanostructured system upon reduction of the oxide precursor. This question could be addressed  
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19 by XPS with lower kinetic energy and by TEM with higher resolution and without precipitate.  
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25 The recent enhanced CO<sub>2</sub> selectivity correlation to grain boundaries in vapor deposited Cu on  
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27 carbon nanotubes<sup>20</sup> could also be due to subsurface oxygen since the samples were exposed to air  
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29 prior to the electrocatalytic measurements. Similarly the enhanced CO adsorption energy on  
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31 reduced oxide-derived Cu is correlated to the amount of grain boundaries.<sup>25</sup> This could point to  
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33 the fact that subsurface oxygen could preferentially be more stable and enhancing the reactivity  
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35 of Cu metallic sites at grain boundaries.  
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39 Several studies have shown a preference for C-C-coupling on Cu(100) facets<sup>8,46</sup> and Li *et al.*  
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41 proposed a break in scaling relations, caused by the stabilization of \*C<sub>2</sub>O<sub>2</sub> on the particular  
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43 surface geometry of Cu(100).<sup>47</sup> Furthermore, it was shown that an increased local pH, caused by  
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45 the consumption of protons in the pores of a rough catalyst, also promotes C-C-coupling.<sup>17,18</sup>  
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47 However, studies which investigated those effects particularly for oxide-derived copper came to  
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49 the conclusion, that neither the exposed facets<sup>15</sup> nor the increase in local pH<sup>19</sup> can sufficiently  
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51 explain the extraordinary properties of these catalysts. An increased CO binding energy due to  
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53 subsurface oxygen can thus be the crucial puzzle piece which makes oxide-derived copper so  
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55 special for CO<sub>2</sub> reduction.  
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3 In conclusion, we have shown by *in situ* APXPS and quasi *in situ* EELS a correlation between  
4 favorable electrochemical properties of an oxide-derived copper catalyst for the electrochemical  
5 reduction of carbon dioxide, such as higher activity and higher yields of ethylene versus methane  
6 formation, and the existence of subsurface oxygen which is formed by oxidation-reduction  
7 cycles. Both APXPS and EELS prove the absence of residual copper oxide in the reduced  
8 electrocatalyst. We propose a mechanism in which subsurface oxygen increases the CO binding  
9 energy to the catalyst by reducing the  $\sigma$ -repulsion. This mechanism kinetically favors C-C-bond  
10 formation due to a higher CO coverage on the catalyst. However, this could only occur if some  
11 special sites in the near-surface region sufficiently inhibit diffusion to the first layer where  
12 oxygen would subsequently be protonated to water. This finding provides further insight on the  
13 “oxide-derived copper story” as it contributes to the explanation of the outstanding change in  
14 electrochemical properties due to oxidation and reduction of copper.  
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## 35 ASSOCIATED CONTENT

### 36 37 38 **Supporting Information.**

39 The Supporting Information is available free of charge.

40  
41 Spectra, illustrations, detailed discussions of side aspects, experimental and computational  
42 details (PDF)  
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## 49 AUTHOR INFORMATION

### 50 51 **Notes**

52 The authors declare no competing financial interests.  
53  
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