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Dynamic Changes in the Structure, Chemical State and Catalytic Selectivity of Cu Nanocubes during CO₂ electroreduction: Size and Support Effects

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Abstract

In situ and *operando* spectroscopic and microscopic methods were used to gain insight into the correlation between the structure, chemical state, and reactivity of size- and shape-controlled ligand-free Cu nanocubes (Cu-cubes) during CO₂ electroreduction (CO₂RR). Dynamic changes in the morphology and composition of Cu-cubes supported on carbon were monitored under potential control via electrochemical atomic force microscopy, X-ray absorption fine-structure spectroscopy and X-ray photoelectron spectroscopy. Under reaction conditions, the roughening of the nanocube surface, disappearance of the (100) facets, formation of pores, loss of Cu and reduction of CuO_x species observed were found to lead to a suppression of the selectivity for multi-carbon products (i.e. C₂H₄ and ethanol) versus CH₄. A comparison with Cu-cubes supported on Cu foils revealed an enhanced morphological stability and persistence of Cu(I) species under CO₂RR. Both factors are held responsible for the higher C₂/C₁ product ratio observed for the Cu cubes/Cu as compared to Cu cubes/C. Our findings highlight the importance of the structure of the active nanocatalyst but also its interaction with the underlying substrate in CO₂RR selectivity.

Introduction

Since a number of years there is an increasing interest in trying to understand the parameters that govern the electrochemical reduction of CO₂ (CO₂RR) to design catalysts with high selectivity for valuable chemicals and fuels.^[1] Among these parameters, the catalyst structure and chemical state are of particular importance.^[2–4] Compared to polycrystalline Cu electrodes, nanostructured Cu catalysts have shown a significantly improved CO₂RR performance, attributed to grain boundaries^[5,6], Cu(100) facets^[7–12], increased roughness^[13], defects^[14–16], low-coordinated sites^[17–19], and the presence of subsurface oxygen and Cu(I) species^[20–29].

Previous studies^[9–11] on Cu single crystals have shown the improved C-C coupling performance of (100) facets, which was further confirmed by the high selectivity towards ethylene observed on cubic-shaped Cu catalysts^[7,8,16,21,30–36]. However, the presence of (100) facets is not the only factor responsible for the superior activity and selectivity of cubic-shaped Cu catalysts, with surface roughness, subsurface oxygen and Cu(I) species or Cu/Cu(I) interfaces formed and/or stabilized under reaction conditions also playing a very important role.^[21] The function of oxygen in such structures is particularly intriguing, since on Cu(100) surfaces it was discussed to contribute to the formation of oxygenated hydrocarbons^[37]. The complexity arising from the multiple factors affecting the catalytic performance of cubic-shaped Cu catalysts requires a systematic study of the evolution of their structure and oxidation state under *operando* CO₂RR conditions.

This work focuses on the understanding of the relative importance of the different factors responsible for specific selectivity trends observed for Cu catalysts during CO₂RR, namely, the presence and stability of Cu(100) facets, defect sites, and the content of Cu(I) species and/or subsurface oxygen. By electrochemically growing ligand-free Cu-cubes on C supports and comparing with analogous samples supported on Cu foils, we were able to reveal the intrinsic behavior of the cubic-shaped Cu NPs catalysts and their dynamic evolution under CO₂RR conditions.

Results and Discussion

Size-dependent changes in the morphology and composition of Cu-cube samples electrodeposited on carbon paper were investigated *ex situ* via scanning electron microscopy (SEM) and energy dispersive X-ray spectroscopy (EDX) before and after CO₂RR, Figures 1, S1 and S2 (Suppl. docs). In all samples the Cu-cubes were found to decrease in size in the course of the first 1 h of the reaction. Furthermore, in this process the originally flat facets and sharp edges of the Cu-cubes were found to roughen, resulting in a porous nanocube surface, Fig. 1.

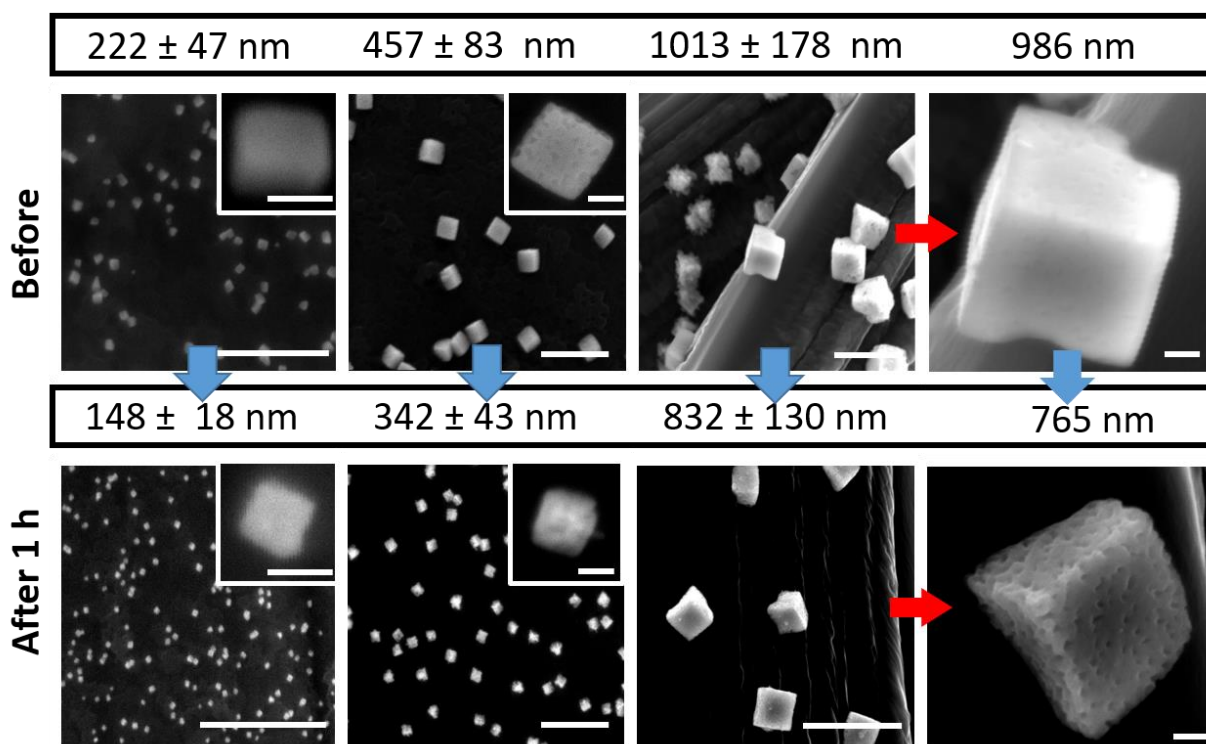


Figure 1: SEM images from size-controlled Cu-cube samples electrodeposited on carbon paper acquired before and after CO₂RR at -1.05 V for 1 h. The scale bars in the main panels are 2 μm, those in the insets and in the images of the last column are 200 nm.

Operando electrochemical atomic force microscopy (EC-AFM) measurements carried out on a sample with ~100 nm large cubes also revealed significant changes in their morphology already prior to the actual CO₂RR. Sample immersion in the aqueous 0.1 M KHCO₃ electrolyte at open circuit potential was observed to lead to the formation of “cracks” on the cube facets. These cracks are likely the result of mechanical stress taking place during the solvation of Cl⁻ ions while being transferred from the cubes to the electrolyte. SEM-EDX data revealed that Cl

was present throughout the entire Cu volume in the as prepared samples and therefore, the loss of Cl leads to drastic structural changes in the entire cube volume.

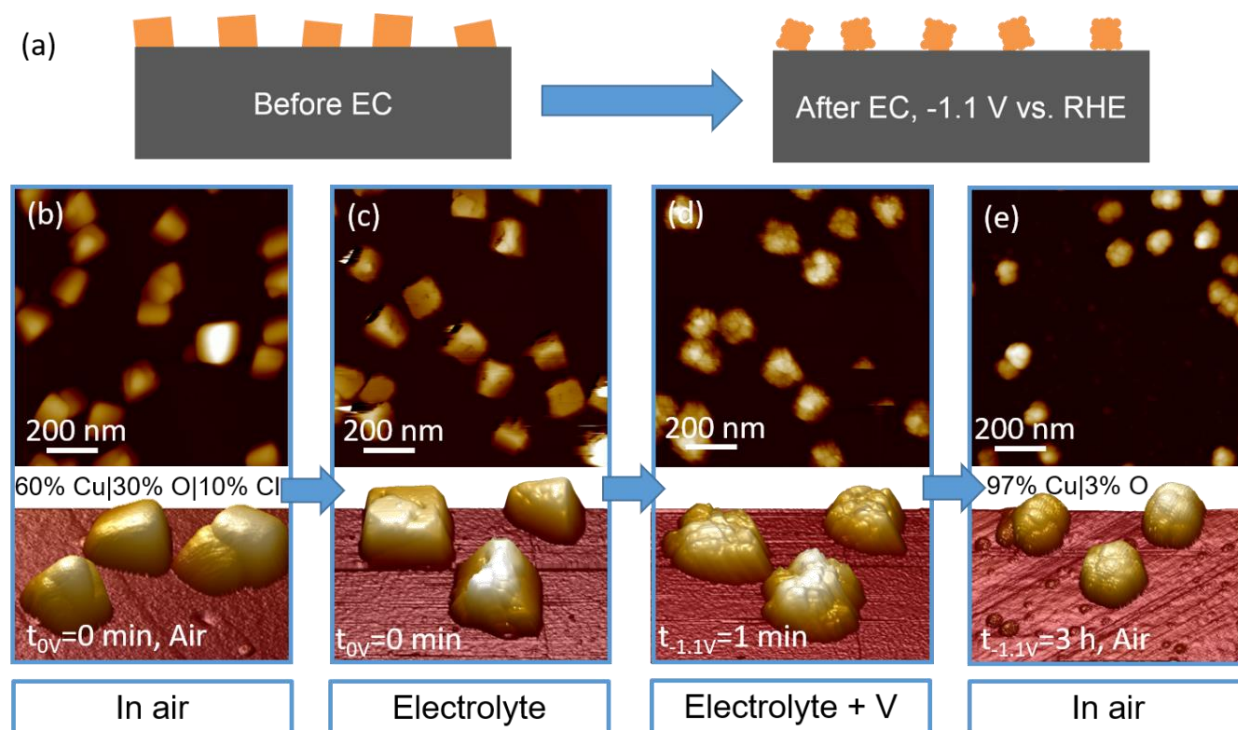


Figure 2: (a) Schematic representation of the morphological evolution of Cu-cube catalysts. (b-e) AFM images of Cu-cubes electrodeposited on highly oriented pyrolytic graphite (HOPG) acquired in air (b), and *operando* EC-AFM measurements in a CO_2 -saturated 0.1 M KHCO_3 aqueous solution at open circuit potential (c), at -1.1 V vs RHE in the same electrolyte for 1 min (d), and after 3 h under the same conditions as in (d) and subsequent air exposure (e).

A 1 min potential pulse at -1.1 V vs RHE resulted in the further roughening of the nanocube surface, loss of sharp corners and edges, and a decrease in the edge length of approximately 10 %. The latter is assigned to the initial reduction of Cu_2O to Cu as well as some loss of Cu. After 3h CO_2RR at -1.1 V vs RHE, an additional 10% decrease of the NP size was observed, with rough spherical-like NP shapes.

X-ray induced Cu LMM Auger electron spectra (XAES) were acquired from pristine C-supported Cu-cubes (220 nm) before and after CO_2RR for 1 h at -1.1 V vs RHE in an ultrahigh vacuum (UHV) XPS system directly interfaced to an electrochemical cell. Thus, the sample could be transferred in vacuum to avoid re-oxidation.^[21] Additional Cu-2p XPS data are also

included in Figure S3. A surface composition of the pristine C-supported Cu-cubes of 52% Cu₂O, 22% CuCl₂, 23% CuCl, and 3% metallic Cu was obtained before reaction, Fig. 3. After the electrochemical treatment, the cubes are mostly reduced, with only 4.6% Cu₂O detected. Interestingly, a higher content of Cu(I) species (13%) was detected after CO₂RR on similarly synthesized Cu-cubes supported on a Cu-foil, and no Cu(I) at all in the pristine Cu foil support after CO₂RR (Fig. S4). The latter finding highlights the key role of having a Cu cube/Cu foil interaction for the stabilization of Cu(I) species.

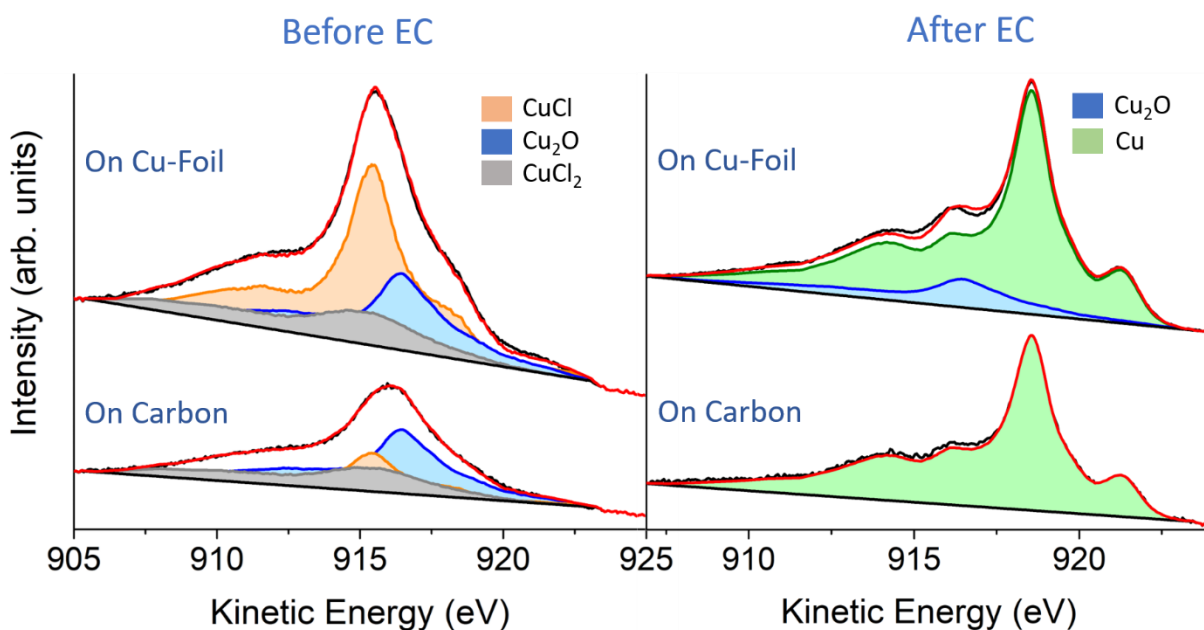


Figure 3: *Quasi in situ* Cu Auger LMM XAES spectra of 220 nm Cu-cubes electrodeposited on C paper and 250 nm cubes deposited on a Cu-foil acquired before (left) and after 1 hour of CO₂RR at -1.1 V vs RHE (right).

To gain further insight into the stability of Cl, Cu(I) species, and subsurface oxygen in our Cu-cubes under CO₂RR, *operando* X-ray absorption fine-structure (XAFS) measurements were conducted on 120 nm Cu-cubes, Figures 4 and 5 (see also Figs. S5, S6, and S7 in the SI for more details). As compared to XPS and XAES, probing about 10 nm below the surface, XAFS is a bulk sensitive technique that reveals the overall change in the structure and composition of the Cu-cubes. The as prepared state of the Cu-cubes shows a mixture of Cu₂O and CuCl_x, which almost immediately changes fully into Cu₂O upon becoming in contact with the electrolyte.

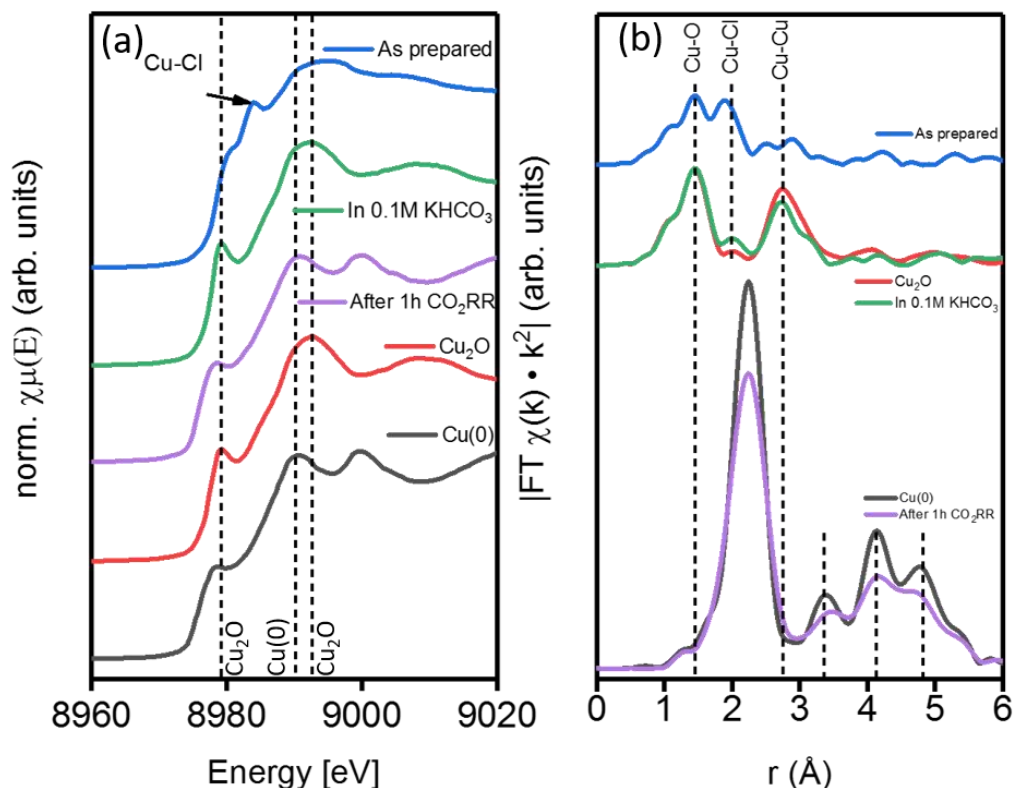


Figure 4: Normalized XANES region (a) and Fourier-transform magnitudes of k^2 -weighted EXAFS data of pristine Cu-cubes (b) measured as prepared, in 0.1 M KHCO_3 at open circuit potential and in the same electrolyte after 1 h of CO_2RR at -1.1V vs RHE.

In CO_2 -saturated 0.1 M KHCO_3 , the spectrum resembles greatly the Cu_2O reference, with only a very small difference where the Cl-feature was previously observed and a small decrease of the Cu-O feature. After 1 h at -1.1 V vs RHE, the spectrum obtained for the Cu-cubes resembles that of metallic Cu. The Fourier Transform (FT) data obtained for the Cu-cubes in the electrolyte after 1 h CO_2RR at -1.1 V vs RHE are less intense in comparison to a bulk Cu-foil, indicating a lower atomic coordination and/or enhanced disorder. From the first shell Cu-Cu fit after 1 h of applied potential, a coordination number (CN) of 10.8 was obtained for the Cu-cubes (vs 12 expected for bulk fcc-Cu). The lower CN values might be indicative of the formation and growth of pores on the Cu-cubes under reaction conditions or the partial disruption of the nanocubes, leading to an increased surface roughness. The *operando* EC-AFM data also revealed the formation of cracks (defects) on the Cu-cubes even before applying the potential once they were exposed to the electrolyte. The porous structure seen by SEM can also be a result of the

progressive CuO_x reduction during the reaction. Under reaction conditions, these pores as well as the cube corner and edges appear more prone to dissolution than the higher coordinated flat facets.

Since Cu(I) species^[37] have been suggested to play an important role in CO_2RR selectivity, in order to better detect small amounts of Cu-O species an additional analysis of the XAFS data by wavelet-transform^[38] (WT) was applied (Fig. 5). Details on this analysis are given in the Suppl. documents. After processing the measured spectra and doing the Morlet WT, the spectra were normalized to the highest signal, allowing subtraction of measured reference materials. The Cu-cubes on C in their as prepared state display a large feature at $R \sim 1.4 \text{ \AA}$ and $k \sim 3.9 \text{ \AA}^{-1}$ which represents the Cu-O shell, whereas at a slightly higher distance ($R \sim 2 \text{ \AA}$, $k \sim 7 \text{ \AA}^{-1}$) the interaction of Cu with the heavier Cl atoms is visible. For the as prepared sample, only a faint Cu-Cu 1st and 2nd neighbor interaction is observed after the subtraction of a Cu_2O reference from the pristine Cu-cubes, indicating that the sample consists almost exclusively of Cu_2O and CuCl_x . In the electrolyte, the CuCl_x feature immediately disappears and a Cu-O-Cu feature arises at $R \sim 3 \text{ \AA}$ and $k \sim 4 \text{ \AA}^{-1}$. In addition, the magnitude of the Cu-O component decreases slightly. After 1 hour of CO_2RR , the most prominent feature is that of the Cu-Cu first shell ($R \sim 2.3 \text{ \AA}$, $k \sim 7 \text{ \AA}^{-1}$). The pristine Cu-cubes resemble almost exactly the metallic Cu reference after 1 h of CO_2RR . A deconvolution of the Cu XANES region of pristine Cu-cubes in 0.1 M KHCO_3 with Cu, Cu_2O , and CuO standards, Fig. S7, yields a composition of 99% Cu and 1% (within the error margin) Cu_2O after 1 h of CO_2RR . CuO could not be reasonable fitted. These findings are in good accordance with the XPS surface analysis indicating that for the Cu cube/C system, no Cu_2O species remain after 1 h of reaction either at the surface (XPS) or in sub-surface regions (XAFS), which is in clear contrast with previous findings for Cu-cubes/Cu-foil (Fig. 3).^[21,22]

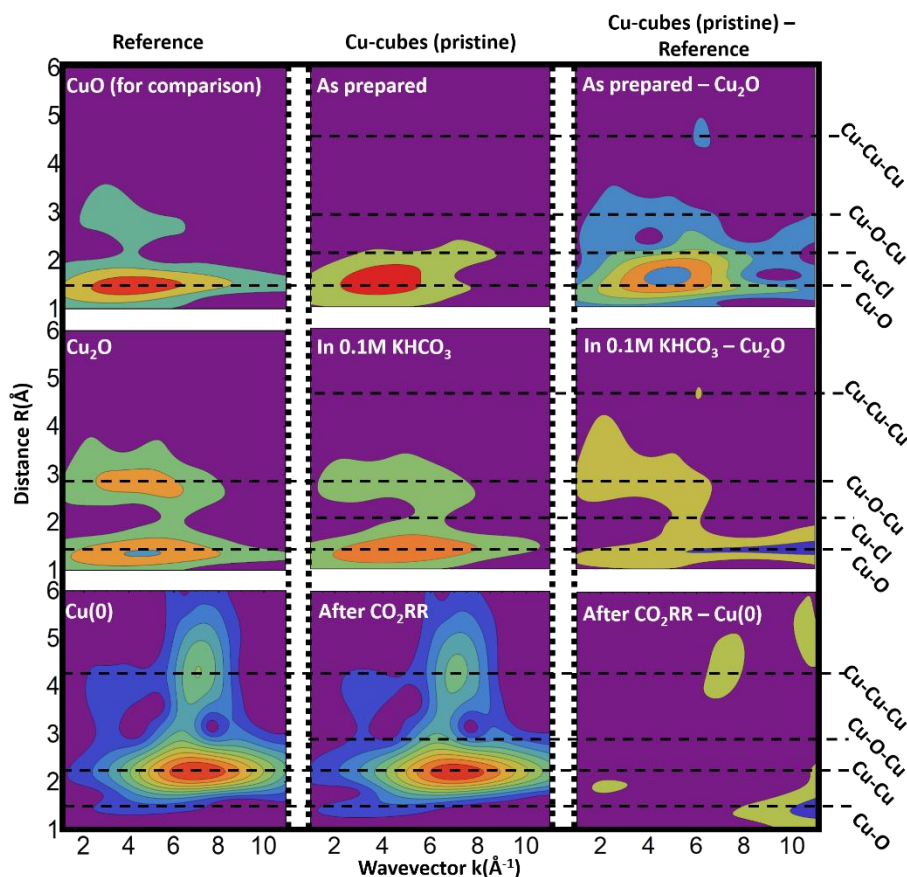


Figure 5: Wavelet transform of Cu-cubes as prepared, in 0.1 M KHCO_3 , and after 1 h of CO_2RR as well as the bulk references used. The height is normalized to 1 and the subtracted images scaled by a factor of two for better visibility.

After observing drastic changes in the morphology and chemical composition of the samples under *operando* reaction conditions within the first 1-3 hours, the electrochemical performance was studied in order to gain insight into structure/chemical-state/reactivity correlations. The background signal from the C-paper support was subtracted from all data presented as shown in Fig. S8. A comparison between HER Faradaic efficiency (parasitic reaction) and the sum of the Faradaic efficiency of all products from CO_2RR is displayed in Fig. 6. Detailed information on the different products detected can be found in Figure 7 for the 220 nm Cu-cubes and in Figs. S9, S10 and S11. Similar trends for small Cu-Au cubic NPs were reported in Ref.^[39].

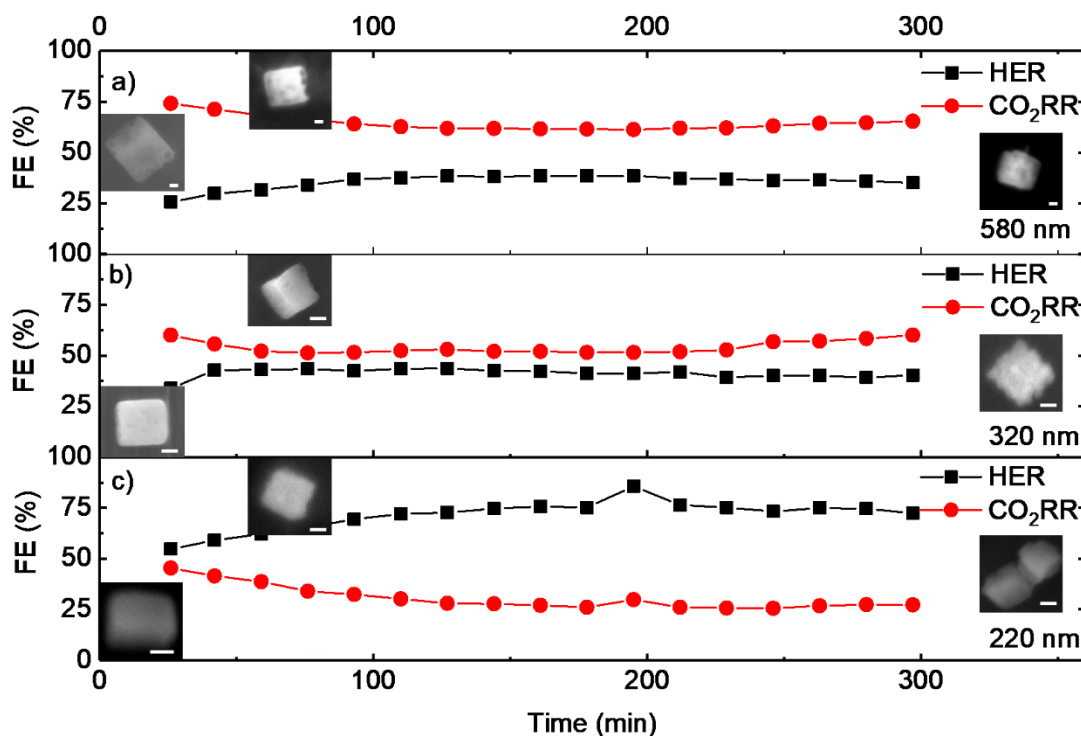


Figure 6: Faradaic efficiency for CO₂RR and HER at approx. -1.05 V vs RHE recorded during 5 h for samples with different Cu-cube sizes (580 nm, 320 nm and 220 nm). The insets display SEM images of typical Cu-cubes measured after different reaction times. The size of the scale bars is: 100 nm.

For the stability tests, Cu-cubes of three different initial sizes were investigated at -1.05 V vs RHE, Fig. 6. This potential was chosen since we expect to see the highest selectivity towards C₂-C₃ products.^[40] The total CO₂RR FE was found to increase with increasing Cu-cube size, and achieves an initial value of ~75% on the 580 nm cubes. This is assigned to the enhanced morphological stability of the larger cubes. A second general trend, independent of size, is a decrease in Faradaic efficiency for CO₂RR and increase in HER over time. An exception is the 220 nm Cu-cubes sample, which starts already with higher Faradaic efficiency for HER over CO₂RR. The smaller cube size on the weakly binding C support leads to a lower stability of the Cu-cubes, leading to the loss of material into the electrolyte and possible subsequent re-deposition as small clusters on the support and Cu-cubes. Such low-coordinated structures were found to result in an increase in the H₂ and CO production and decrease in the formation of hydrocarbons, i.e., they favor HER over CO₂RR.^[41,42] This correlates with the observed

morphological changes shown in Figs. 1 and 2, where a rounding of corners and edges and roughening of the facets is observed.

Figure 7(a) displays the FE data for the pristine 220 nm Cu-cubes on C measured as a function of the applied potential after 1 h of CO₂RR. The production of C₂ and C₃ products peaks around -1.0 V vs RHE, whilst hydrogen evolution is the lowest at this potential. At potentials more negative than -1.0 V vs RHE, C₂/C₃ products decline, whilst methane production keeps increasing.

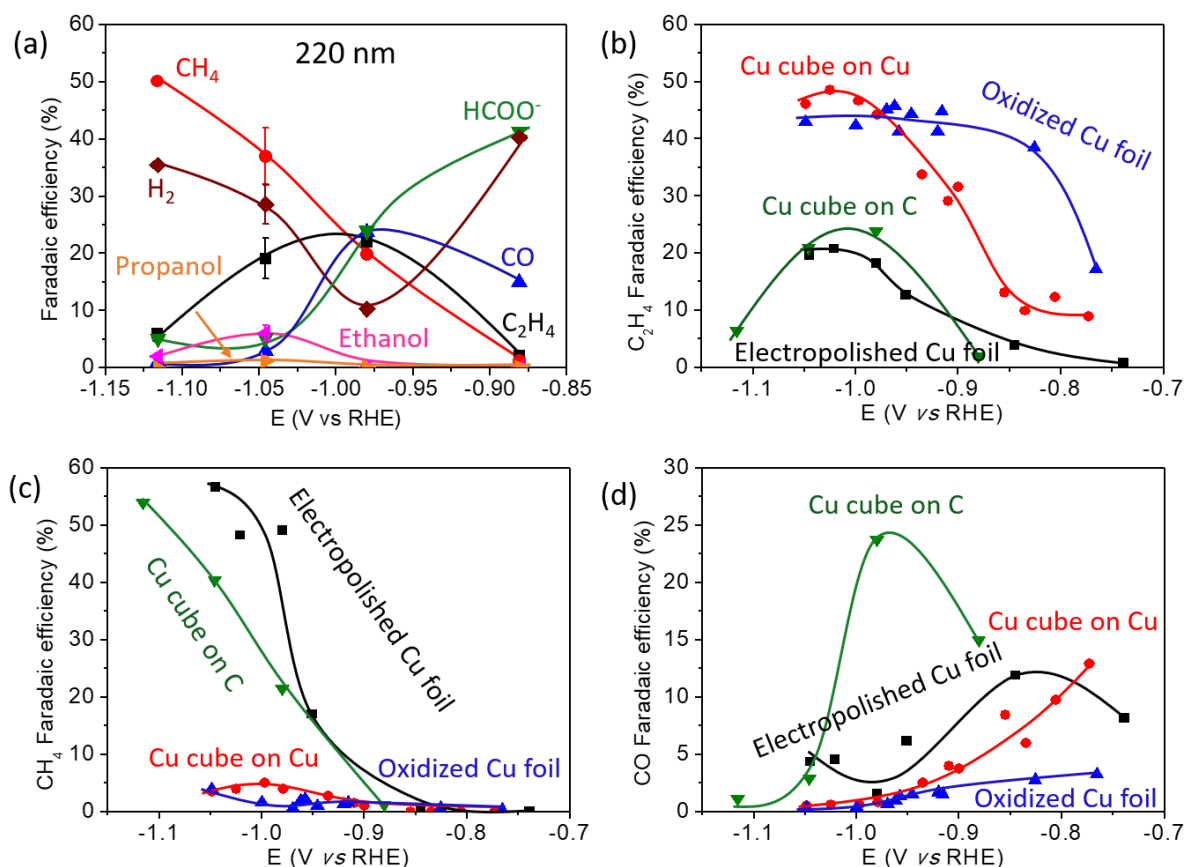


Figure 7: (a) Faradaic efficiency of pristine 220 nm Cu-cubes on C as a function of applied potential obtained after 1 h of CO₂RR. Faradaic efficiencies for C₂H₄ (b), CH₄ (c), and CO (d) are also shown for the pristine 220 nm Cu-cubes on C, 250 nm Cu-cubes on Cu foil and two reference Cu foils electropolished and O₂-plasma treated (20, 20 W, 400 mTorr^[21,22]).

The FE for C₂H₄, CH₄ and CO as a function of the applied potential for the pristine 220 nm Cu-cubes on C together with similarly synthesized 250 nm Cu-cubes supported on a Cu foil as

well as electropolished and O₂-plasma-treated Cu foils are shown in Figure 7(b-d). A very strong support-effect is evident, with the Cu-cubes grown in the weakly-interacting C support displaying a higher overpotential and much lower FE for C₂-C₃ products than the similarly-sized Cu-cubes on the Cu foil. In fact, as can be seen in Fig.7(c), the production of CH₄ of the Cu-cubes on C resembles the metallic electropolished Cu foil, although the CO production of the Cu-cubes on C is much higher, Fig. 7(d). It is remarkable that the pristine Cu-cubes on the Cu-foil, which were found to better stabilize Cu(I) species under reaction conditions (Fig. 3), behave similarly to the O₂-plasma treated Cu foils. Therefore, we can confirm a direct correlation between the stabilization of Cu(I)/Cu interfaces (Cu-cubes on Cu-foil) and the selectivity for C₂-C₃ products, since metallic Cu-cubes supported on C are more selective for C₁ products. This is however not the only reason for the change in the selectivity. Even more important are the drastic structural changes observed for the Cu-cubes on C. The roughening of the Cu(100) facets, partial loss of the cubic shape and formation of pores as well as detachment from the surface play a critical role in the selectivity switch reported here.

2. Conclusion

A simple electrochemical method for the synthesis of cubic-shaped nanoparticles of tunable size supported on carbon substrates is presented here. Dynamic morphological and chemical transformations of Cu-cubes during CO₂RR were monitored using *operando* EC-AFM and X-ray absorption fine-structure spectroscopy (XAFS). Drastic changes in the cube morphology were found to take place for the Cu-cubes on C under CO₂RR conditions, including the roughening and loss of (100) facets, loss of Cu atoms from edge and corner sites, and the reduction of CuO_x species.

The selectivity for CO₂RR versus HER was found to decrease with decreasing cube size, which was assigned to more drastic changes in the cube morphology taking place under CO₂RR conditions over smaller cubes. In contrast with the findings previously reported for Cu-cubes grown on Cu foils, a surprisingly high selectivity for CH₄ as compared to C₂H₄ was observed when deposited on C. The observed morphological instability of the Cu-cubes on C versus those deposited on Cu, together with the absence of stable Cu(I) species in the former samples are considered responsible for the preferred selectivity for C₁ products.

Experimental

A mixture of 5 mM copper sulfate-pentahydrate ($\text{CuSO}_4 \cdot 5\text{H}_2\text{O}$) and 5 mM potassium chloride (KCl) was used as starting material. Electrochemical cycling between an oxidizing (+0.55 V vs. RHE) and a reducing potential (+0.22 V vs. RHE) with varying number of cycles (1-100), depending on the desired cube size and coverage, lead to the electrodeposition of size- and shape-controlled Cu-cubes with a narrow size distribution. Initially, a potential of -0.2 V vs Ag/AgCl was held for 8 s with a subsequent ramp to +0.4 V vs Ag/AgCl for 4 s at a ramp rate of 700 mV/s. Returning to the initial potential completes the cyclic voltammetry. Cu-cube sizes ranging from 80 nm to 1,2 μm were obtained depending on the 1 mM-100 mM KCl and $\text{CuSO}_4 \cdot 5\text{H}_2\text{O}$ concentration, number of cycles, and applied potential. High surface area carbon paper was used as substrate (Toray Carbon Paper TGP-H-060). The Cu-cubes on Cu-foil were prepared as described in Ref. ^[21].

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3. References

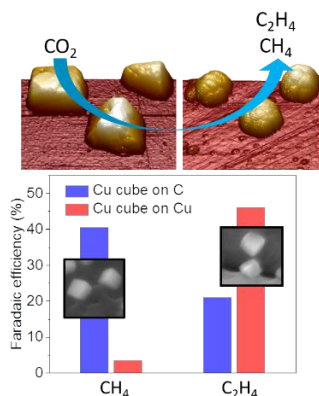
- [1] Y. Hori, A. Murata, R. Takahashi, S. Suzuki, *J. Chem. Soc. Chem. Commun.* **1988**, 109, 17.
- [2] H. Mistry, A. S. Varela, S. Kühn, P. Strasser, B. R. Cuenya, *Nat. Rev. Mater.* **2016**, 1, 16009.
- [3] R. Kortlever, J. Shen, K. J. P. Schouten, F. Calle-Vallejo, M. T. M. Koper, *J. Phys. Chem. Lett.* **2015**, 6, 4073–4082.
- [4] G. O. Larrazábal, A. J. Martín, J. Pérez-Ramírez, *J. Phys. Chem. Lett.* **2017**, 8, 3933–3944.
- [5] C. W. Li, M. W. Kanan, *J. Am. Chem. Soc.* **2012**, 134, 7231–7234.
- [6] A. M. Appel, *Nature* **2014**, 508, 460–461.
- [7] F. S. Roberts, K. P. Kuhl, A. Nilsson, *Angew. Chem. Int. Ed.* **2015**, 54, 5179–5182.
- [8] A. Loiudice, P. Lobaccaro, E. A. Kamali, T. Thao, B. H. Huang, J. W. Ager, R. Buonsanti, *Angew. Chem. Int. Ed.* **2016**, 55, 5789–5792.
- [9] K. J. P. Schouten, E. Pérez Gallent, M. T. M. Koper, *ACS Catal.* **2013**, 3, 1292–1295.
- [10] F. Calle-Vallejo, M. T. M. Koper, *Angew. Chem.* **2013**, 125, 7423–7426.
- [11] J. H. Montoya, C. Shi, K. Chan, J. K. Nørskov, *J. Phys. Chem. Lett.* **2015**, 6, 2032–2037.
- [12] D. Kim, C. S. Kley, Y. Li, P. Yang, *Proc. Natl. Acad. Sci.* **2017**, 114, 10560–10565.
- [13] R. Kas, R. Kortlever, H. Yilmaz, M. T. M. Koper, G. Mul, *ChemElectroChem* **2015**, 2, 354–358.
- [14] W. Tang, A. A. Peterson, A. S. Varela, Z. P. Jovanov, L. Bech, W. J. Durand, S. Dahl, J. K. Nørskov, I. Chorkendorff, *Phys. Chem. Chem. Phys.* **2012**, 14, 76–81.
- [15] R. B. Sandberg, J. H. Montoya, K. Chan, J. K. Nørskov, *Surf. Sci.* **2016**, 654, 56–62.
- [16] I. Zegkinoglou, A. Zendegani, I. Sinev, S. Kunze, H. Mistry, H. S. Jeon, J. Zhao, M. Y.

- Hu, E. E. Alp, S. Piontek, et al., *J. Am. Chem. Soc.* **2017**, *139*, 14360–14363.
- [17] R. Reske, H. Mistry, F. Behafarid, B. Roldan Cuenya, P. Strasser, *J. Am. Chem. Soc.* **2014**, *136*, 6978–6986.
- [18] H. Mistry, F. Behafarid, R. Reske, A. S. Varela, P. Strasser, B. Roldan Cuenya, *ACS Catal.* **2016**, *6*, 1075–1080.
- [19] H. Mistry, R. Reske, P. Strasser, B. Roldan Cuenya, *Catal. Today* **2017**, *288*, 30–36.
- [20] H. Mistry, A. S. Varela, C. S. Bonifacio, I. Zegkinoglou, I. Sinev, Y. W. Choi, K. Kisslinger, E. A. Stach, J. C. Yang, P. Strasser, et al., *Nat. Commun.* **2016**, *7*, 12123.
- [21] D. Gao, I. Zegkinoglou, N. J. Divins, F. Scholten, I. Sinev, P. Grosse, B. Roldan Cuenya, *ACS Nano* **2017**, *11*, 4825–4831.
- [22] D. Gao, F. Scholten, B. Roldan Cuenya, *ACS Catal.* **2017**, *7*, 5112–5120.
- [23] C. Reller, R. Krause, E. Volkova, B. Schmid, S. Neubauer, A. Rucki, M. Schuster, G. Schmid, *Adv. Energy Mater.* **2017**, *7*, 1602114.
- [24] A. Eilert, F. Cavalca, F. S. Roberts, J. Osterwalder, C. Liu, M. Favaro, E. J. Crumlin, H. Ogasawara, D. Friebel, L. G. M. Pettersson, et al., *J. Phys. Chem. Lett.* **2017**, *8*, 285–290.
- [25] S. Lee, D. Kim, J. Lee, *Angew. Chem. Int. Ed.* **2015**, *54*, 14701–14705.
- [26] D. Ren, Y. Deng, A. D. Handoko, C. S. Chen, S. Malkhandi, B. S. Yeo, *ACS Catal.* **2015**, *5*, 2814–2821.
- [27] H. Xiao, W. A. Goddard, T. Cheng, Y. Liu, *Proc. Natl. Acad. Sci.* **2017**, *114*, 6685–6688.
- [28] M. Favaro, H. Xiao, T. Cheng, W. A. Goddard, J. Yano, E. J. Crumlin, *Proc. Natl. Acad. Sci.* **2017**, *114*, 6706–6711.
- [29] X. Wang, A. S. Varela, A. Bergmann, S. Kühn, P. Strasser, *ChemSusChem* **2017**, *10*, 4642–4649.
- [30] Z. W. Ulissi, M. T. Tang, J. Xiao, X. Liu, D. A. Torelli, M. Karamad, K. Cummins, C.

- Hahn, N. S. Lewis, T. F. Jaramillo, et al., *ACS Catal.* **2017**, *7*, 6600–6608.
- [31] X. Liu, J. Xiao, H. Peng, X. Hong, K. Chan, J. K. Nørskov, *Nat. Commun.* **2017**, *8*, 15438.
- [32] Z. W. She, J. Kibsgaard, C. F. Dickens, I. Chorkendorff, J. K. Nørskov, T. F. Jaramillo, *Science* **2017**, *355*, eaad4998.
- [33] W. Ju, A. Bagger, G. P. Hao, A. S. Varela, I. Sinev, V. Bon, B. Roldan Cuenya, S. Kaskel, J. Rossmeisl, P. Strasser, *Nat. Commun.* **2017**, *8*, 944.
- [34] P. De Luna, R. Quintero-Bermudez, C.-T. Dinh, M. B. Ross, O. S. Bushuyev, P. Todorović, T. Regier, S. O. Kelley, P. Yang, E. H. Sargent, *Nat. Catal.* **2018**, *1*, 103–110.
- [35] K. Jiang, R. B. Sandberg, A. J. Akey, X. Liu, D. C. Bell, J. K. Nørskov, K. Chan, H. Wang, *Nat. Catal.* **2018**, *1*, 111–119.
- [36] B. Eren, R. S. Weatherup, N. Liakakos, G. A. Somorjai, M. Salmeron, *J. Am. Chem. Soc.* **2016**, *138*, 8207–8211.
- [37] C. S. Le Duff, M. J. Lawrence, P. Rodriguez, *Angew. Chem. Int. Ed.* **2017**, *56*, 12919–12924.
- [38] J. Timoshenko, A. Kuzmin, *Comput. Phys. Commun.* **2009**, *180*, 920–925.
- [39] J. Monzó, Y. Malewski, R. Kortlever, F. J. Vidal-Iglesias, J. Solla-Gullón, M. T. M. Koper, P. Rodriguez, *J. Mater. Chem. A* **2015**, *3*, 23690–23698.
- [40] Y. Hori, A. Murata, R. Takahashi, *J. Chem. Soc. Faraday Trans. 1 Phys. Chem. Condens. Phases* **1989**, *85*, 2309.
- [41] H. Mistry, R. Reske, Z. Zeng, Z.-J. Zhao, J. Greeley, P. Strasser, B. R. Cuenya, *J. Am. Chem. Soc.* **2014**, *136*, 16473–16476.
- [42] F. Behafarid, J. Matos, S. Hong, L. Zhang, T. S. Rahman, B. Roldan Cuenya, *ACS Nano* **2014**, *8*, 6671–6681.

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A variety of *In situ* and *operando* spectroscopic and microscopic methods were used to gain insight into the correlation between the structure, chemical state, and reactivity of size- and shape-controlled ligand-free Cu nanocubes during CO_2 electroreduction. Dynamic changes in the morphology and composition of Cu nanocubes supported on carbon were monitored under potential control and correlated to product selectivity.



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