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#### Article

# Controllable CO adsorption determines ethylene and methane productions from CO<sub>2</sub> electroreduction

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#### ABSTRACT

Among all CO<sub>2</sub> electroreduction products, methane (CH<sub>4</sub>) and ethylene (C<sub>2</sub>H<sub>4</sub>) are two typical and valuable hydrocarbon products which are formed in two different pathways: hydrogenation and dimerization reactions of the same CO intermediate. Theoretical studies show that the adsorption configurations of CO intermediate determine the reaction pathways towards  $CH_4/C_2H_4$ . However, it is challenging to experimentally control the CO adsorption configurations at the catalyst surface, and thus the hydrocarbon selectivity is still limited. Herein, we seek to synthesize two well-defined copper nanocatalysts with controllable surface structures. The two model catalysts exhibit a high hydrocarbon selectivity toward either  $CH_4$  (83%) or  $C_2H_4$  (93%) under identical reduction conditions. Scanning transmission electron microscopy and X-ray absorption spectroscopy characterizations reveal the low-coordination  $Cu^0$  sites and local  $Cu^0/Cu^+$  sites of the two catalysts, respectively. CO-temperature programed desorption, *in-situ* attenuated total reflection Fourier transform infrared spectroscopy and density functional theory studies unveil that the bridge-adsorbed CO (CO<sub>B</sub>) on the low-coordination  $Cu^0$  sites is apt to be hydrogenated to CH<sub>4</sub>, whereas the bridge-adsorbed CO plus linear-adsorbed CO ( $CO_B + CO_L$ ) on the local  $Cu^0/Cu^+$  sites are apt to be coupled to  $C_2H_4$ . Our findings pave a new way to design catalysts with controllable CO adsorption configurations for high hydrocarbon product selectivity.

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#### 1. Introduction

 $CO_2$  electroreduction ( $CO_2RR$ ), powdered by renewable electricity, provides a promising way to reduce greenhouse gas net emission, store the intermittent renewable clean energy and generate industrial feedstock [1,2]. Up to now, only copper (Cu) catalysts can give a rich gamut of single-carbon and multi-carbon products [3,4]. However, the reduction pathways often furcate to form a wide range of products, which are affected by both the chemical properties of catalysts and reduction reaction conditions [5].

Among all reduction products, only a few primary products like CO and HCOOH can be achieved with a high selectivity (i.e., Faradic efficiency (FE)) of >95% [6,7]. The production of either CO or HCOOH is related to a two-electron process, typically starting from

\* Corresponding authors. *E-mail addresses:* bozhang@fudan.edu.cn (B. Zhang), penghs@fudan.edu.cn (H. Peng). relative fixed CO<sub>2</sub> adsorption configurations [8]. For instance, the C-bonded CO<sub>2</sub> adsorption configurations on Au, Ag, Pd surfaces [9–11] undergo a proton-coupled electron transfer process to \*COOH for CO formation, whereas O-bonded CO<sub>2</sub> adsorptions on Bi, Sn and Pb surfaces [6,12] transform to \*OCHO for HCOOH production. The simple production pathways with controlled CO<sub>2</sub> adsorption configurations lead to the high selectivity of CO or HCOOH.

In contrast, the hydrocarbon products are derived from the \*CO intermediate and subsequent more complicated reaction pathways [9,13], but \*CO absorption configurations on the Cu catalysts are not well controllable, so the hydrocarbon product selectivity is not as high as the selectivity of CO or HCOOH. The reduction of \*CO intermediate mainly results in C<sub>1</sub> productions such as CH<sub>4</sub> via a hydrogenation pathway and C<sub>2</sub> productions such as C<sub>2</sub>H<sub>4</sub> via a dimerization pathway [14]. Theoretical studies have shown that different \*CO configurations and adsorption energies on the

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**Fig. 1.** (Color online) Morphology characterizations. (a), (b) TEM images of the MP-Cu and EP-Cu catalysts. Insets show a uniform particle size of ~15 nm for both catalysts. (c), (d) STEM images of a representative catalyst particle of EP-Cu and MP-Cu, revealing the  $Cu^0$  phase with abundant step sites in MP-Cu while abundant local  $Cu^0/Cu^+$  sites in EP-Cu.

Cu catalysts [15,16] can result in different energy barriers of CO hydrogenation or dimerization reactions [17–19].

The CO adsorption configuration and adsorption energy rely on a number of factors, such as the surface structures of catalysts which can be affected by composition [20], particle size [21], grain boundary [22], facet [23] and so on, and reduction reaction conditions of applied potentials [24], electrolyte species [25], local pH [26], electrolyte concentration [27], etc. For instance, a lowcoordination Cu is potential to facilitate the translation of \*CO to \*CHO, and thus facilitate the C<sub>1</sub> pathway [23,28]. By contrast, the coupling reaction of \*CO is highly related to the local valence structure of Cu in which the local Cu<sup>0</sup>/Cu<sup>+</sup> site is reported to promote the dimerization and therefore enhance the C<sub>2</sub> production [20,29].

It is still challenging to control the variate that affects the \*CO absorption configurations on Cu experimentally, thus it remains difficult to modulate the reaction pathways and final products [21,30]. Especially, the formations of CH<sub>4</sub> and C<sub>2</sub>H<sub>4</sub> are always concomitant on the Cu catalysts when the reaction parameters are similar, which makes it even more difficult to distinguish the contributions of \*CO adsorption properties. It is therefore necessary to obtain well-defined model catalysts with controllable \*CO absorption configurations that can catalyze CO<sub>2</sub> towards highly selective CH<sub>4</sub> and C<sub>2</sub>H<sub>4</sub> productions in identical reduction conditions.

Herein, using a controlled variation method, we develop two catalysts with controlled and distinct active sites. The low-coordination Cu catalysts show a high  $CH_4$  selectivity of 83% while the Cu/Cu<sub>2</sub>O catalysts exhibit a high  $C_2H_4$  selectivity of 93% under identical reduction conditions. We study the CO adsorption properties and establish a correlation between the CO adsorption and reduction pathways on these two model catalysts, with the aid of CO temperature-programed desorption (TPD), *in-situ* attenuated total reflection Fourier transform infrared spectroscopy

(ATR-FTIR) and density functional theory calculations. These results show that the bridge-adsorbed CO (CO<sub>B</sub>) on the low-coordination Cu<sup>0</sup> site is apt to be hydrogenated to form CH<sub>4</sub>, whereas the bridge-adsorbed CO plus linear-adsorbed CO (CO<sub>B</sub> + CO<sub>L</sub>) on the local Cu<sup>0</sup>/Cu<sup>+</sup> site is apt to be coupled to form C<sub>2</sub>H<sub>4</sub>.

#### 2. Experimental

The two catalysts were prepared via an epoxide-assisted hydrolysis method and were activated during CO<sub>2</sub>RR. The chemical structures of the catalysts were studied by scanning transmission electron microscopy (STEM) and X-ray absorption spectroscopy (XAS). The CO adsorption properties were studied by TPD and ATR-FTIR. The details of the preparation and characterization can be found in Supplementary materials (online).

#### 3. Results and discussion

#### 3.1. Structures characterizations

Inspired by the chemical structure modulating capability of halogens on the Cu catalyst [31,32], we sought to introduce halogen modifiers into the catalyst precursors to construct different Cu catalysts with either low-coordination or local Cu<sup>0</sup>/Cu<sup>+</sup> sites. Experimentally, we successfully synthesized two Cu catalysts derived from bromine-doped (Figs. S1–S3 online) and iodine-doped copper oxychlorides (Figs. S4–S6 online), respectively [7,33] (see Supplementary materials (online) for details): one contains a low-coordination Cu<sup>0</sup> phase for potentially high CH<sub>4</sub> selectivity, denoted as methane production copper (MP-Cu); the other possesses Cu<sup>0</sup>/Cu<sup>+</sup> sites for potentially high C<sub>2</sub>H<sub>4</sub> selectivity,

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**Fig. 2.** (Color online) XAS characterizations of the MP-Cu and EP-Cu. (a) Cu K-edge XANES spectra of MP-Cu versus time, showing the valence states of Cu from initial 2+ to the stabilized 0. (b) Linear combination fit results of MP-Cu XANES at the Cu K-edge acquired in 90 min using Cu, Cu<sub>2</sub>O and CuO NPs as standards (Fitting range: 8970–9030 eV). (c) Cu K-edge XANES spectra of EP-Cu versus time, showing the valence states of Cu from initial 2+ to the stabilized positively charged Cu. (d) Linear combination fit results of EP-Cu XANES at the Cu K-edge acquired in 90 min using Cu, Cu<sub>2</sub>O and CuO NPs as standards (Fitting range: 8970–9030 eV).

denoted as ethylene production copper (EP-Cu). The two catalysts were activated using *in-situ* electrochemical reduction processes. No halogen residual was presented in the activated catalysts as shown in the X-ray photoelectron spectroscopy (XPS) and energy dispersive spectroscopy (EDS) results (Figs. S7 and S8 online).

To study the structures of the catalysts, we firstly carried out morphology characterizations. Fig. 1a and b represent the transmission electron microscopy (TEM) images of the two catalysts. Both MP-Cu and EP-Cu are nano-sized particles with ~15 nm diameter (Figs. S9 and S10 online), from which we excluded the influences of catalyst particle size on the reduction performance. High-resolution STEM images show distinct crystal structures of the two catalysts. The MP-Cu shows only one Cu<sup>0</sup> phase with surface step sites (Fig. 2c) while EP-Cu shows two phases containing both Cu<sup>0</sup> and Cu<sup>+</sup>, indicating the abundant local Cu/Cu<sup>+</sup> sites (Fig. 2d). Taken the TEM results together, we preliminarily conclude that the two catalysts were synthesized as designed.

To study the electronic structures of the two catalysts, we performed the Cu K-edge XAS measurements (see Supplementary materials for details (online)). By comparing the absorption thresholds of the X-ray absorption near edge structure (XANES), we extracted the valence states of the catalysts [33,34] (Fig. S11 online). The absorption threshold of MP-Cu shifts from Cu<sup>2+</sup> (8985.1 eV) to Cu<sup>0</sup> (8979.0 eV) and becomes stable (Fig. 2a) after a 30-min reduction process, indicating MP-Cu almost contains only Cu<sup>0</sup> species. The linear combination fit results of XANES of MP-Cu confirm the metallic Cu nature during CO<sub>2</sub>RR (Fig. 2b, see Supporting materials (online) for details). Extended X-ray absorption fine structure (EXAFS) of MP-Cu at the Cu K-edge reveals that the sample contains almost no Cu–O path but dominant Cu-Cu path (R = 2.54 Å). MP-Cu shows a low Cu-Cu coordination number of ~7, which is much lower than that of Cu foil (with a Cu-Cu coordination number of ~12), indicating a potential of MP-Cu towards highly selective CH<sub>4</sub> production from CO<sub>2</sub>RR (Fig. S12a, Table S1 online) [21,28]. For EP-Cu, the absorption threshold is stabilized at 8979.4 eV, revealing an intermediate valence between Cu<sup>+</sup> (8980.3 eV) and Cu<sup>0</sup> (Fig. 2c). The linear combination fit results of XANES of EP-Cu at the Cu K-edge suggest that EP-Cu contains 47% of Cu<sup>0</sup> and 53% of Cu<sup>+</sup> (Fig. 2d). The Cu K-edge EXAFS of EP-Cu shows an intense Cu–O path (R = 1.85 Å) similar to that of  $Cu_2O$  [35] (Fig. S12b online), which further confirms the  $Cu/Cu^+$ structure that is supposed to be active in C<sub>2</sub>H<sub>4</sub> production [20,29]. Both morphology and XAS studies show that the two catalysts have similar particle size but controllable surface chemical structures, and therefore can work as model catalysts for hydrocarbon production.

#### 3.2. CO<sub>2</sub> electroreduction performance

We then evaluated the CO<sub>2</sub> reduction performances of the two catalysts. CO<sub>2</sub>RR tests were carried out using 0.05 mol L<sup>-1</sup> KHCO<sub>3</sub> aqueous solution as electrolytes. All reduction conditions were controlled to eliminate the undesired influences of experimental variables (Table S2 online). Fig. 3 represents the CO<sub>2</sub>RR results of the MP-Cu and EP-Cu catalysts. The two catalysts show distinct selectivities towards CH<sub>4</sub> and C<sub>2</sub>H<sub>4</sub>, respectively. The MP-Cu shows 59% ± 1% CH<sub>4</sub> FE at -1.71 V versus reversible hydrogen electrode (vs. RHE) (Figs. 3a and S13a online). The corresponding carbon selectivity of CH<sub>4</sub> comparing to all carbon-containing products reaches up to 83% (Figs. 3b and S13b online). In contrast, the

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**Fig. 3.** (Color online) CO<sub>2</sub>RR performance of MP-Cu and EP-Cu. (a) FE distribution of reduction products on MP-Cu and (b) carbon selectivity distribution comparing to all carbon containing products of MP-Cu at optimized potential. (c) FE distribution of reduction products on MP-Cu and (d) carbon selectivity distribution comparing to all carbon containing products of MP-Cu at optimized potential. (e) FE distribution of reduction products on MP-Cu versus applied potential. (f) FE distribution of reduction products on EP-Cu versus applied potential.

commercial Cu without the low-coordination site show a much lower CH<sub>4</sub> selectivity under the same CO<sub>2</sub> reduction conditions, suggesting the intrinsic high CH<sub>4</sub> selectivity of MP-Cu (Fig. S14 online). At the same applied potential, the FE and carbon selectivity of  $C_2H_4$  produced on EP-Cu are 70% ± 1% and 93%, respectively (Fig. 3c and d). Furthermore, Figs. 3e, f and S15, S16 (online) represent the CH<sub>4</sub> and C<sub>2</sub>H<sub>4</sub> FEs of MP-Cu and EP-Cu at various applied potentials. Under all applied potentials, MP-Cu shows a high CH<sub>4</sub> selectivity while EP-Cu shows a high C<sub>2</sub>H<sub>4</sub> selectivity. Thus, we conclude that the MP-Cu and EP-Cu catalysts have intrinsic high selectivities towards CH<sub>4</sub> and C<sub>2</sub>H<sub>4</sub>, respectively, compared to previously reported catalysts (Fig. S17 online). The two catalysts also present robust stabilities without an obvious FE decrease after running CO<sub>2</sub>RR for 9000 s (Fig. S18 online). The local pH is a parameter that may cause differences in selectivity and it mainly depends on the electrode geometry, electrolyte, and current density [26,36]. In this study, the electrode geometry and electrolyte are identical in the two reduction processes. As a result, the difference in local pH mainly comes from the applied current density. To study the influence of local pH, we further measured the cyclic voltammetry curves of the two catalysts (Figs. S19-S21 online). The results show similar current densities of the two catalysts under the same applied potential, excluding the contribution of local pH on the reduction pathways in this study.

#### 3.3. CO adsorption study

To bridge the catalytic structures and  $CO_2$  reduction performance, the CO adsorption properties of the two catalysts were studied. Generally, the adsorbed CO differs in adsorption configurations (linear, bridge, and multi adsorption) and in adsorption energy [37,38]. The adsorption energy of bridge-adsorbed CO (CO<sub>B</sub>) is stronger than that of linear-adsorbed CO (CO<sub>L</sub>) but weaker than that of multi-adsorbed CO (CO<sub>M</sub>). We firstly used CO-TPD to evaluate the adsorption strength of CO on different catalysts. As shown in the TCD results, at a temperature range lower than 550 °C, MP-Cu shows only one obvious desorption peak at 481 °C that can be attributed to CO<sub>B</sub>, while EP-Cu shows two desorption peaks at 453 and 218 °C that can be attributed to CO<sub>B</sub> and CO<sub>L</sub>, respectively (Fig. 4a) [39,40]. The peaks at temperature exceeding 550 °C can be ascribed to CO<sub>M</sub> that is firmly bonded on the catalyst (Fig. S22 online) and presumably inactive during the reduction process [41,42].

Furthermore, surface sensitive *in-situ* ATR-FTIR spectroscopy was applied to study the surface adsorbed CO under different applied potentials. It is apparent that both MP-Cu and EP-Cu show an adsorption peak at ~1900 cm<sup>-1</sup> in associated with CO<sub>B</sub>, while EP-Cu shows an additional adsorption peak at ~2080 cm<sup>-1</sup> attributable to the weakly adsorbed CO<sub>L</sub> (Fig. 4b and c) [38,42,43]. From the CO adsorption studies, both CO-TPD and *in-situ* ATR-FTIR results suggest the consistent conclusion that there is only one CO adsorption configuration on MP-Cu (CO<sub>B</sub>) while there are two CO adsorption configurations on EP-Cu (both CO<sub>B</sub> and CO<sub>L</sub>).

To further study the relationship between the Cu oxidation state and CO adsorption, we further carried out CO-TPD measurements of commercial Cu and Cu<sub>2</sub>O NPs as controls. As shown in Fig. S23 (online), each sample shows only one dominate desorption peak, i.e., 210  $^{\circ}$ C (CO<sub>B</sub>) for Cu<sub>2</sub>O NPs and 430  $^{\circ}$ C (CO<sub>L</sub>) for Cu NPs before 550  $^{\circ}$ C (see Supplementary materials for details online). It can be seen from the CO-TPD results that the CO<sub>B</sub> adsorption energy of MP-Cu is stronger than that of Cu NPs, which can be ascribed to its lower Cu-Cu coordination numbers. Taken together

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**Fig. 4.** (Color online) CO adsorption studies of EP-Cu and MP-Cu. (a) CO TPD results of EP-Cu, MP-Cu and the catalysts support carbon paper (CP), showing only  $CO_B$  on MP-Cu while both  $CO_L$  and  $CO_B$  on EP-Cu. (b) *In-situ* ATR-FTIR study of MP-Cu versus applied potential, representing only  $CO_B$  on MP-Cu. (c) *In-situ* ATR-FTIR study of EP-Cu versus applied potential, revealing both  $CO_L$  and  $CO_B$  on EP-Cu. (d) Optimized CO adsorption configuration and relative vibration frequency on step site (bridge, left side) and on oxidized site (linear, right side) after removing the beneath Cu to show the sub-surface oxygen from DFT results (side views). (e) Illustration of different CO adsorption configurations towards either hydrogenation or dimerization reaction pathways.

with previous structure results, it can be concluded that MP-Cu containing only a low-coordination  $Cu^0$  phase shows only  $CO_B$  in accordance with commercial Cu NPs, and EP-Cu containing both  $Cu^0$  phase and  $Cu^+$  phases possess both  $CO_B$  and  $CO_L$  and the peak positions are adjacent to relative desorption peaks of Cu NPs and  $Cu_2O$  NPs. Thus, we experimentally conclude that CO is apt to adsorb on the  $Cu^0$  sites with a strong bridge configuration while on the  $Cu^+$  sites with a weak linear configuration.

To further understand our experimental results, we carried out DFT calculations with generalized gradient approximations (GGA) of Perdew-Burke-Ernzerhof (PBE) functional to investigate the CO adsorption property and possible active sites on MP-Cu and EP-Cu. As indicated from the CO adsorption results, only strong  $CO_B$  binding sites exist on MP-Cu. The strong binding sites, other than rhombi sites or square sites from basal planes, are likely step sites (low-coordination sites) as shown in Figs. 1c, 4d and S24a (online). We, therefore, carried out DFT calculations to verify the vibration frequency of CO on this step site, which leads to a prediction of 1891 cm<sup>-1</sup>, very close to the experimental observations of 1898 to 1911 cm<sup>-1</sup>. In our previous work, we determined that this step site can greatly reduce the formation energy of \*CHO [44] from the

reduction of \*CO intermediate, the first step of CO hydrogenation, which is known as both the potential determining step or rate determining step in the pathway of  $CH_4$  formation. Thus, the presence of step site can greatly catalyze the formation of  $CH_4$ , which explains the highly selective  $CH_4$  production as observed experimentally.

In our previous work, we found that partially reduced Cu<sub>2</sub>O generates Cu sites carrying partial charges (Cu<sup> $\delta^+$ </sup>), which can provide a fairly strong linear CO binding site for the further CO coupling reaction [45]. The atomic structure of one of the representative sites is shown in Figs. 4d and S24b (online). From DFT calculations, we found the C–O stretching of CO on this site is 2064 cm<sup>-1</sup>, very close to the experimental observations of 2086 to 2090 cm<sup>-1</sup>. We, therefore, conclude that the second ATR-FTIR peak appearing in EP-Cu attributes to the linear-adsorbed CO on the Cu<sup>+</sup> site, which plays an important role in C–C coupling as we demonstrated in our previous work: Cu<sup>+</sup> interacts weakly with CO, providing a suitable CO binding for CO coupling, which is beneficial for CO coupling to form C<sub>2</sub> products when combines with another strong CO binding site [46]. This also well explains the experimentally observed high selectivity of C<sub>2</sub>H<sub>4</sub> on EP-Cu.

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#### 4. Conclusions

In summary, under identical reduction conditions, a high CH<sub>4</sub> selectivity was achieved on MP-Cu consist of a low-coordination Cu<sup>0</sup> phase, and a high C<sub>2</sub>H<sub>4</sub> selectivity was realized using EP-Cu containing abundant Cu<sup>0</sup>/Cu<sup>+</sup> sites. The CO adsorption studies showed that the strongly adsorbed CO<sub>B</sub> on the low-coordination Cu<sup>0</sup> sites was apt to be hydrogenated to produce CH<sub>4</sub>, and the co-existence of CO<sub>B</sub> on the Cu<sup>0</sup> site and CO<sub>L</sub> on the Cu<sup>+</sup> site were apt to be coupled to produce C<sub>2</sub>H<sub>4</sub> (Fig. 4e). The DFT results were highly consistent with experimental results to further verify our conclusions. Our results expose the crucial factors that cause variation in reduction pathways for hydrocarbon productions and may guide future design of catalysts for both CO<sub>2</sub> and CO reductions.

#### **Conflict of interest**

The authors declare that they have no conflict of interest.

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#### **Author contributions**

Huisheng Peng and Bo Zhang supervised the project. Haipeng Bai, Shangyu Li and Zhenyu Zhou carried out all the experiments, characterization and  $CO_2$  reduction measurements. Tao Cheng guided the DFT part and Miao Xie carried out the detailed calculation. Yujin Ji and Youyong Li also discussed the DFT calculation part. Zhiyou Zhou and Shigang Sun carried out CO adsorption studies. All authors discussed the results and assisted for the manuscript preparation.

#### **Appendix A. Supplementary materials**

Supplementary materials to this article can be found online at https://doi.org/10.1016/j.scib.2020.06.023.

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