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## Stabilizing Cu<sup>0</sup>–Cu<sup>+</sup> sites by Pb-doping for highly efficient CO<sub>2</sub> electroreduction to C<sub>2</sub> products†

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The electrochemical CO<sub>2</sub> reduction reaction (CO<sub>2</sub>RR) can convert CO<sub>2</sub> to C<sub>2</sub> hydrocarbons and oxygenate over Cu-based catalysts, and has great potential to store renewable energy and close the carbon cycle. Developing a facile method to modify the local electronic structure of Cu is a useful way to design efficient catalysts. Herein, we design a Pb-doping Cu<sub>2</sub>O catalyst with controllable Cu<sup>0</sup>–Cu<sup>+</sup> sites. The catalyst generated a high C<sub>2</sub> faradaic efficiency (FE) of 83.9% with a current density of 203.8 mA cm<sup>-2</sup> at –1.1 V vs. RHE in a flow cell. *In situ* X-ray absorption spectroscopy and Raman spectroscopy revealed that the Pb doping in Cu<sub>2</sub>O could stabilize the Cu<sup>0</sup>–Cu<sup>+</sup> structure and enhance the CO adsorption and C–C coupling, leading to high activity for C<sub>2</sub> product formation. Theoretical calculations also show that Pb doping could reduce the energy barrier for both CO<sub>2</sub> activation and C–C coupling processes.

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### Introduction

The electrochemical CO<sub>2</sub> reduction reaction (CO<sub>2</sub>RR) can convert CO<sub>2</sub> into valuable fuels or chemicals, which have great potential to mitigate the greenhouse effect, alleviate the energy crisis, and store intermittent electrical energy.<sup>1–4</sup> C<sub>2</sub> products including ethylene, ethanol, and acetic acid have higher energy density and economic value per unit mass compared with their C<sub>1</sub> counterparts.<sup>5–9</sup> Current research has paid increasing attention to the enhancement of conversion and selectivity for CO<sub>2</sub>-to-C<sub>2</sub> products. Cu-based catalysts have been widely studied owing to their excellent performance in the electrochemical CO<sub>2</sub>RR to C<sub>2+</sub> products.<sup>10–15</sup> Previous studies show that the Cu<sup>0</sup>–Cu<sup>+</sup> sites in Cu-based catalysts are the key active sites for producing the C<sub>2+</sub> products, which are favorable for CO<sub>2</sub> activation and C–C coupling.<sup>16–20</sup> Therefore, it is of interest to modify the local electronic structure of Cu to achieve high activity and selectivity for CO<sub>2</sub>-to-C<sub>2</sub> products.

Unfortunately, the surface Cu<sup>+</sup> is usually unstable, and it is prone to be reduced to Cu<sup>0</sup> at the high applied reducing potentials during the CO<sub>2</sub>RR, leading to the loss of the performance of the CO<sub>2</sub>RR to give C<sub>2</sub> products.<sup>16, 21–25</sup> Hence, it is extremely important to stabilize Cu<sup>+</sup> in the catalysts.<sup>26,27</sup> Some strategies such as surface modification,<sup>17</sup> electropolishing,<sup>28</sup> and doping<sup>29</sup> can stabilize the surface Cu<sup>+</sup> species. Among them, introducing a p-block metal dopant into Cu has been shown to be a promising strategy. It has been reported that the introduction of a p-block metal can effectively stabilize Cu<sup>+</sup> in Cu-based catalysts.<sup>30–33</sup> Not only that, some p-block metal elements such as In, Sn, Bi, and Pb have O affinity and can reduce the reaction energy barrier of the CO<sub>2</sub>RR, as well as show high overpotentials for the H<sub>2</sub> evolution reaction (HER).<sup>34–37</sup> For example, Xie *et al.* screened Cu-based bimetallic catalysts for the CO<sub>2</sub>RR to form C<sub>2+</sub> products and found that Sn, In, Pb, and Bi in Cu-based bimetallic catalysts exhibited performances for inhibiting the HER.<sup>35</sup> Bai *et al.* used first-principles calculations to find that Sn doping into Cu-based catalysts can effectively suppress the HER without altering the activity toward CO<sub>2</sub> reduction, thereby improving the FE of ethanol.<sup>36</sup> Li *et al.* found that introducing Sn can enhance the surface oxophilicity of Cu–Sn alloy catalysts, which plays an important role in guiding the protonation of the key oxygenic intermediate and transforming CO<sub>2</sub> into ethanol.<sup>34</sup> Wang *et al.* developed Cu–Bi bimetallic aerogels as catalysts, which successfully improved the faradaic efficiency (FE) of ethylene in the CO<sub>2</sub>RR.<sup>37</sup> Therefore, developing facile methods to construct p-block metal doped Cu catalysts with controllable Cu<sup>0</sup>–Cu<sup>+</sup> sites is an effective way to realize the CO<sub>2</sub>-to-C<sub>2</sub> products with high efficiency.

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In this work, we have designed a Pb-doped  $\text{Cu}_2\text{O}$  electrocatalyst for highly efficient  $\text{CO}_2$  reduction to  $\text{C}_2$  products. It was discovered that the FE of  $\text{C}_2$  products could reach 83.9% with good stability. Experimental and density functional theory (DFT) studies indicated that the doping of Pb in the catalyst can stabilize the  $\text{Cu}^0\text{-Cu}^+$  structure, which could reduce the energy barrier of  $\text{CO}_2$  activation and C-C coupling. As a result, it can improve the performance of the  $\text{CO}_2\text{RR}$  to form  $\text{C}_2$  products.

## Results and discussion

The detailed fabrication procedures of the electrocatalysts are discussed in the Methods section. In brief, as shown in Fig. 1A, 1 mmol of copper sulfate, 5 mL of ethanol, and 2 mL of oleic acid were added into 15 mL of deionized water and heated to 80 °C. Then, 5 mL of 1 M NaOH solution and 5 mL of 2 M glucose solution were added into the flask. The obtained reddish-brown precipitate was  $\text{Cu}_2\text{O}$ .<sup>38</sup> Subsequently, the  $\text{Cu}_2\text{O}$  powder was evenly dispersed in the lead acetate solution and stirred for a long time without light and air. The mass fraction of Pb doped into  $\text{Cu}_2\text{O}$  can be controlled with different stirring times. After stirring, we can obtain a series of  $\text{Pb/Cu}_2\text{O-x}$  catalysts, where  $x$  is the mass fraction of Pb in the

catalysts measured by inductively coupled plasma optical emission spectroscopy (ICP-OES). The measured mass fraction of Pb was 1.2%, 2.1%, 3.4%, and 6.2%, respectively.  $\text{Pb/Cu}_2\text{O-x}$  was taken and uniformly loaded on a gas diffusion electrode (GDE) and electrochemically activated in a 3 M KOH aqueous solution. Finally, the e- $\text{Pb/Cu}_2\text{O-x}$  GDE was obtained, where e stands for “after electrochemical activation”.

The structure of e- $\text{Pb/Cu}_2\text{O-x}$  was confirmed through powder X-ray diffraction (XRD) characterization, which showed the diffraction peaks of Cu,  $\text{Cu}_2\text{O}$ , and PbO (Fig. S1†). This result preliminarily indicated the co-existence of  $\text{Cu}^0$  and  $\text{Cu}^+$  in the catalysts. The scanning electron microscopy (SEM) and transmission electron microscopy (TEM) images of e- $\text{Pb/Cu}_2\text{O-3.4\%}$  are shown in Fig. 1B and C. We can see that e- $\text{Pb/Cu}_2\text{O-3.4\%}$  has a sheet-like morphology with an average diameter ranging from 400 to 600 nm. Other e- $\text{Pb/Cu}_2\text{O-x}$  had similar morphological characteristics (Fig. S2†). The high-resolution TEM (HRTEM) images in Fig. 1D and Fig. S3† show that the spacings of the lattice fringe for all of the e- $\text{Pb/Cu}_2\text{O-x}$  were 0.25 nm and 0.21 nm, which can be assigned to the (111) plane of  $\text{Cu}_2\text{O}$  and (111) plane of Cu, respectively.<sup>39,40</sup> Elemental distribution mappings illustrated a uniform distribution of Cu, O, and Pb over the entire architecture (Fig. 1E).

X-ray photoelectron spectroscopy (XPS) analysis was performed to study the composition and chemical nature of the

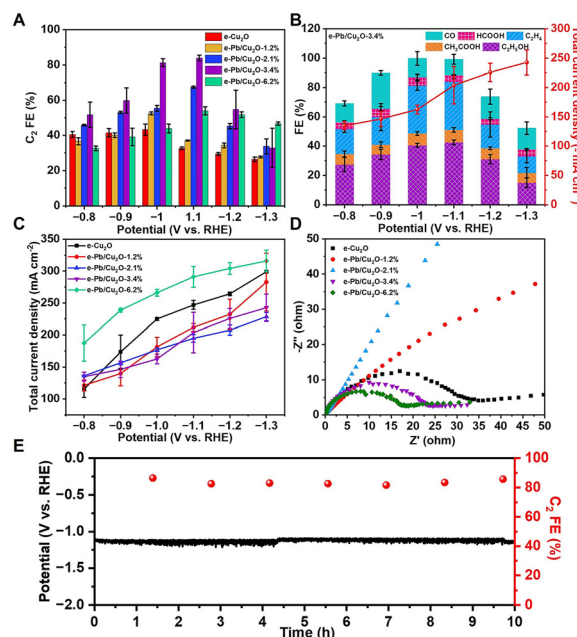


**Fig. 1** (A) Schematic illustration of the preparation of e- $\text{Pb/Cu}_2\text{O-x}$ . (B) SEM image of e- $\text{Pb/Cu}_2\text{O-3.4\%}$ . (C and D) TEM and HRTEM images of e- $\text{Pb/Cu}_2\text{O-3.4\%}$ . (E) Elemental mappings of e- $\text{Pb/Cu}_2\text{O-3.4\%}$ . (F) Cu LMM XPS spectra of e- $\text{Pb/Cu}_2\text{O-3.4\%}$ . (G) Pb 4f XPS spectra of e- $\text{Pb/Cu}_2\text{O-3.4\%}$ . (H) Cu K-edge XANES spectra of e- $\text{Pb/Cu}_2\text{O-3.4\%}$ . (I) Pb L-edge XANES spectra of e- $\text{Pb/Cu}_2\text{O-3.4\%}$ .

catalysts. The spectra of Cu 2p for e-Pb/Cu<sub>2</sub>O-3.4% (Fig. S4†) displayed the fitting peaks, which were attributed to Cu<sup>+</sup> or Cu<sup>0</sup> (932.1 eV and 951.9 eV),<sup>41,42</sup> and the Auger LMM spectra of Cu further clarified two fitting peaks at 569.9 eV and 567.9 eV that belonged to Cu<sup>+</sup> and Cu<sup>0</sup> (Fig. 1F).<sup>42,43</sup> The spectra of Pb 4f (Fig. 1G) displayed the two fitting peaks of Pb at 138.5 and 143.2 eV, indicating that the valence of Pb in e-Pb/Cu<sub>2</sub>O-3.4% was +2.<sup>44</sup> The same conclusions can be obtained from the XPS spectra of other e-Pb/Cu<sub>2</sub>O-*x* samples (Fig. S5–S7†). Furthermore, X-ray absorption spectroscopy (XAS) was used to explore the detailed electronic structures of e-Pb/Cu<sub>2</sub>O-3.4%. The X-ray absorption near-edge structure (XANES) spectra showed that the near-edge absorption energy (*E*<sub>0</sub>) of Cu K-edge for e-Pb/Cu<sub>2</sub>O-3.4% was between those of Cu and Cu<sub>2</sub>O (Fig. 1H). It can be concluded that the average valence of Cu in e-Pb/Cu<sub>2</sub>O-3.4% was between 0 and +1.<sup>45–47</sup> The coordination structures for Cu can be confirmed by the extended X-ray absorption fine structure (EXAFS) spectra (Fig. S8A and S9A†). The Cu–O coordination of Cu<sub>2</sub>O and e-Pb/Cu<sub>2</sub>O-3.4% were at the same peak (1.52 Å). The peak of the Cu–Cu coordination (2.61 Å) of e-Pb/Cu<sub>2</sub>O-3.4% was between Cu and Cu<sub>2</sub>O. These results further indicated the coexistence of Cu<sub>2</sub>O and Cu structures in e-Pb/Cu<sub>2</sub>O-3.4%.

Similarly, the *E*<sub>0</sub> of Pb L-edge was almost the same as that of PbO (Fig. 1I), demonstrating that the valence of Pb in e-Pb/Cu<sub>2</sub>O-3.4% was +2.<sup>48</sup> The EXAFS spectra of Pb (Fig. S8B and S9B†) also showed that the Pb–O and Pb–Pb coordination was similar to those of PbO. These results are consistent with the data of XRD, XPS, and HRTEM.

The CO<sub>2</sub>RR electrocatalytic activity of the as-prepared catalysts was carried out in a flow cell. In a typical experiment, the 3 M KOH aqueous solution was used as the electrolyte. <sup>1</sup>H nuclear magnetic resonance (NMR) spectroscopy and gas chromatography (GC) were adopted to determine and quantify the liquid and gas products from the CO<sub>2</sub>RR. We can detect ethanol, acetate, formate, and trace methanol in liquid products, and H<sub>2</sub>, CO, C<sub>2</sub>H<sub>4</sub>, and trace CH<sub>4</sub> in gas products. It can be found that the total FE of C<sub>2</sub> products with e-Pb/Cu<sub>2</sub>O-*x* showed volcano-shaped dependence on the applied potentials (Fig. 2A). Significantly, e-Pb/Cu<sub>2</sub>O-3.4% had the best activity with the highest C<sub>2</sub> product FE. The maximum C<sub>2</sub> product FE could reach 83.9% with a current density of 203.8 mA cm<sup>-2</sup> at -1.1 V vs. RHE (Fig. 2B and C), which is comparable to those of many reported systems (Table S1†). The FEs of C<sub>2</sub>H<sub>4</sub>, ethanol, and acetate were 32.8%, 42.5%, and 8.6%, respectively. Different Pb contents in the catalysts can affect the CO<sub>2</sub>-to-C<sub>2</sub> product performance obviously (Fig. S10†). The FE of C<sub>2</sub> products increased with the increase of Pb. However, when the mass fraction of Pb increased to 6.2%, the yield of CO increased, and the FE of the C<sub>2</sub> product was only 54.0%. In addition, different Pb contents in the catalysts can also affect the HER in the CO<sub>2</sub>RR (Fig. S11†). The FE of H<sub>2</sub> was reduced until the Pb increased to 3.4%. Then the H<sub>2</sub> FE increased when the mass fraction of Pb increased to 6.2%. We then performed electrochemical impedance spectroscopy (EIS) to investigate the electrode/electrolyte interface properties. The



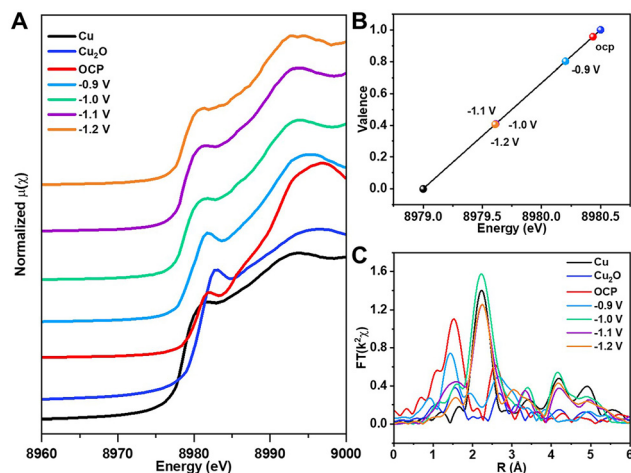
**Fig. 2** (A) The FE for C<sub>2</sub> production over various catalysts at different applied potentials. (B) The FE of different products and total current density for e-Pb/Cu<sub>2</sub>O-3.4% at different applied potentials. (C) The total current density for e-Cu<sub>2</sub>O, e-Pb/Cu<sub>2</sub>O-1.2%, e-Pb/Cu<sub>2</sub>O-2.1%, e-Pb/Cu<sub>2</sub>O-3.4% and e-Pb/Cu<sub>2</sub>O-6.2% at different applied potentials. (D) Electrochemical impedance spectra for e-Cu<sub>2</sub>O, e-Pb/Cu<sub>2</sub>O-1.2%, e-Pb/Cu<sub>2</sub>O-2.1%, e-Pb/Cu<sub>2</sub>O-3.4%, and e-Pb/Cu<sub>2</sub>O-6.2% at the open circuit potential (OCP) shown as a Nyquist plot. (E) Stability test for e-Pb/Cu<sub>2</sub>O-3.4% at a constant current density of 200 mA cm<sup>-2</sup>.

Nyquist plots were obtained by running the experiments at an open circuit potential (Fig. 2D). e-Pb/Cu<sub>2</sub>O-3.4% had lower interfacial charge transfer resistance, which ensures a faster electron transfer during the reaction. These results indicated that appropriate Pb doping in e-Pb/Cu<sub>2</sub>O-*x* could improve the C<sub>2</sub> product selectivity.

The stability of e-Pb/Cu<sub>2</sub>O-3.4% in the flow cell system at a constant current density of 200 mA cm<sup>-2</sup> is shown in Fig. 2E. There was no obvious decay in the potential and FE of C<sub>2</sub> products during the 10 h test. e-Pb/Cu<sub>2</sub>O-3.4% after the reaction, denoted as e-Pb/Cu<sub>2</sub>O-3.4%-R, was examined by TEM and XRD. The morphology and structure of the catalysts did not change notably, further indicating its excellent stability (Fig. S12 and S13†). The fine structure was analyzed by XAS (Fig. S14†). The valence and structure of Cu and Pb in e-Pb/Cu<sub>2</sub>O-3.4%-R were almost the same as those in e-Pb/Cu<sub>2</sub>O-3.4%. These results suggest that the structure of the catalyst can remain stable during the CO<sub>2</sub>RR.

To explore the behavior of e-Pb/Cu<sub>2</sub>O-3.4% in the CO<sub>2</sub>RR, we carried out detailed experimental analysis. *In situ* XAS was first performed to monitor the valence change and coordination environment of Cu under different potentials. In Fig. 3A, the Cu K-edge *E*<sub>0</sub> of e-Pb/Cu<sub>2</sub>O-3.4% reflects the change in the oxidation state. Compared with the data of the Cu K-edge *E*<sub>0</sub> of Cu foil and Cu<sub>2</sub>O, we can determine the valence of Cu in e-Pb/Cu<sub>2</sub>O-3.4% at each applied potential





**Fig. 3** (A) *In situ* XANES spectra at Cu K-edge for e-Pb/Cu<sub>2</sub>O-3.4% at different applied potentials. (B) The average oxidation state of Cu in e-Pb/Cu<sub>2</sub>O-3.4% at different applied potentials from Cu K-edge XANES. (C) R spatial data of Cu K-edge EXAFS spectra for e-Pb/Cu<sub>2</sub>O-3.4% at different applied potentials.

(Fig. 3B). It can be seen that the average valence of Cu in e-Pb/Cu<sub>2</sub>O-3.4% decreased from +0.95 at OCP to +0.8 at -0.9 V vs. RHE, and finally it would be stable around +0.4 under a higher potential. This indicated that the local electronic structure of Cu with positive valence sites could be retained during the CO<sub>2</sub>RR.<sup>45–47</sup> From the EXAFS spectra in Fig. 3C, it can be seen that the Cu–Cu and Cu–O coordination is close to those in Cu<sub>2</sub>O at -0.9 V vs. RHE, suggesting that e-Pb/Cu<sub>2</sub>O-3.4% could still maintain the Cu<sub>2</sub>O structure, but a small amount of Cu<sup>0</sup> appeared. When the applied potential increased to a value higher than -1.0 V vs. RHE, the Cu–Cu coordination of e-Pb/Cu<sub>2</sub>O-3.4% was close to that of Cu foil, and the Cu–O coordination of e-Pb/Cu<sub>2</sub>O-3.4% was close to that of Cu<sub>2</sub>O. The complete Cu<sub>2</sub>O structure could not be maintained in the catalysts, and a part of Cu<sup>+</sup> was reduced to Cu<sup>0</sup>. These results prove that the Cu<sup>0</sup>–Cu<sup>+</sup> structure exists in e-Pb/Cu<sub>2</sub>O-3.4% during the electrochemical CO<sub>2</sub>RR.

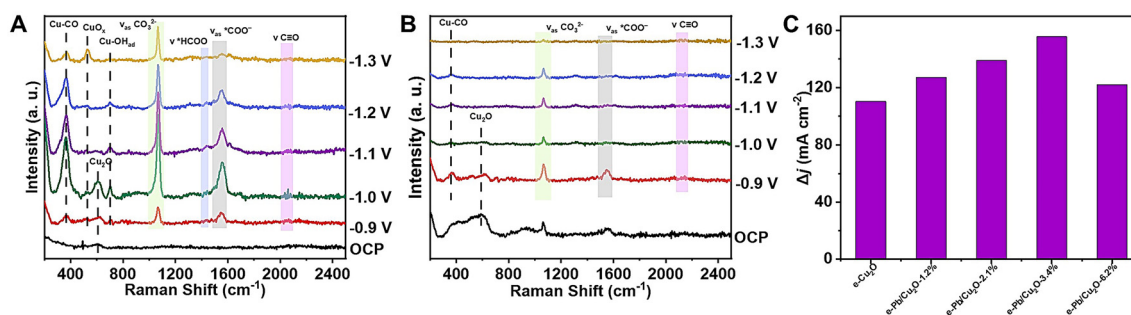
*In situ* Raman spectroscopy measurements were conducted to identify the possible intermediates. As shown in Fig. 4A, the

Raman peaks around 2060 cm<sup>-1</sup>, 1556 cm<sup>-1</sup>, and 1436 cm<sup>-1</sup> can be attributed to \*CO, \*COO<sup>-</sup>, and \*HCOO<sup>-</sup>, respectively.<sup>15, 49–53</sup> These important reaction intermediates were adsorbed on the surface of e-Pb/Cu<sub>2</sub>O-3.4% during the CO<sub>2</sub>RR, which could promote the occurrence of C–C coupling and increase the FE of C<sub>2</sub> products. The peak at 603 cm<sup>-1</sup> was attributed to the Cu<sub>2</sub>O structure.<sup>50,53,54</sup> However, it would disappear when the applied potential was higher than -1.1 V vs. RHE. Moreover, the new characteristic peak appeared at 525 cm<sup>-1</sup>, which can be attributed to CuO<sub>x</sub>.<sup>50,53,54</sup> It suggested the existence of the Cu<sup>0</sup>–Cu<sup>+</sup> structure in the catalysts. The signal of Cu–CO was also detected at 361 cm<sup>-1</sup>, indicating that the generated CO was adsorbed on the catalyst.

For comparison, we also obtained the *in situ* Raman spectra over e-Cu<sub>2</sub>O (Fig. 4B). At a potential of -1.1 V vs. RHE, we cannot observe the CuO<sub>x</sub> characteristic peak at 525 cm<sup>-1</sup>, indicating that Cu<sub>2</sub>O was reduced to Cu during the reaction.<sup>55</sup> Compared with the Cu<sup>0</sup>–Cu<sup>+</sup> structure, Cu<sup>0</sup> had a poor performance for C–C coupling. Therefore, the doping of Pb in the catalyst could stabilize the Cu<sup>0</sup>–Cu<sup>+</sup> structure and improve the catalytic performance for CO<sub>2</sub>-to-C<sub>2</sub> products.

According to Raman data, CO is an important intermediate in CO<sub>2</sub>-to-C<sub>2</sub> products. The increase of CO adsorption on the catalysts can increase the C<sub>2</sub> product FE. The detailed experimental procedures of the gas electroresponse experiments are discussed in the Methods section. The gas electroresponse experiments clearly showed the capacity for the adsorption of CO on e-Pb/Cu<sub>2</sub>O-x and e-Cu<sub>2</sub>O (Fig. 4C).<sup>56</sup> Compared with other materials, e-Pb/Cu<sub>2</sub>O-3.4% showed the best capacity of adsorbing CO. This indicated that moderate Pb doping can improve the adsorption of a \*CO intermediate, leading to high catalytic performance for C<sub>2</sub> product formation.

To better understand the reaction mechanism, we performed correlative theoretical calculations through DFT. All the simulated data and detailed procedures are shown in the ESI.† Based on the materials characterization, we have established the structural model of e-Pb/Cu<sub>2</sub>O-3.4%. For e-Pb/Cu<sub>2</sub>O-3.4%, a Cu(111) and Cu<sub>2</sub>O(111) facet heterostructure was used as a model and a part of Cu in Cu<sub>2</sub>O was replaced with Pb (Fig. S15†). We named it Cu(111)–Cu<sub>2</sub>O(111)–Pb. Previous studies showed that the adsorption of \*CO on Cu<sup>+</sup> is stronger



**Fig. 4** (A) *In situ* Raman spectra at different applied potentials for e-Pb/Cu<sub>2</sub>O-3.4%. (B) *In situ* Raman spectra at different applied potentials for e-Cu<sub>2</sub>O. (C) The results of CO adsorption responses for the catalysts.

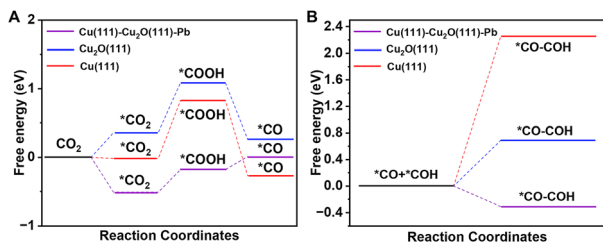


Fig. 5 (A) Gibbs free-energy diagrams for CO<sub>2</sub> activation on different simulated models. (B) Gibbs free-energy diagrams for C–C coupling on different simulated models.

than that on Cu<sup>0</sup>.<sup>28,57</sup> The Cu<sub>2</sub>O(111) facet and Cu(111) were used to compare with Cu(111)–Cu<sub>2</sub>O(111)–Pb to explore the effect of Pb doping.

The catalytic pathway of CO<sub>2</sub> activation is illustrated in Fig. 5A. It can be seen that the formation of \*COOH from \*CO<sub>2</sub> on the three models was highly endergonic and acted as the rate-determining step. The Gibbs free energy for \*CO<sub>2</sub> to \*COOH on Cu(111)–Cu<sub>2</sub>O(111)–Pb was only about 0.33 eV, which was much lower than that on Cu<sub>2</sub>O(111) (0.73 eV) and Cu(111) (0.85 eV). This indicated that the doping of Pb greatly reduced the reaction energy barrier of \*CO<sub>2</sub> to \*COOH, which contributed to easier CO<sub>2</sub> activation. The C–C coupling process was studied subsequently (Fig. 5B). We have compared different coupling pathways (\*CO–\*CO, \*CO–\*COH, \*CO–\*CHO, and \*CHO–\*CHO) and found that the \*CO–\*COH coupling had a lower Gibbs free energy. The Gibbs free energy of this coupling process on Cu(111)–Cu<sub>2</sub>O(111)–Pb was –0.31 eV, while it would be highly endergonic on Cu<sub>2</sub>O(111) (0.69 eV) and Cu(111) (2.26 eV).

Therefore, by combining the above simulation data, it is found that Pb doping can be beneficial not only for the activation of CO<sub>2</sub> but also the improvement of the C–C coupling, leading to the enhancement of the CO<sub>2</sub>RR to form C<sub>2</sub> products.

## Conclusions

In summary, a series of Pb-doped Cu<sub>2</sub>O catalysts have been successfully designed and synthesized for the efficient electrochemical CO<sub>2</sub>RR to form C<sub>2</sub> products. e-Pb/Cu<sub>2</sub>O-3.4% exhibited the highest performance with a C<sub>2</sub> product FE of 83.9% in the flow cell. The catalyst also showed good stability in 10 h. *In situ* characterization revealed that Pb doping could stabilize the Cu<sup>0</sup>–Cu<sup>+</sup> structure in e-Pb/Cu<sub>2</sub>O-3.4% during the electrochemical CO<sub>2</sub>RR, which had strong adsorption ability for the \*CO intermediate. DFT calculations suggested that the doping of Pb could reduce the reaction energy barrier of \*CO<sub>2</sub> to \*COOH and C–C coupling processes simultaneously. This work provides a facile strategy for the design of Cu-based catalysts to improve the production of C<sub>2</sub> products from the CO<sub>2</sub>RR. We believe that it may inspire new exploration of electrocatalyst design in the future.

## Author contributions

X.D.M., X.F.S., and B.X.H. proposed the project, designed the experiments, and wrote the manuscript; M.X.D. performed the whole experiments; X.N.S., L.B.Z., L.M.W., J.Q.F., S.H.J., X.X.T., and L.X. performed the analysis of experimental data; X.F.S. and B.X.H. supervised the whole project.

## Data availability

All experimental data are available in the ESI.†

## Conflicts of interest

The authors declare no competing financial interests.

## Acknowledgements

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