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# The spatial distribution of cobalt phthalocyanine and copper nanocubes controls the selectivity towards C<sub>2</sub> products in tandem electrocatalytic CO<sub>2</sub> reduction†

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The coupling of CO-generating molecular catalysts with copper electrodes in tandem schemes is a promising strategy to boost the formation of multi-carbon products in the electrocatalytic reduction of CO<sub>2</sub>. While the spatial distribution of the two components is important, this aspect remains underexplored for molecular-based tandem systems. Herein, we address this knowledge gap by studying tandem catalysts comprising Co-phthalocyanine (CoPc) and Cu nanocubes (Cu<sub>cub</sub>). In particular, we identify the importance of the relative spatial distribution of the two components on the performance of the tandem catalyst by preparing CoPc-Cu<sub>cub</sub>/C, wherein the CoPc and Cu<sub>cub</sub> share an interface, and CoPc-C/Cu<sub>cub</sub>, wherein the CoPc is loaded first on carbon black (C) before mixing with the Cu<sub>cub</sub>. The electrocatalytic measurements of these two catalysts show that the faradaic efficiency towards C<sub>2</sub> products almost doubles for the CoPc-Cu<sub>cub</sub>/C, whereas it decreases by half for the CoPc-C/Cu<sub>cub</sub>, compared to the Cu<sub>cub</sub>/C. Our results highlight the importance of a direct contact between the CO-generating molecular catalyst and the Cu to promote C–C coupling, which hints at a surface transport mechanism of the CO intermediate between the two components of the tandem catalyst instead of a transfer *via* CO diffusion in the electrolyte followed by re-adsorption.

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## Introduction

The electrochemical CO<sub>2</sub> reduction reaction (CO<sub>2</sub>RR) is an attractive route for CO<sub>2</sub> recycling, wherein value-added chemicals and fuels are produced from CO<sub>2</sub> while also storing electricity generated from renewable energy sources.<sup>1,2</sup> Among all catalysts, Cu-based materials provide suitable intermediate binding energies to drive CO<sub>2</sub>RR towards highly reduced products (*e.g.* methane, ethylene, ethanol), which are of interest due to their high energy densities.<sup>3,4</sup> To date, tremendous effort has been dedicated to increasing the selectivity towards targeted products of the reaction, which produces over sixteen different products in various yields when performed over a polycrystalline Cu foil.<sup>5</sup>

Cu-based tandem schemes have emerged as a valid strategy to enhance the selectivity of CO<sub>2</sub>RR towards multicarbon products (C<sub>2+</sub>) by decoupling the CO<sub>2</sub> to CO and the CO to C<sub>2+</sub> reduction steps.<sup>6–9</sup> Previous work has demonstrated that increasing the local

concentration of CO and/or the surface coverage of adsorbed CO (\*CO) decreases the energetic barrier for C–C coupling, which is the rate-determining step towards C<sub>2+</sub> products on Cu surfaces.<sup>10–17</sup> In addition to their intrinsic catalytic properties (*i.e.* turnover frequency, selectivity, overpotential), the relative spatial arrangement of the CO-producing component and of the Cu catalyst plays an important role in defining the efficiency of such tandem systems.<sup>9,18–20</sup> For example, researchers have synthesized reverse core–shell structures, with the CO-producing catalyst (Ag) in the core and the Cu as a shell, which was found to maximize CO utilization and enhance the C<sub>2+</sub> activity compared to core–shell structures with Cu in the core.<sup>19</sup> In an alternative approach, similar benefit was obtained by rationally segmenting the Ag and Cu components in gas-diffusion electrodes.<sup>20</sup> In both cases, the spatial distribution of the two catalysts impacts the modality of CO transport to the active sites, which can occur either *via* surface diffusion or *via* sequential adsorption following transport through the gas phase or the electrolyte. Thus, optimization of the tandem catalyst configuration emerges as an important parameter to modulate the CO utilization efficiency.

Most studies on the design of tandem systems have so far focused on the coupling of Cu with a second metallic domain (Zn,<sup>21,22</sup> Au<sup>23</sup> or Ag<sup>24–27</sup>) as the CO-generating catalyst. Molecular catalysts, such as cobalt phthalocyanine<sup>28</sup> and iron

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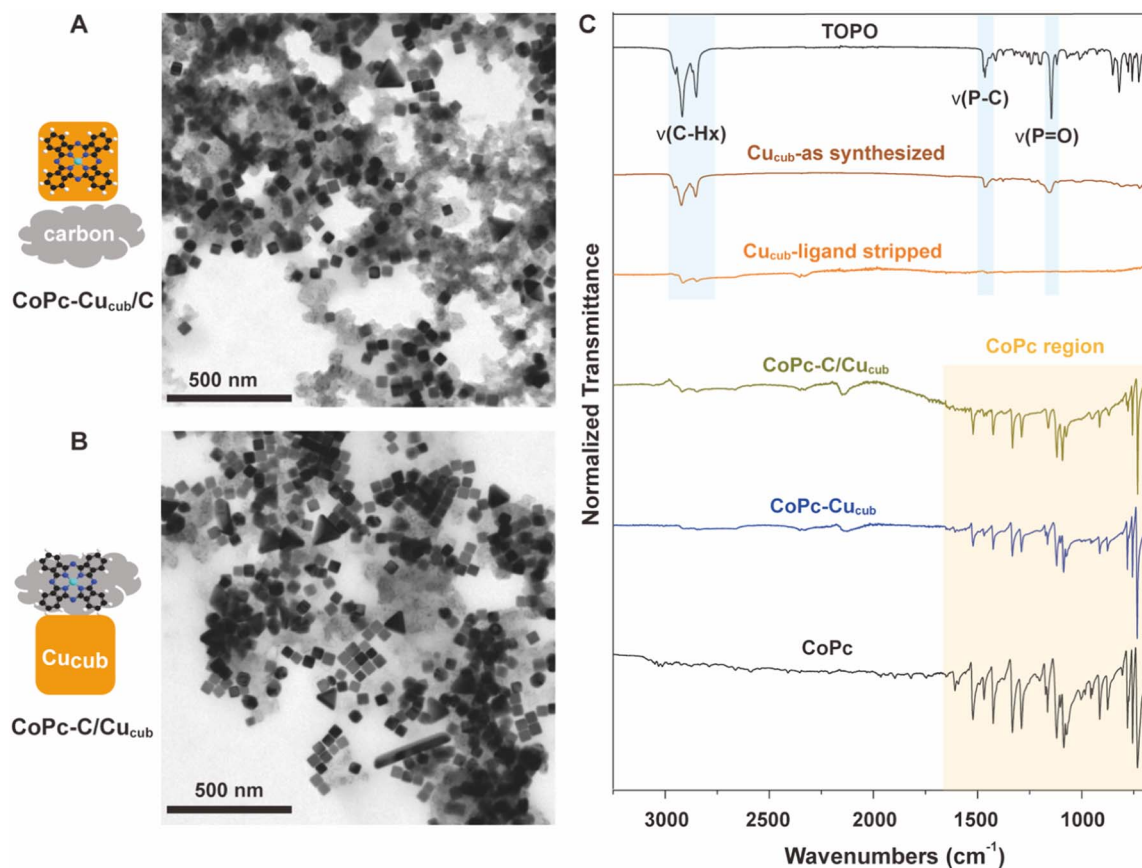
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porphyrin<sup>29,30</sup> have also been successfully employed as the CO-generating catalyst in the context of tandem schemes. Compared to metals, molecular catalysts offer additional versatility for varying the local CO concentration and production rate thanks to the superior chemical tunability *via* synthetic modifications of molecules *versus* materials.<sup>30</sup> Despite their promise, no experimental work has investigated yet the influence of the spatial arrangement of the components on the catalytic performance of molecular-Cu based tandem systems.

In this work, we investigate the impact of a direct interface between the molecular catalyst and the Cu electrode on the C–C coupling efficiency. We use colloiddally dispersible Cu nanocubes ( $\text{Cu}_{\text{cub}}$ ) enclosed by (100) facets as model catalysts. Their high surface to volume ratio, which is ideal for exploring surface phenomena, as well as their intrinsic selectivity towards ethylene production, justify this choice.<sup>31–33</sup> To work alongside  $\text{Cu}_{\text{cub}}$ , we select cobalt phthalocyanine (CoPc) as the CO-producing molecular catalyst. In addition to being highly active and stable when immobilized on conductive substrates for  $\text{CO}_2\text{RR}$ , the reduction potential window matches the applied potentials of  $\text{Cu}_{\text{cub}}$  in aqueous solutions.<sup>34–37</sup> Additionally, CoPc readily

adsorbs on different surfaces, including carbon and metallic Cu, *via* electrostatic interactions, thanks to the Pc planar structure.<sup>38–40</sup> Finally, both catalysts (*i.e.* the CoPc and the  $\text{Cu}_{\text{cub}}$ ) can be produced in the form of inks, which facilitates the manipulation of their surface interactions. We exploit these intrinsic characteristics of the  $\text{Cu}_{\text{cub}}$  and of the CoPc to create tandem catalysts comprising CoPc- $\text{Cu}_{\text{cub}}$ /C, where the CoPc shares a direct interface with the  $\text{Cu}_{\text{cub}}$ , and CoPc-C/ $\text{Cu}_{\text{cub}}$ , where the CoPc is first adsorbed on carbon black and then mixed with the  $\text{Cu}_{\text{cub}}$ . We find that the CoPc- $\text{Cu}_{\text{cub}}$ /C catalyst exhibits a faradaic efficiency (FE) and partial current density for  $\text{C}_2$  products which are double the values measured for pristine  $\text{Cu}_{\text{cub}}$ /C without the molecular component present. In contrast, the FE for  $\text{C}_2$  products decrease by half for the CoPc-C/ $\text{Cu}_{\text{cub}}$  compared to the  $\text{Cu}_{\text{cub}}$ /C. These results demonstrate that CO utilization is much more efficient in the CoPc- $\text{Cu}_{\text{cub}}$ /C, where a close contact between the two components of the tandem catalyst exists. Thus, we propose that surface transport, which requires a direct interface, dominates over electrolyte-mediated diffusion-readsorption and is the favorable pathway to maximize the CO utilization efficiency and  $\text{C}_2$  product formation in these molecular-based tandem catalysts.



**Fig. 1** (A and B) Bright field TEM images of CoPc- $\text{Cu}_{\text{cub}}$ /C (A) and CoPc-C/ $\text{Cu}_{\text{cub}}$  (B), confirming that the structure of  $\text{Cu}_{\text{cub}}$  is retained in both cases following catalyst preparation. The synthesized tandem catalyst structures are schematically indicated on the left. (C) FTIR spectra of TOPO ligand, the as-synthesized  $\text{Cu}_{\text{cub}}$ , ligand-stripped  $\text{Cu}_{\text{cub}}$ , CoPc- $\text{Cu}_{\text{cub}}$ , CoPc-C/ $\text{Cu}_{\text{cub}}$  and CoPc. These results demonstrate the successful removal of TOPO ligand from  $\text{Cu}_{\text{cub}}$  and the retention of CoPc within both catalyst assemblies.



## Results and discussion

The CoPc-Cu<sub>cub</sub>/C catalyst was obtained by first mixing the CoPc molecules with the Cu<sub>cub</sub> in dimethylformamide (DMF), followed by the addition of carbon black into CoPc-Cu<sub>cub</sub>. Prior to mixing, the native ligands of the Cu<sub>cub</sub> were removed *via* solvent washing, which allowed them to become dispersible in DMF. The molecular catalyst was added in excess to the washed Cu<sub>cub</sub> in order to maximize the Cu surface coverage (see Experimental section in ESI† for details). The amount of the adsorbed molecular catalyst (~1 nmol per cm<sup>2</sup> per working electrode) was quantified by inductively coupled plasma-optical emission spectrometry (ICP-OES) (Table S1†). The CoPc-C/Cu<sub>cub</sub> catalyst was prepared by adsorbing the same total amount of CoPc on the carbon black prior to mixing with the Cu<sub>cub</sub>. For reference, carbon black was also added to the individual Cu<sub>cub</sub> and CoPc systems (Cu<sub>cub</sub>/C and CoPc/C) to enable a direct comparison among each of the different systems.

Fig. 1 reports the transmission electron microscopy (TEM) and Fourier transform infrared spectroscopy (FTIR) characterization of the as-obtained catalysts. TEM reveals the intact cubic shape of the Cu<sub>cub</sub> in both CoPc-Cu<sub>cub</sub>/C (Fig. 1A) and CoPc-C/Cu<sub>cub</sub> (Fig. 1B), which proves that the synthesis procedure does not impact the Cu<sub>cub</sub> morphology. The low contrast material in both images is the carbon black. The FTIR data presented in Fig. 1C show that, first of all, the washed Cu<sub>cub</sub> (Cu<sub>cub</sub>-ligand stripped) has ligand-free surfaces. Indeed, the intensity of the –

CH– stretching at 2915 cm<sup>-1</sup> is negligible compared to the as-synthesized sample and the P=O vibration is completely absent. Both vibrational modes are representative of the native trioctylphosphine oxide ligands of the Cu<sub>cub</sub> (area shaded in blue), confirming their removal from the Cu nanoparticles. The characteristic peaks of CoPc (area shaded in yellow: with Pc ring at 753 cm<sup>-1</sup>, pyrrole C–N asymmetric stretch at approximately 1086 cm<sup>-1</sup>, and the isoindole and pyrrole stretch at 1200–1500 cm<sup>-1</sup>) are present in the CoPc-Cu<sub>cub</sub> sample even after thoroughly washing, proving that the CoPc is strongly adsorbed on the Cu<sub>cub</sub> surface, likely due to the expected electrostatic interactions. Similarly, the characteristic IR peaks of CoPc are present for the CoPc-C/Cu<sub>cub</sub> catalyst. The background signal in the region from 700 cm<sup>-1</sup> to 2000 cm<sup>-1</sup>, is attributed to the carbon black (Fig. S1†).

We further characterized the catalysts by high-angle annular dark field scanning transmission electron microscopy (HAADF-STEM) coupled with energy-dispersive X-ray (EDX) spectroscopy to learn more about the spatial distributions of the different components (Fig. 2). For the case of the CoPc-Cu<sub>cub</sub> catalyst, the Cu and Co signals are co-localized on the Cu<sub>cub</sub> and no free CoPc molecules are present on the grid (Fig. 2A and B). By contrast, for the case of CoPc-C/Cu<sub>cub</sub> catalyst, the CoPc is present only on the carbon black and not on the Cu<sub>cub</sub> (Fig. 2C and D). These results confirm the successful assembly of two different catalyst motifs according to the designed structures depicted by the schematic illustrations in Fig. 1A.

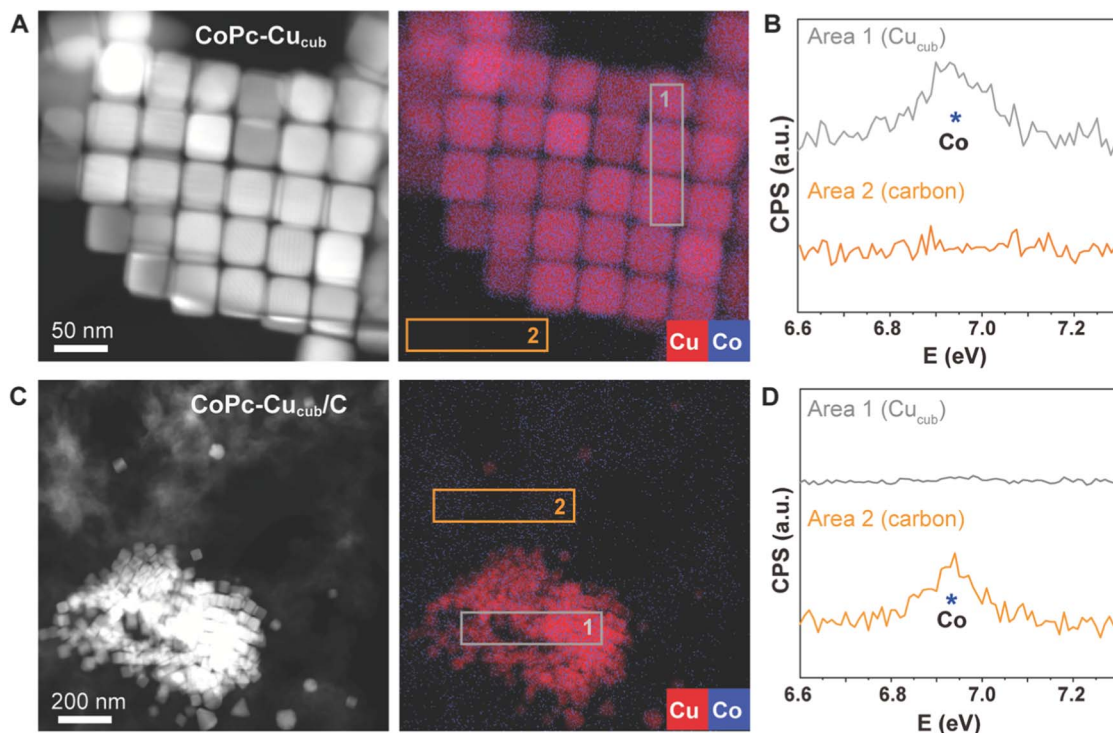
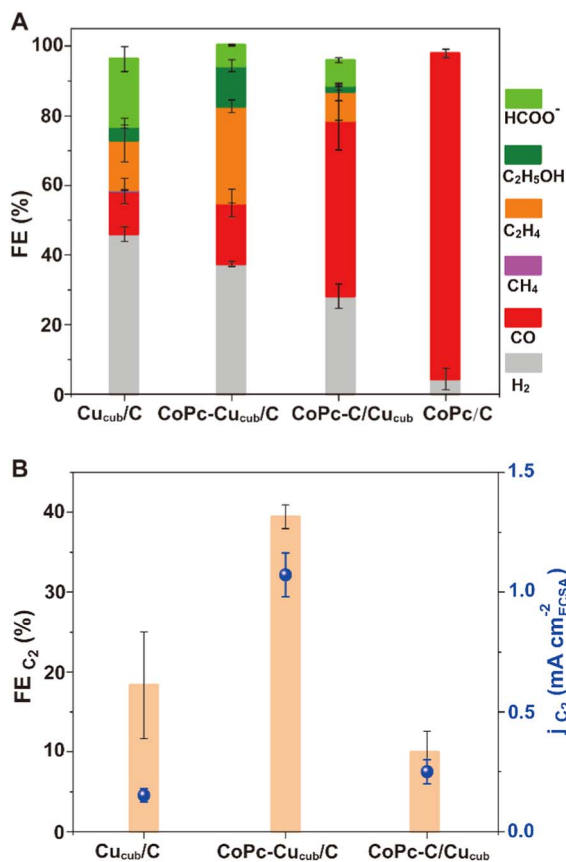


Fig. 2 (A and C) HAADF-STEM images and corresponding EDX elemental maps, along with (B and D) EDX spectra for CoPc-Cu<sub>cub</sub>/C (A and B) and CoPc-C/Cu<sub>cub</sub> (C and D). The EDX maps show the spatial distributions of Cu (red) and Co (blue). The carbon is the low contrast material in (C) and is not colored for the sake of clarity. The EDX spectra report the intensity of the Co signals in two different areas of the map, which correspond to the Cu<sub>cub</sub> (area 1) and to the carbon (area 2), as indicated on the colored maps.





**Fig. 3** (A) FEs for all gaseous products (*i.e.*, H<sub>2</sub>, CO, CH<sub>4</sub>, C<sub>2</sub>H<sub>4</sub>) and the main liquid products (*i.e.*, formate, C<sub>2</sub>H<sub>5</sub>OH), (B) FE<sub>C<sub>2</sub></sub> and corresponding partial current density  $j_{C_2}$  normalized by the ECSA for the Cu<sub>cub</sub>/C, CoPc-Cu<sub>cub</sub>/C, CoPc-C/Cu<sub>cub</sub> and CoPc/C. The loading of Cu<sub>cub</sub> was 20 μg cm<sup>-2</sup> and the loading of CoPc anchored on the Cu<sub>cub</sub> and on the carbon black was 1.1 nmol cm<sup>-2</sup> and 0.9 nmol cm<sup>-2</sup>, respectively. These data were collected in an H-cell configuration using 0.1 M KHCO<sub>3</sub> as the electrolyte with the working electrode poised at -1.0 V vs. RHE for 1 h. The reported values are an average of three independent experiments with error bars indicating the standard deviations. Compared to the reference Cu<sub>cub</sub>/C, a significant increase of FE<sub>C<sub>2</sub></sub> is observed for CoPc-Cu<sub>cub</sub>/C, while a decrease is observed for CoPc-C/Cu<sub>cub</sub>.

Having completed the characterization of the two catalysts, we evaluated the behavior of CoPc-Cu<sub>cub</sub>/C and CoPc-C/Cu<sub>cub</sub> towards CO<sub>2</sub>RR in a H-cell system, using CO<sub>2</sub>-saturated 0.1 M aqueous KHCO<sub>3</sub> as the supporting electrolyte (Fig. 3). We chose an electrochemical potential of -1.0 V vs. RHE (reversible hydrogen electrode) for initial comparison based on the previous knowledge that Cu<sub>cub</sub> exhibits a significant selectivity for C<sub>2</sub>H<sub>4</sub> and that CoPc possesses maximum activity for CO near this working potential. For reference, the product distributions for the tandem catalyst assemblies are also compared to those possessing only the individual components, Cu<sub>cub</sub>/C and CoPc/C. All reported current densities are normalized by the electrochemically active surface area (ECSA) (Fig. S2†).

The product distribution analysis (Fig. 3A), which is represented by the FE values, reveals that the major products observed for Cu<sub>cub</sub>/C are still present for the CoPc-Cu<sub>cub</sub>/C, but

with a clear increase of the FE for C<sub>2</sub> products (*i.e.* ethylene + ethanol). However, for the CoPc-C/Cu<sub>cub</sub>, CO production notably increases while the other products are suppressed. Importantly, FE<sub>H<sub>2</sub></sub> is suppressed for both CoPc-Cu<sub>cub</sub>/C and CoPc-C/Cu<sub>cub</sub> compared to the Cu<sub>cub</sub>/C. As expected, the major product for CoPc/C is CO, with minor production of H<sub>2</sub>, indicating that the activity of the heterogenized molecular component is preserved.

To facilitate the comparison of C<sub>2</sub> products, Fig. 3B reports the FE<sub>C<sub>2</sub></sub> and the corresponding partial current densities ( $j_{C_2}$ ) of all Cu<sub>cub</sub>-based systems. The CoPc-Cu<sub>cub</sub>/C clearly shows an enhancement of C<sub>2</sub> products with the highest FE of 39.4% and  $j_{C_2}$  of 1.07 mA cm<sup>-2</sup>. In contrast, the FE<sub>C<sub>2</sub></sub> for CoPc-C/Cu<sub>cub</sub> is only 10%, much lower than that of the Cu<sub>cub</sub>/C, which is 18.3%. The  $j_{C_2}$  are instead comparable and equal to 0.2 mA cm<sup>-2</sup> for CoPc-C/Cu<sub>cub</sub> and 0.16 mA cm<sup>-2</sup> for the Cu<sub>cub</sub>.

The high FE<sub>CO</sub> and the suppressed FE<sub>C<sub>2</sub></sub> for CoPc-C/Cu<sub>cub</sub> suggests that CO released from the molecule diffuses away rather than being utilized by the Cu<sub>cub</sub>. Longer reaction times up to 10 hours do eventually result in a slight increase of FE<sub>C<sub>2</sub></sub> and  $j_{C_2}$  for CoPc-C/Cu<sub>cub</sub> (Fig. S3†), which might result from the overall increasing concentration of CO in the electrolyte. However, the performance of CoPc-C/Cu<sub>cub</sub> remain always lower than CoPc-Cu<sub>cub</sub>/C and Cu<sub>cub</sub>/C. Decreasing the loading of the CoPc does decrease the FE<sub>CO</sub> but does not improve the FE<sub>C<sub>2</sub></sub> compared to CoPc-Cu<sub>cub</sub>/C with same CoPc loading and to Cu<sub>cub</sub>/C (Fig. S4†), which indicates that the CO transfer from CoPc-C to Cu<sub>cub</sub> does not efficiently occur in these catalysts.

To further investigate the mechanism behind the observed catalytic activity differences, we investigated the CoPc-Cu<sub>cub</sub>/C and CoPc-C/Cu<sub>cub</sub> CO<sub>2</sub>RR performance over a wide range of applied potentials and compared them to those of Cu<sub>cub</sub>/C and CoPc/C (Fig. 4). We note that the CoPc stability decreases and the selectivity for methane increases at more negative voltages (Fig. S5†). Consistent with the data in Fig. 3, CoPc-Cu<sub>cub</sub>/C exhibits a similar potential-dependent product distribution compared to Cu<sub>cub</sub>/C, but with an increased C<sub>2</sub> product yield and suppressed hydrogen generation across the entire potential range (Fig. 4A). In particular, the FE<sub>C<sub>2</sub></sub> and the corresponding partial current density of the CoPc-Cu<sub>cub</sub>/C are consistently higher than the values measured for the Cu<sub>cub</sub>/C and increase at more cathodic potentials (Fig. 4B). The best-performing CoPc-Cu<sub>cub</sub>/C possess a FE<sub>C<sub>2</sub></sub> of 48% with a partial current density of 1.5 mA cm<sup>-2</sup> at -1.05 V vs. RHE, which is 1.7 times the FE<sub>C<sub>2</sub></sub> of the Cu<sub>cub</sub>/C. In contrast, for the CoPc-C/Cu<sub>cub</sub> assembly, CO is the major product and the FE<sub>C<sub>2</sub></sub> is reduced over the whole potential range compared to Cu<sub>cub</sub>/C (Fig. 4D). Furthermore, the partial current density of the C<sub>2</sub> products in CoPc-C/Cu<sub>cub</sub> and Cu<sub>cub</sub>/C is always similar, which indicates that the Cu<sub>cub</sub> intrinsic selectivity is unaffected by the CoPc molecules (Fig. 4D). We also note that the total current densities of the tandem catalysts are higher compared to those of either the Cu<sub>cub</sub> or the CoPc alone, which indicates that both components are active (Fig. S8†). Furthermore, the overall C<sub>2</sub> production rate is higher in the tandem catalysts compared to the Cu<sub>cub</sub> alone (Fig. S8†), which is similar to what has been reported for some of the metallic/Cu tandem catalysts.<sup>41,42</sup> We highlight that



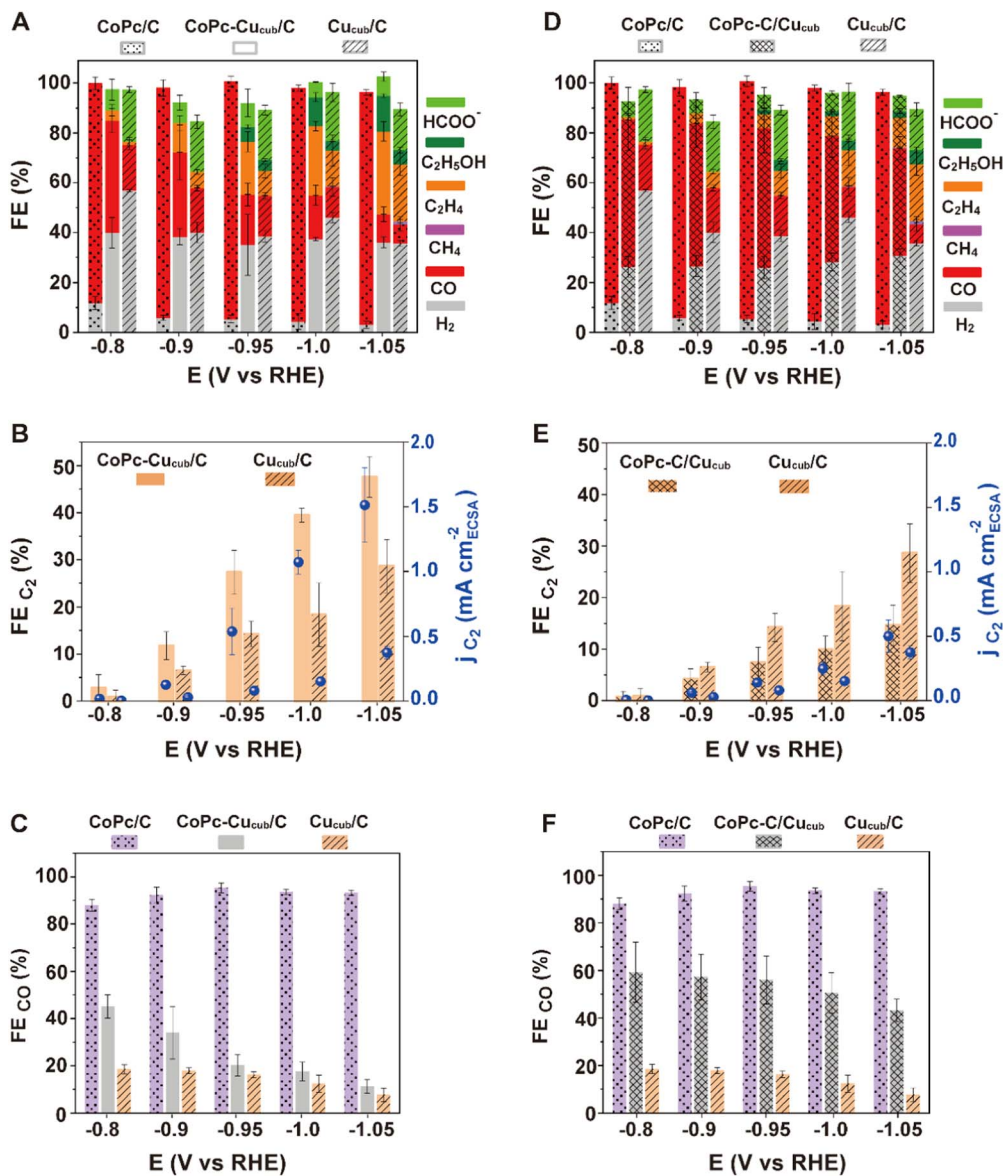


Fig. 4 (A and D) FEs for all gaseous products (*i.e.*, H<sub>2</sub>, CO, CH<sub>4</sub>, C<sub>2</sub>H<sub>4</sub>) and the main liquid products (*i.e.*, formate, C<sub>2</sub>H<sub>5</sub>OH), (B and E) FEs and partial current densities of C<sub>2</sub> products (*i.e.* C<sub>2</sub>H<sub>4</sub> + C<sub>2</sub>H<sub>5</sub>OH),  $j_{C_2}$ , normalized by the ECSA and (C and F) FEs for CO as a function of potential for Cu<sub>cub</sub>/C, CoPc-Cu<sub>cub</sub>/C, CoPc-C/Cu<sub>cub</sub> and CoPc/C. The loading of Cu<sub>cub</sub> was 20 μg cm<sup>-2</sup> and the loading of CoPc anchored on the Cu<sub>cub</sub> and on the carbon black was 1.1 nmol cm<sup>-2</sup> and 0.9 nmol cm<sup>-2</sup>, respectively. These data were collected in a H-cell system using 0.1 M KHCO<sub>3</sub> as the electrolyte for 1 h. The reported values are an average of three independent experiments with error bars indicating the standard deviations.

morphology and composition of all catalysts were maintained after CO<sub>2</sub>RR at -1.05 V vs. RHE for 1 hour (Fig. S6†).

To understand whether the C-C coupling enhancement in the CoPc-Cu<sub>cub</sub>/C catalyst is consistent with a tandem mechanism, we plotted the FE<sub>CO</sub> of the different systems at various electrochemical potentials (Fig. 4C and F). The FE<sub>CO</sub> for the CoPc-Cu<sub>cub</sub>/C decreases as the potential becomes more cathodic (Fig. 4C), concomitant with the C<sub>2</sub> production increase. This correlation is consistent with a tandem mechanism in which the CO formed on CoPc is consumed by Cu<sub>cub</sub>. By contrast, the FE<sub>CO</sub> of the CoPc-C/Cu<sub>cub</sub> catalyst remains approximately constant and is similar to that of CoPc/C across the entire potential range (Fig. 4F), proving that the CO formed on CoPc is

not consumed by Cu<sub>cub</sub> and the sequential tandem reaction does not occur.

To further validate the conclusion that the obtained results are due to a tandem effect, we performed control experiments to investigate whether other factors could be involved. XPS revealed that the electronic properties of the catalysts are not impacted by interactions between the different components and are, therefore, not responsible for the observed catalytic differences (Fig. S8 and Table S2†). In addition, we verified that neither the phthalocyanine ligand nor the Co alone on the Cu surface can provide the enhanced C-C coupling observed for CoPc-Cu<sub>cub</sub>/C (Fig. S9–S11†). Finally, no major changes in hydrophobicity, which could justify hydrogen suppression, were observed upon



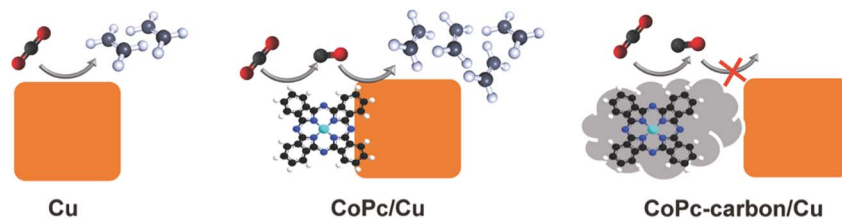


Fig. 5 Schematic illustration summarizing the effect of the spatial distribution of CO-producing molecular catalysts and Cu catalysts in tandem systems. The close spatial coupling and direct interface between the two components is crucial for enhancing C–C coupling *via* enhanced CO generation in the vicinity of Cu.

adsorption of the CoPc on the Cu<sub>cub</sub> (Fig. S12†). Overall, the tandem effect emerges as the main reason underlying the C–C coupling enhancement observed in the CoPc-Cu<sub>cub</sub>/C catalysts. Finally, encouraged by the results measured in a H-cell configuration, we included the spatial configured systems into a gas-fed flow cell, comprising the CoPc-Cu<sub>cub</sub>/C and the CoPc-C/Cu<sub>cub</sub> supported on a gas diffusion electrode as the cathode, in order to realize CO<sub>2</sub> to C<sub>2</sub> conversion at high current density (Fig. S13†). In agreement with the data discussed above, the CoPc-Cu<sub>cub</sub>/C catalyst demonstrated enhanced C<sub>2</sub> production compared to the Cu<sub>cub</sub>/C reaching a FE for C<sub>2</sub> of 62% with a partial current density of 125 mA cm<sup>-2</sup> in 1 M KHCO<sub>3</sub>.

Fig. 5 summarizes the results of this study and the proposed mechanism responsible for the observed electrocatalytic activity of molecule-copper tandem catalysts. We find that the close vicinity and direct interface between the CO-generating molecules and the Cu surface is essential to promote C–C coupling *via* the tandem effect. When the molecular component is spatially separated from the copper, the sequential tandem process does not occur and the overall multicarbon production is suppressed. These results suggest that the tandem mechanism in these catalysts is enabled by surface diffusion of the CO rather than CO diffusion in the electrolyte and sequential reabsorption.

## Conclusions

In summary, we have demonstrated that tandem catalysts must be carefully engineered to promote efficient C–C coupling. In particular, our results reveal that a direct interface between the CO-generating molecular catalyst and the Cu is a key factor for efficient CO utilization by the copper. Indeed, we have reported a CO<sub>2</sub>-to-C<sub>2</sub> conversion with FE of 48% at –1.05 V vs. RHE for CoPc-Cu<sub>cub</sub>/C catalysts, which is 1.7 times higher than that of the Cu<sub>cub</sub>/C when tested under the same conditions. In contrast, the CoPc-C/Cu<sub>cub</sub> assembly, in which the Cu<sub>cub</sub> and the CoPc were physically separated, exhibited a dramatically reduced FE<sub>C<sub>2</sub></sub> compared to the Cu<sub>cub</sub>. Overall, these findings advance our fundamental understanding of the performance-driving parameters necessary for constructing more efficient hybrid molecule/nanocrystal tandem catalysts in the future. For example, the discovery that a close spatial coupling is crucial for efficient tandem catalysts suggests that their integration into gas diffusion electrodes operating at high current density may

benefit from functional groups to anchor the molecular catalysts on the surface of the nanocatalysts.

## Data availability

Data for this paper, including TEM images, FT-IR, HAADF-STEM images and corresponding EDX and electrochemical characterization, are available at Zenodo at: <https://doi.org/10.5281/zenodo.7327628>.

## Author contributions

M. W. carried out the synthesis of the tandem catalysts and electrocatalytic testing for CO<sub>2</sub>RR. A. L. performed X-ray photoelectron spectroscopy, HAADF-STEM images and corresponding EDX and contributed to daily discussions. V. O. and I. D. S. contributed to discussions. R. B. supervised this work and coordinated the project. All the authors contributed to the writing of the manuscript.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

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