Nanostructured Copper-Based Electrodes Electrochemically Synthesized on a Carbonaceous Gas Diffusion Membrane with Catalytic Activity for the Electrocroduction of CO₂

Martina Serafini, Federica Mariani, Andrea Fasolini, Erika Scavetta, Francesco Basile,* and Domenica Tonelli*

Cite This: ACS Appl. Mater. Interfaces 2021, 13, 57451−57461

ABSTRACT: In this work, four different 4 cm²-sized nanostructured Cu-based electrocatalysts have been designed by a one-step electrodeposition process of Cu metal on a three-dimensional carbonaceous membrane. One consisted of Cu₀, and the other three were obtained by further simple oxidative treatments. Morphological, structural, and electrochemical investigations on the four materials were carried out by scanning electron microscopy, Raman spectroscopy, X-ray diffraction, linear sweep voltammetry, and potential-controlled electrolysis. All the electrocatalysts showed promising catalytic activities toward CO₂ electroreduction in liquid phase, with a remarkable selectivity toward acetic acid achieved when using the oxidized materials. In particular, the best electrocatalytic activity was observed for the Cu₂O-Cu₀ catalyst, working at a relatively low potential (−0.4 V vs RHE), which exhibited a stable and low current density of 0.46 mA cm⁻² and a productivity of 308 μmol gcat⁻¹ h⁻¹. These results were attributed to the nanostructured morphology that is characterized by many void spaces and by a high surface area, which should guarantee a large number of CuI and Cu₀ catalytic active sites. Moreover, kinetic analyses and preliminary studies about catalyst regeneration highlighted the stability of the best-performing catalyst.

KEYWORDS: nanostructure, Cu-based electrocatalyst, CO₂ conversion, electroreduction, acetic acid, gas diffusion layer

1. INTRODUCTION

The global effort in pursuing effective decarbonization strategies is depicting new scenarios toward alternative production chains. In particular, CO₂ capture and utilization play a key role in view of a substantial technological revolution in the way that energy is produced, stored, and converted nowadays from and among different energy sources and vectors. The electrochemical reduction of CO₂ under liquid- or gas-phase conditions, is a reaction by which hydrocarbons can be obtained at ambient temperature and atmospheric pressure, thus closing the carbon cycle with the production of added-value commodity chemicals, using electricity derived preferably from renewables. In this context, the fascinating concept of solar-driven chemistry, realized with photochemical systems either including inorganic¹⁻⁷ or organic biohybrid¹⁺ components, is attracting much attention thanks to the biomimetic approach toward a sustainable process, which culminates in the concept of artificial leaves.²⁻⁹ In such a system, light is absorbed at a photoanode to generate electricity that is used in situ for the electroreduction of CO₂ to chemicals, thus ideally eliminating the need for any external power supply. Of particular interest is the direct production of methanol or acetic acid, which are otherwise obtained in a high-temperature, multistep process from methane-derived syngas (MeOH) and successive carbonylation (CH₃COOH).⁴ To date, the design of lightweight, low-cost, and low-power consuming electrocatalytic platforms for high-throughput CO₂ electrocatalytic reduction (CO₂ER) represents a major challenge for future integration in the aforementioned artificial leaf. In this context, this work aims to develop novel active materials for the CO₂ER.¹¹ Due to its molecular structure, CO₂ is particularly stable from a thermodynamic point of view, and its chemical transformation into products generally requires harsh temperature and pressure conditions. Conversely, a high overpotential is required for carbon dioxide electroreduction,¹¹ which generally suffers from sluggish kinetics, multiphase rate-
limiting steps, and poor selectivity, as well as intermediate-sharing, multiple reaction pathways, whose mechanisms are still under debate. On the other hand, experimental parameters such as the catalyst properties, local pH, chemical nature of the electrolyte solution, and applied potential dramatically affect the reaction and, consequently, the product distribution.5,12,13

Due to its unique ability to catalyze CO2ER toward a number of hydrocarbons, aldehydes, and alcohols requiring more than two electron transfers with substantial Faradaic efficiencies (FE), Cu has been overall recognized as the gold standard among pure metal catalysts since 1985.13–15

In particular, Cu-based electrode interfaces are able to stabilize the chemisorbed CO2− radical anions and CO species, which are the key intermediates in the initial phase of the catalytic CO2 conversion and in the subsequent process of hydrocarbon and alcohol formation, respectively. This not only increases the CO surface coverage, thereby promoting those CO2ER pathways involving C−C coupling reactions, but also blocks the reaction sites for the parasitic hydrogen evolution reaction (HER).16,17 Recent studies concerning liquid-phase CO2ER have reported on the fundamental role of nano- and microstructures,18–21 as well as the redox state22,23 of Cu-based cathode materials in improving the catalyst’s selectivity and activity. It has been shown that metallic Cu obtained after reduction of CuO, which was formed by electropolishing and subsequent thermal annealing of a polycrystalline Cu foil, displayed enhanced current densities and Faradaic efficiencies with respect to the pristine Cu foil.24 The more active Cu catalyst was produced in situ during CO2ER at −0.5 V vs RHE and led to formate production. The effect of the orientation and thickness of CuO films electrodeposited on commercial Cu plates for CO2ER performed at −1.1 V vs RHE was investigated by online electrochemical mass spectrometry and XRD.25 On the one hand, the selectivity toward the main product (formic acid) was demonstrated to largely depend on the film thickness rather than the orientation. On the other hand, the reaction was found to be actually catalyzed by in situ formation of nanostructured Cu domains during the progress of the reaction itself. The influence of the CuO nanoparticle (NP) morphology synthesized by a hydrothermal method was studied during CO2ER at −1.7 V vs RHE.25 Again, in situ reduction to metallic Cu was reported, leading to good selectivity toward ethanol, while different electrocatalytic activities were observed depending on the initial CuO NP morphologies. Dendritic Cu catalysts electrodeposited on technical Cu mesh supports were also studied for CO2 electroreduction showing good Faradaic selectivity to formate and ethylene at −0.7 and −1.1 V vs RHE, respectively.17 Subsequent thermal annealing led to C1 > alcohol formation (EtOH and n-PrOH at −1.0 V vs RHE). Thermal and electrochemical activations of copper foils to form oxide-based catalysts for CO2ER have been recently compared by Giri et al.26 Interestingly, the authors observed a relevant difference between the oxidation methods and showed, by XPS and XRD analyses, that all oxides reduced back to Cu0 within the first few minutes of the reaction. Higher currents and Faradaic efficiencies were obtained from the activated samples and were ascribed to the increased surface area after Cu activation. Overall, these works suggest that the CO2ER actually proceeds on Cu0 sites that form during the first stages of the reaction. Nevertheless, the redox state and morphology of the starting oxides affect the reaction outcome in terms of selectivity and the chemical nature of the products.

In other recent studies, the morphological evolution of electrochemically deposited cuprous oxide nanocubes during CO2ER has been monitored using liquid cell transmission electron microscopy27 (in situ TEM). Interestingly, pulsed electrolysis conditions were chosen to continuously regenerate Cu(I) species, and the combination of Cu(100) domains, defect sites, and surface Cu2O was found to enhance the CO2ER reaction pathway leading to C2+ products.23 Similarly, a high C2 Faradaic efficiency (80%) was reported toward acetic acid and ethanol using a 3D dendritic Cu/CuO composite electrochemically synthesized on a Cu foil, upon application of a reduction potential as low as −0.4 V vs RHE.28

Efforts in the replacement of bulk metal cathodes with lightweight and low-cost carbonaceous supports, especially large-area gas diffusion electrodes, represent a step forward in the overall design of artificial photosynthesis-oriented systems. The first attempts involving Cu-based catalysts for CO2ER date back to the 90s and were motivated by the increased mass fixation of CO2 provided by the porous and 3D structure of gas diffusion layers (GDLs).29,30 In more recent years, only a few works on liquid-phase CO2ER catalysts based on Cu0 NPs, either alone31 or deposited on CNTs,32,33 and loaded on GDLs have been reported, where formic and acetic acids were the major products. However, to date the role of the Cu morphology and redox state on CO2ER product distribution has not yet been investigated on GDL supports, which is a crucial step to pave the way for the use of such low-cost and high-surface area supports in the challenging electrochemical reaction.

In this work, a set of nanostructured Cu catalysts have been loaded over a carbonaceous GDL (Toray carbon paper, shortly named CP) by simple and highly reproducible procedures, including first the Cu0 electrodeposition and then a chemical or electrochemical oxidative treatment, leading to the formation of a pristine Cu0/CP and its oxidized forms Cu2O-Cu0/CP, CuO-Cu0/CP, and Cu(OH)2-Cu0/CP. The materials have been exploited for CO2ER in liquid phase to produce acetic acid as the main product. The catalytic performances in terms of productivity and selectivity toward formic and acetic acids depend on the electrocatalyst morphology and the Cu redox state, which can be finely tuned modifying the experimental parameters of the oxidative treatment.

2. MATERIALS

2.1. Chemicals and Materials. Toray carbon paper (TGP-H-60), Nafion membrane N-115 (0.125 mm-thick, ≥0.90 meq/g exchange capacity), copper tape, and ammonium nitrate were purchased from Alfa Aesar. Copper nitrate trihydrate (Cu(NO3)2·3H2O), sulfuric acid (96.0–97.2%), potassium hydroxide, hydrogen peroxide, ammonium persulfate, potassium hydrogen carbonate, and phenol were purchased from Sigma-Aldrich. Pure carbon dioxide (99.9%) was acquired from Rivoira S.l. Deuterium oxide (99.96%) was purchased from Eur isotop. Gas sampling bags (Tedlar bags) were obtained from Supelco. All chemicals were of reagent grade or higher.

2.2. Apparatus. The copper electrodeposition and the electrochemical characterizations of the support were carried out in a conventional three-electrode cell, and all potentials were controlled by a potentiostat (CH Instrument 660 C). Toray carbon paper (4 cm2-sized) was used as the working electrode, while a saturated calomel electrode (SCE) and a Pt gauze were used as the reference and counter electrodes, respectively. With the exception of the electrocatalytic tests, all the potentials were quoted vs SCE. The morphology and the structure of the catalysts were investigated by scanning electron microscopy (SEM) using an E-SEM Zeiss EVO 50 series instrument. Energy-dispersive X-ray spectroscopy (EDS) measurements were...
performed with an Oxford INCA system equipped with a 30 mm² silicon drift detector. Raman spectra were recorded with a Renishaw Raman RM1000 equipped with a Leica DMLM optical microscope. The excitation wavelength came from an argon laser adjusted at 514.5 nm with an output power of 25 mW. This power was reduced as needed by neutral density filters in order to prevent sample damage. The crystalline phases and the dimensions of the crystallites were analyzed by X-ray diffraction spectroscopy (XRD) using a PW1050/81 diffractometer (Phillips/Malvern, Royston, UK) coupled with a graphite monochromator in the diffracted beam and controlled by a PW1710 unit (Cu Kα, λ = 0.15418 nm). The electrochemical CO₂ reduction reaction tests were carried out in a low-volume, two-compartment electrochemical (H-type) cell (Pine Reasearch Instrumentation, Inc.). The GDLs coated with the catalysts were used as the working electrodes, and a Ag/AgCl (KCl sat.) electrode and a Pt gauze were used as the reference and counter electrodes, respectively. ¹H NMR spectra were recorded by means of an Inova 600 spectrometer (600 MHz, D₂O) coupled with a triple resonance probe. Gaseous products were analyzed by a Thermo Focus GC with a carbon molecular sieve column (CARBOSPHERE 80/100 6 ft × 0.32 cm) equipped with a TCD detector.

2.3. Pretreatment of the Carbonaceous Support. A total geometrical surface area of 4 cm² of Toray carbon paper (CP) was obtained from a 19 × 19 cm foil. Before copper deposition, the carbonaceous membrane was soaked in 1 M H₂SO₄ followed by treatment in pure EtOH for well-defined times, whose optimization will be discussed in Section 3.1.2.

2.4. Electrodeposition of Cu₀ on the Carbon Paper Electrode. The electrodeposition of metal copper was performed in a conventional three-electrode cell by applying a constant potential of −0.4 V vs SCE for 500 s. The electrolytic solution was composed of 0.15 M Cu(NO₃)₂·3H₂O and 0.70 M NH₄NO₃. The pretreated CP was employed as the working electrode and coupled with a Pt gauze and the SCE as counter and reference electrodes, respectively. The as-obtained Cu₀ thin layer was carefully washed in monodistilled H₂O and then dried at 333 K overnight. The pristine metal copper electrocatalyst was named Cu₀/CP.

2.5. Oxidative Treatment of the Pristine Cu₀/CP. Once the Cu₀ electrocatalyst had been obtained, the copper oxidation state was tuned by means of simple chemical and electrochemical methods. Three different oxidized catalysts were obtained as follows.

(a) The CuₓO−Cu₀/CP catalyst was achieved by applying a constant potential of −0.4 V vs SCE for 720 s in 0.5 M KOH aqueous solution, obtaining a thin CuₓO film over the metal copper deposit.

(b) The Cu(OH)₂−Cu₀/CP and the Cu₂O−Cu₀/CP catalysts were chemically obtained. Indeed, Cuₓ/Cu₀/CP electrodes were soaked for 20 min in two different 2.5 M NaOH solutions adding either 0.1 M (NH₄)₂S₂O₈ in order to obtain the copper hydroxide or 0.1 M H₂O₂ to obtain a mixture of copper oxides, along with a copper deposit.

The morphologies and the structures of each electrocatalyst were thoroughly investigated by SEM-EDS, Raman, and X-ray diffraction analyses.

2.6. Liquid-Phase CO₂ Electroreduction Tests. The electrochemical reduction tests were performed in an H-type cell equipped with a proton exchange polymeric membrane (Nafion-115). Both sides of the cell were filled with 0.3 M KHCO₃, as the electrolyte. The cathodic side was presaturated with a constant flow of pure CO₂ (20 mL min⁻¹) for 30 min, which was maintained at 5 mL min⁻¹ during the reaction. A Ag/AgCl electrode and a Pt gauze were located in the anodic part of the cell, while the electrocatalyst was placed in the cathodic side (Figure 1). All the applied potentials used for the CO₂ ER are referred to the reversible hydrogen electrode (RHE).

The liquid products were analyzed by a quantitative ¹H NMR analysis, adding phenol as the internal standard and deuterium oxide to provide an internal lock signal. Gaseous products were collected in a gas sampling bag for all the duration of the electrocatalytic tests and analyzed by gas chromatography. When different reaction times were investigated, a new catalyst was electrodeposited each time.

3. RESULTS AND DISCUSSION

3.1. Carbonaceous Support for the Catalysts. 3.1.1. Morphology and XRD Characterization. Toray carbon paper was chosen as the conductive support on which the electrocatalyst was deposited. Mostly known for its large use as a gas diffusion layer (GDL) in the proton exchange membrane fuel cells (PEMFC), the CP structure is composed of three main features. The long, thin, and randomly distributed fibers provide good electrical conductivity and act as the active phase during electrodeposition, while the void regions among them assure a good carbon dioxide transfer, in both liquid and gas phases. The third feature is ascribed to a carbon binder, which guarantees strength and durability of the fibers. The bare CP was morphologically investigated by SEM (Figure S1a). The majority of the carbon fibers can be individually visualized with an average diameter of 7 μm, but also, fibers linked together by the carbon binder can be distinguished, forming larger areas of conductive fibers. Moreover, X-ray diffraction analysis was conducted in order to evaluate the crystalline structure of the carbon support. The diffraction pattern, which is given in Figure S1b, was compared to the standard ICSD. The reflection pattern was assigned to hexagonal graphite (ICSD no. 98-005-3029), which shows the most intense signal at 26.5° and another one that is less intense at 43.6°, corresponding to the (002) and (004) planes, according to Miller indices, respectively.

3.1.2. Optimization of the Carbon Gas Diffusion Layer Pretreatment. The cleaning procedure of the catalyst support is a fundamental step to obtain a reproducible and well-adherent coating. In this study, an already reported pretreatment was adopted that guaranteed (i) an effective cleaning of the carbon support from organic and inorganic impurities without altering the structure of the fibers, (ii) an increase in its hydrophilicity, (iii) good reproducibility of the electrodeposition process, and (iv) short preparation times. H₂SO₄ (1 M) and pure EtOH sequential baths were used for this purpose. We tested five treatment combinations in both acidic and alcoholic environments upon variation of the residence times. The effect of the pretreatment was investigated by Raman spectroscopy, and attention was focused on the two typical bands of the carbonaceous materials: the “D band” (1354 cm⁻¹), which represents the vibration of the defects inside the crystalline structure, and the “G band” (1581 cm⁻¹) that is related to the
C–C vibrations of graphite (Figure S2a). The ratio between them, known as the "R value," indicates the degree of ordered graphite crystallites inside the structure. Figure S2b displays the R value (I_D/I_G) related to the five samples coming from the different pretreatments. Overall, the Toray carbon paper after acidic and alcoholic treatments for 2 and 1 h, respectively, reported the lowest ratio (R value = 0.19), which means that the disordered carbonaceous components on the surface were reduced, thus favoring a greater contribution of the crystalline structure. Therefore, the aforementioned pretreatment was chosen for the GDL support. Finally, an electrochemically active surface area (ECSA) evaluation was performed using cyclic voltammetry (CV) for the chosen pretreated CP. A potential range of 100 mV centered around the open-circuit value was set, and several CVs with different scan rates were recorded, obtaining an average ECSA of 39 cm² and a roughness factor (RF) of 9.75. The increase in the ECSA in comparison to the geometrical area is fully explained by the fiber structure of the paper.

3.2. Copper-Based Catalysts Electrodeposited on the GDL. 3.2.1. Synthesis and Morphological/Structural Characterization of the Cu⁰/CP Electrocatayst. The electrodeposition of a thin layer of metal copper over the carbonaceous membrane was performed by applying a constant voltage for a well-defined time, as reported in Figure 2a. The reaction that occurred at the CP, used as the working electrode, is the following

\[
\text{Cu(NO}_3\text{)}_2 + 2e^- \rightarrow \text{Cu}^0 + 2\text{NO}_3^-
\]

The progress of the reaction was investigated recording SEM images at different times in order to follow the evolution of the Cu⁰ deposition over the carbon fiber. The nucleation of sub-micrometric copper particles started after 1.5 s, as shown in Figure 2b, and a maximum current density of 21.7 mA cm⁻² was recorded. Afterward, the current density began to decrease and reached a constant value of 15.6 mA cm⁻² due to the diffusion processes, while the particles were still growing (Figure S3) until the end of the deposition process.

In addition, as the particles started to increase their dimensions, it can be noticed that crystal agglomerates rather than large single crystals mainly formed. After 500 s, a homogeneous covering of the conducting fibers was achieved, and the metal copper deposit showed good adhesion to the 3D carbonaceous substrate (Figure S4). Moreover, the reproducibility of the Cu mass was investigated, obtaining an average of 10 ± 1 mg for 20 electrodepositions. Interestingly, the Cu⁰ coating occurred only on the superficial fibers, leaving the underlying ones completely uncovered. Due to the poor interactions of carbon dioxide with the bare fibers, this fact could facilitate the passage of CO₂ through the 3D...
According to Dongare et al., the current density recorded with the N2-saturated electrolyte is ascribable to the contribution of the carbon dioxide reduction. The Faradaic efficiency of the catalyst electroreduction itself coupled with the hydrogen evolution reaction (HER) takes place is slightly anticipated in the presence of CO2 (from −1.1 to −0.95 V vs SCE). According to Dongare et al., the current density recorded with the N2-saturated electrolyte is ascribable to the catalyst electroreduction itself coupled with the hydrogen evolution. On the other hand, the maximum current density recorded after the onset potential shown in the red curve (CO2-saturated electrolyte) was slightly higher, probably due to the contribution of the carbon dioxide reduction. The Faradaic peaks observed at −0.42 (red curve) and −0.55 V vs SCE (black curve) can be associated with the reduction of surface Cu oxides locally formed during the LSV characterization. Due to the lower pH of the CO2-saturated electrolyte, the reduction of oxidized Cu species was facilitated, occurring at a less cathodic potential. Figure 3b shows several chronoamperometric curves recorded at a Cu0/CP electrode during CO2ER in different potential-controlled electrolysis conditions with CO2 continuously flowing. Herein, the current densities rapidly decreased from the very first step of the reaction until a plateau value, which linearly increased as the applied potential was more cathodic ($R^2 = 0.998$). Differently from other literature reports, the potential increase did not lead to significant disturbance in the recorded current.

3.3. Oxidative Functionalization and Characterization of the Cu0/CP-Derived Electrocatalysts. Once the Cu0-based gas diffusion electrode was fully investigated, simple chemical and electrochemical methods were employed to selectively change the copper redox couple acting as the active phase. A first functionalization of the Cu0 layer was achieved by performing a potential-controlled electrochemical oxidation at −0.4 V vs SCE in 0.5 M KOH as the electrolyte for 12 min, leading to the CuOx-Cu0/CP catalyst. The potential for electrochemical oxidation was selected on the basis of a cyclic voltammetry study on the pristine Cu0/CP catalyst, as shown in Figure S6, which was in accordance with the literature. Two further copper oxide-based electrocatalysts were obtained by simple chemical oxidations, soaking the pristine Cu0/CP layer in strong oxidizing mixtures for a relatively short time. The species produced on the pristine metal copper layer were Cu(OH)2 and CuO, depending on the chemical nature of the oxidant. A comprehensive morphological and structural characterization of the copper-derived samples is reported in Figure 4.

If compared to the metal copper deposit described above, then the SEM images highlight different morphologies, which are typical of the oxidized species, while the corresponding Raman and X-ray diffraction patterns reveal the nature of each catalyst. In Figure 4, the red-evidenced catalyst was confirmed to be CuOx-Cu0/CP by XRD, Raman analyses, and SEM images revealing fibers fully covered by thin and dense flaps with a thickness lower than 100 nm and arranged perpendicularly with

Figure 4. (a) SEM images, (b) Raman spectra, and (c) X-ray diffraction patterns of the oxidized copper species derived from the pristine Cu0/CP: CuOx-Cu0/CP (red line), CuO−Cu0/CP (green line), and Cu(OH)2−Cu0/CP (blue line).
respect to the fibers. The electrochemical oxidative treatment has converted the spherical metal copper particles into a segmented nanosheet-like coating characterized by a strong increase in both the surface area of the active material and the void spaces. The X-ray diffraction pattern showed typical reflections at 36.6, 42.5, and 61.7° (ICSD no. 98-006-2467), referred to the crystalline planes (111), (002), and (220), respectively, likewise of the pristine Cu0/CP catalyst crystal structure. Moreover, compared to the crystallite size of the metal copper deposit, the average dimension of the cuprite crystallites was only 3 nm larger, with an average value of around 39 nm. In addition, the Raman spectrum displayed typical peaks of cuprous oxide at 147, 212, and 641 cm−1 with much lower intensities than those recorded for the other catalysts, thus suggesting the presence of a very thin oxidized layer over the pristine metal copper. The catalyst obtained by chemical oxidation in a mixture of 30% v/v H2O2 and NaOH (Figure 4, green-evidenced) had a uniform shuttle-like microstructure, similarly to the CuO nanoparticles obtained by a sol–gel method developed by Narsinga Rao et al.44 Contrary to their nanoparticles, which were 800 nm in length with a thickness of 130 nm, the length of a single shuttle was quite higher at around 8−7 μm with a thickness lower than 600 nm. The Raman analysis displayed the presence of a broad peak at 288 cm−1 with a shoulder at 334 cm−1, ascribable to the CuO species,45 in addition to the typical peaks of Cu2O. Moreover, optical microscope images (not shown) revealed the coexistence of weak blue thin fibers growing from the copper deposit along the fibers, similar to the one that will be described just below. Those singular components were hard to analyze because of their smaller sizes, but it was also possible to confirm the presence of Cu(OH)2 species (see below). Therefore, by employing a relatively mild chemical oxidation, we were able to obtain an electrocatalyst in which the surface of the metal copper layer was fully covered by all three copper oxide species Cu2O, CuO, and Cu(OH)2. Due to the small crystallite size of the hydroxide and the bulk nature of the technique, the XRD pattern recorded for the Cu0/CP electrode, but its presence is due to a side reaction (HER) with no carbon dioxide involved, and a fair amount of carbon monoxide is present. On the other hand, considering the liquid phase, H2 NMR analysis revealed the presence of formic acid as the main product and of a small amount of acetic acid. The Faradaic efficiency (FE) was ~80% for hydrogen evolution and ~4%, 7%, and 1% for CO, HCOO−, and CH3COO− production, respectively.

When using the other three catalysts based on the oxidized species of copper, the same products and an analogous behavior were pointed out, but the FE for the hydrogen evolution decreased to a value of ~50%, regardless of the employed catalyst. The product distribution as a function of the applied potentials is reported in Figure S7. It is worth noting that −1.1 V was reported as the potential value at which the highest amount of CO2 reduction products can be obtained, as stated by Kuhl et al., and, in our case, the best formic acid productivity and selectivity were obtained. Differently, acetic acid was the major product obtained at the least cathodic potential (−0.4 V vs RHE). Moreover, in the latter condition, the lowest amount of H2 as a side product was detected. Therefore, these two potentials leading to the most interesting reaction outcomes were investigated during the following tests with the oxidized electrocatalysts. The resulting product distributions are shown in Figure S5, and the recorded current densities are reported in Table S1.

Comparing the results at the two voltages, the first evidence that stands out is the different product distribution. Indeed, by setting the most cathodic potential (−1.1 V vs RHE), hydrogen is observed as the main gaseous product, especially for the pristine Cu0/CP electrode, but its presence is due to a side reaction (HER) with no carbon dioxide involved, and a fair amount of carbon monoxide is present. On the other hand, considering the liquid phase, H2 NMR analysis revealed the presence of formic acid as the main product and of a small amount of acetic acid. The Faradaic efficiency (FE) was ~80% for hydrogen evolution and ~4%, 7%, and 1% for CO, HCOO−, and CH3COO− production, respectively.

Weighing the other three catalysts based on the oxidized species of copper, the same products and an analogous behavior were pointed out, but the FE for the hydrogen evolution decreased to a value of ~50%, regardless of the employed catalyst.

The product distribution as a function of the applied potentials is reported in Figure S7. It is worth noting that −1.1 V was reported as the potential value at which the highest amount of CO2 reduction products can be obtained, as stated by Kuhl et al., and, in our case, the best formic acid productivity and selectivity were obtained. Differently, acetic acid was the major product obtained at the least cathodic potential (−0.4 V vs RHE). Moreover, in the latter condition, the lowest amount of H2 as a side product was detected. Therefore, these two potentials leading to the most interesting reaction outcomes were investigated during the following tests with the oxidized electrocatalysts. The resulting product distributions are shown in Figure S5, and the recorded current densities are reported in Table S1.

Comparing the results at the two voltages, the first evidence that stands out is the different product distribution. Indeed, by setting the most cathodic potential (−1.1 V vs RHE), hydrogen is observed as the main gaseous product, especially for the pristine Cu0/CP electrode, but its presence is due to a side reaction (HER) with no carbon dioxide involved, and a fair amount of carbon monoxide is present. On the other hand, considering the liquid phase, H2 NMR analysis revealed the presence of formic acid as the main product and of a small amount of acetic acid. The Faradaic efficiency (FE) was ~80% for hydrogen evolution and ~4%, 7%, and 1% for CO, HCOO−, and CH3COO− production, respectively.

When using the other three catalysts based on the oxidized species of copper, the same products and an analogous behavior were pointed out, but the FE for the hydrogen evolution decreased to a value of ~50%, regardless of the employed catalyst.

The product distribution as a function of the applied potentials is reported in Figure S7. It is worth noting that −1.1 V was reported as the potential value at which the highest amount of CO2 reduction products can be obtained, as stated by Kuhl et al., and, in our case, the best formic acid productivity and selectivity were obtained. Differently, acetic acid was the major product obtained at the least cathodic potential (−0.4 V vs RHE). Moreover, in the latter condition, the lowest amount of H2 as a side product was detected. Therefore, these two potentials leading to the most interesting reaction outcomes were investigated during the following tests with the oxidized electrocatalysts. The resulting product distributions are shown in Figure S5, and the recorded current densities are reported in Table S1.

Comparing the results at the two voltages, the first evidence that stands out is the different product distribution. Indeed, by setting the most cathodic potential (−1.1 V vs RHE), hydrogen is observed as the main gaseous product, especially for the pristine Cu0/CP electrode, but its presence is due to a side reaction (HER) with no carbon dioxide involved, and a fair amount of carbon monoxide is present. On the other hand, considering the liquid phase, H2 NMR analysis revealed the presence of formic acid as the main product and of a small amount of acetic acid. The Faradaic efficiency (FE) was ~80% for hydrogen evolution and ~4%, 7%, and 1% for CO, HCOO−, and CH3COO− production, respectively.

When using the other three catalysts based on the oxidized species of copper, the same products and an analogous behavior were pointed out, but the FE for the hydrogen evolution decreased to a value of ~50%, regardless of the employed catalyst.

The product distribution as a function of the applied potentials is reported in Figure S7. It is worth noting that −1.1 V was reported as the potential value at which the highest amount of CO2 reduction products can be obtained, as stated by Kuhl et al., and, in our case, the best formic acid productivity and selectivity were obtained. Differently, acetic acid was the major product obtained at the least cathodic potential (−0.4 V vs RHE). Moreover, in the latter condition, the lowest amount of H2 as a side product was detected. Therefore, these two potentials leading to the most interesting reaction outcomes were investigated during the following tests with the oxidized electrocatalysts. The resulting product distributions are shown in Figure S5, and the recorded current densities are reported in Table S1.

Comparing the results at the two voltages, the first evidence that stands out is the different product distribution. Indeed, by setting the most cathodic potential (−1.1 V vs RHE), hydrogen is observed as the main gaseous product, especially for the pristine Cu0/CP electrode, but its presence is due to a side reaction (HER) with no carbon dioxide involved, and a fair amount of carbon monoxide is present. On the other hand, considering the liquid phase, H2 NMR analysis revealed the presence of formic acid as the main product and of a small amount of acetic acid. The Faradaic efficiency (FE) was ~80% for hydrogen evolution and ~4%, 7%, and 1% for CO, HCOO−, and CH3COO− production, respectively.

When using the other three catalysts based on the oxidized species of copper, the same products and an analogous behavior were pointed out, but the FE for the hydrogen evolution decreased to a value of ~50%, regardless of the employed catalyst.
the greatest amount of acetate was the Cu$_2$O-Cu$^0$/CP. In particular, the produced micromoles were three times higher than the values obtained with the other two electrocatalysts based on oxidized species of copper, and meanwhile, the hydrogen evolution was the lowest. In such a case, the FE was $\sim$8 and 76% for hydrogen and acetate production, respectively, so confirming the good selectivity of the catalyst for CO$_2$ electroreduction to CH$_3$COO$^-$. Therefore, the Cu$_2$O-Cu$^0$/CP represented the most active nanostructured catalyst by applying a relatively low cathodic potential, suitable for the application discussed in the Introduction section. Moreover, other literature reports$^{23,28}$ have found out that the copper redox couple CuI/Cu$^0$ is the most active one in favoring the C$_2$ pathway.

### 3.4.2. Cu$_2$O-Cu$^0$/CP versus Cu$^0$/CP Productivity

The results obtained with the most promising electrocatalyst at the two limit reference potentials were compared, and the productivity toward the formation of the reduced species was investigated (Figure 6). To this purpose, the micromole production of each CO$_2$, HCOO$^-$, and CH$_3$COO$^-$. All the productivities calculated at the two potentials, when working with Cu$_2$O-Cu$^0$/CP or Cu$^0$/CP electrocatalysts, are collected in Table 2 to emphasize the remarkable change of activity and selectivity passing from the reduced to the partly oxidized state of copper. In particular, at both potentials, the total productivity is higher for the Cu$_2$O-Cu$^0$/CP electrocatalyst, and the same holds also for the carbon selectivity toward acetate production, resulting in 8.2% (vs 3.1% for Cu$^0$/CP) at $-1.1$ V and practically 100% at $-0.4$ V. In the following, a tentative explanation of the better performance of the Cu$_2$O-Cu$^0$/CP than the Cu$^0$/CP electrode to produce useful chemicals will be given.

One of the first works that suggested the key role of the Cu$^+$ species in the selective reduction of CO$_2$ was published by Mistry et al.,$^{50}$ who developed Cu electrocatalysts using a plasma treatment that produced a stable cuprous oxide layer (Cu$_2$O) on the surface of a Cu foil. Also, Roberts et al.$^{51}$ reported a copper surface structured in situ using an oxidative–reductive reaction leading to Cu$_2$O, which was subsequently reduced to Cu$^0$ in a cubic structure. This catalyst displayed an exceptional ability to catalyze CO$_2$ reduction favoring C–C coupling, which leads to multcarbon products.

More recently, Zhu et al.$^{28}$ fabricated a 3D dendritic copper–cuprous oxide composite, which displayed a noticeable selectivity to C$_2$ products (acetic acid and ethanol). They attributed the optimum performance of the catalyst to the many exposed active sites in the 3D dendritic structure and to a suitable Cu$^+/Cu^0$ ratio.

Based on these pieces of evidence, the higher activity and the outstanding selectivity of the electrocatalyst Cu$_2$O-Cu$^0$/CP over Cu$^0$/CP can be attributed to the following features, considering that the CO$_2$ reduction is a surface process and its efficiency is related to the density of active sites on the catalyst:

1. One is the proper Cu$^+/Cu^0$ ratio of the catalytic active sites, determined by the electrochemical synthesis above described. In this regard, the coexistence of Cu$^+$ and Cu$^0$ species at the electrocatalyst surface was predicted to lead

![Figure 6. CO$_2$ER productivity using the Cu$_2$O-Cu$^0$/CP electrocatalyst at the two limit reference potentials.](image)
to enhanced *CO dimerization and therefore to C₂ products.⁶²

(2) Another is the structure of the Cu₂O coating, which is preserved also after the application of the cathodic potential at which the CO₂ER occurs (shown in Figure 4a), constituted of a segmented nanosheet morphology. This structure is characterized by a very high surface area of the active material with respect to the spherical metal copper particles and by many void spaces. The high surface area should guarantee a huge number of active sites produced by the reduction of the Cu₂O film to Cu⁰, presumably in the form of non-homogeneous nanoparticles with enhanced grain boundaries, which improve the catalytic performance, as already reported.⁵³

Overall, the nanosheet-like structure is likely to favor the CO₂ mass transport throughout the catalyst surface and should guarantee a vast electrochemical interface for CO₂ activation and reduction. According to the literature, CO₂ adsorbs on the active sites of the catalyst and is reduced to *CO intermediates that can be more strongly bound on the Cu surface and converted into C₂ products by *CO dimerization.⁶¹ That means that *CO dimerization is the key step in the selective reduction of CO₂ to acetic acid, and this process requires a longer residence time than the one necessary to follow the C₁ pathway, which produces HCOOH (2H⁺ and 2e⁻).

Moreover, fiber copper electrodes with a three-dimensional porous hollow structure were introduced by Kas et al.⁵⁴ to study the electrochemical reduction of CO₂, highlighting the importance of the void spaces to remarkably increase the electrocatalytic performance due to the optimal mass transport conditions for those reactions in which at least one gas-phase reactant is involved. The fact that, at the most cathodic potential, a noticeable productivity of CO and HCOOH is observed also at the Cu₂O-Cu⁰/CP electrode, despite the better selectivity for acetate with respect to the Cu⁰/CP electrode, is thus explainable. At high overpotentials, the current density increases due to the higher rate of both CO₂ reduction to CO and HER. Consequently, a significant reduction of the CO₂ concentration at the electrode surface takes place, the reactant is under diffusion control, and the formation of gaseous products limits the formation of multicarbon liquid products, like acetate, at a high concentration.

### 3.4.3. CH₃COO⁻ Production over Time Using Cu₂O-Cu⁰/CP

Although many studies have been carried out to synthesize new materials as catalysts for the electrochemical reduction of CO₂, in most of the cases, the final products are formate and carbon monoxide, which require further reduction to become useful chemicals. Focusing our attention on acetate, which is the only product apart from hydrogen when the catalyst is Cu₂O-Cu⁰/CP and the applied potential is −0.4 V vs RHE, electroreductions of different duration (15 min, 30 min, and 1 h, Figure S8) were performed to study the formation kinetics of this product. Table 3 shows the trend of the produced micromoles during the CO₂ER in 0.3 M KHCO₃ and continuous CO₂ feed.

| time (min) | Jₑ₄₉ (mA cm⁻²) | CH₃COO⁻ (μmol) |
|-----------|----------------|----------------|
| 15        | 0.60           | 0.172          |
| 30        | 0.44           | 0.344          |
| 60        | 0.46           | 3.09           |

Table 3. Acetate Production over Time at −0.4 V vs RHE in 0.3 M KHCO₃ and Continuous CO₂ Flow Using Cu₂O-Cu⁰/CP

constant ratio of the two oxidation states (activation phase of the electrocatalyst). Immediately after this step, the current stabilizes due to the adsorption/reduction of CO₂ on the active sites of the catalyst followed by a slow further decrease, related to the mass transport limitation, until a stable value is recorded. Meanwhile, also the hydrogen evolution reaction occurs since the reduction of CO₂ takes place in aqueous solution, although the applied potential is low and the catalyst displays the lowest activity toward HER with respect to the others investigated in this work. In fact, an evident hydrogen evolution started after ~200 s, which was visible to the naked eyes. Consequently, electrolysis carried out for shorter times refused the most from the initial process related to the catalyst activation and the beginning of the HER, thus resulting in a lower acetate production. By applying the potential of −0.4 V for a longer time (1 h), the acetate production remarkably increased during the second half-hour (2.75 vs 0.344 μmol) caused by a higher activity of the Cu¹/Cu⁰ centers to promote *CO dimerization. However, extending the reaction time to 3 or 5 h, the production of acetate started to decrease. The reasons could be related to a spurious phenomenon due to a possible stripping of the product by the CO₂ stream or to a poisoning of the catalyst surface induced by cathodic deposition of some metal impurities during the CO₂ER or by graphitic carbon species formed via decomposition of intermediates.⁵¹,⁵⁵,⁵⁶

### 3.4.4. Reusability and Stability of the Cu₂O-Cu⁰/CP Electrode

After 1 h of CO₂ER at −0.4 V vs RHE, the reusability of the working electrode was also studied. It was disassembled, washed with bidistilled water, dried under nitrogen, and subjected again to the same electrochemical oxidative process to convert Cu to Cu₂O, as described in the experimental section. Then, it was reused for a second reaction run under the same conditions.

The Faradaic efficiency of acetate during the second CO₂ER resulted in a slight increase (from 76 to 84%), and the productivity slightly decreased from 308 to 245 μmol g⁻¹ h⁻¹. This preliminary experiment, which needs further studies, gives evidence of the possibility to reuse the electrode after a proper activation process, and the Toray carbon paper was not damaged during the disassembling process.

The stability of the optimized catalyst was also investigated over time during a 5 h reaction upon application of −0.4 V vs RHE. The recorded current density plot is shown in Figure S9 and highlights no relevant degradation due to prolonged use, as a relative standard deviation (% RSD) of only 9% on the average Jₑ₄₉ value was obtained during the whole reaction time.

### 3.4.5. Comparison of Cu₂O-Cu⁰/CP Performance with State-of-the-Art Catalysts

In order to compare the performance of the most effective electrocatalyst described in this work, which can reduce CO₂ to liquid products, a short overview of other Cu-based catalysts reported in the literature is shown in Table 4. It was not easy to get the data since the comparison must involve similar experimental conditions, especially the use of a gas diffusion layer based on carbonaceous paper, the
Cu\textsuperscript{0} operating at Cu\textsubscript{0}/CP, maintaining an average crystallite dimension of less and simple oxidative functionalization of the pristine scalability. Herein, a fully covered 4 cm\textsuperscript{2}-sized Cu\textsubscript{0}-based chemical deposition on a carbonaceous gas dielectrocatalyst was successfully obtained by a simple electro-

Table 4. Overview of Cu-based Electro catalysts

| electrocatalyst | potential (V vs RHE) | electrolyte | products | FE (%) | productivity | reference |
|-----------------|----------------------|-------------|----------|--------|--------------|-----------|
| porous Cu\textsuperscript{0} NPs | −0.9 | 0.1 M KHCO\textsubscript{3} | formic acid | 40 | 1460 \(\mu\text{M h}^{-1}\) | 31 |
| dendritic Cu\textsuperscript{0}-Cu\textsuperscript{2}O composite | −0.7 | 0.1 M KCl | acetic acid | 28.6 | 13.2 | 28 |
| Cu\textsuperscript{0} NPs on carbon nanotubes | −1.35 | 0.5 M KHCO\textsubscript{3} | formic acid | 28.5 | 71.5 (C-based) | 33 |
| Cu\textsubscript{2}O/GDL | −0.85 | 0.5 M KHCO\textsubscript{3} | formic acid | 8.1 | 24 \(\mu\text{mol h}^{-1}\) | 57 |
| Cu\textsubscript{2}O-Cu\textsuperscript{0}/GDL | −0.4 | 0.3 M KHCO\textsubscript{3} | acetic acid | 76 | 3.09 \(\mu\text{mol h}^{-1}\) | this work |

application of a constant potential, and a continuous CO\textsubscript{2} feed delivered during the liquid-phase reaction. Furthermore, the parameters used in the literature to describe the performance of the catalysts are different and expressed in a different way.

Some observations can be done from the data in the table. When the catalyst is based on Cu NPs, the main product is formic acid unless the NPs are supported on carbon nanotubes; on the contrary, if the catalyst contains both Cu\textsuperscript{1} and Cu\textsuperscript{0} centers, then the electrocatalytic reduction activity toward acetic acid is favored but depending on the applied potential. In fact, the major selectivity of the electrochemically produced Cu\textsubscript{2}O-Cu\textsuperscript{0} operating at \(~−0.85 \text{ V vs RHE}\) is toward formate, while \(~−0.9 \text{ V vs RHE}\) is toward formic acid.

In conclusion, the possibility to easily produce nanostructured, large-area, and low-cost electrocatalysts with promising catalytic activities was demonstrated in view of industrially relevant applications.

4. CONCLUSIONS

In this research, affordable and easily synthesized nanostructured Cu-based electrodes with notable catalytic activities toward the electroreduction of carbon dioxide at ambient conditions were reported. The combination of noncritical raw materials and reproducible synthesis/activation procedures, which provides a noteworthy and selective production of added-value chemicals from a waste product, represents a good choice for the application in solar-driven chemistry\textsuperscript{2} and future scalability. Herein, a fully covered 4 cm\textsuperscript{2}-sized Cu\textsuperscript{0}-based electrocatalyst was successfully obtained by a simple electrochemical deposition on a carbonaceous gas diffusion membrane. Compared to most of the electrocatalysts employed in the state of the art,\textsuperscript{7,17,23,28} the optimized electrode has a higher geometrical surface area, thus overcoming critical steps of reproducibility and providing a scalable approach. Electrochemical and chemical methods are effective strategies to selectively tune the active phase of the pristine catalyst. Therefore, different copper redox couples were obtained by further and simple oxidative functionalization of the pristine Cu\textsuperscript{0}/CP, maintaining an average crystallite dimension of less than 40 nm. The catalytic activity of each electrocatalyst toward the liquid-phase electrochemical reduction of CO\textsubscript{2} was thoroughly investigated at two limit reference potentials. A selectivity toward the formation of acetic acid as the main liquid product was obtained by applying the lowest voltage (−0.4 V vs RHE) at the Cu\textsubscript{2}O-Cu\textsuperscript{0}/CP electrocatalyst, thus resulting in the most promising material. Indeed, in this case, the most stable current density was observed that may be due to the decrease in the parasitic hydrogen evolution reaction, in favor of useful product formation. Although the reaction mechanisms at the basis of CO\textsubscript{2}ER are still under debate, we can hypothesize that the use of the Cu\textsuperscript{1}/Cu\textsuperscript{0} couple as the active phase preferentially drives the reaction pathway toward C=C bond formation under mild reaction conditions. In conclusion, the possibility to easily produce nanostructured, large-area, and low-cost electrocatalysts with promising catalytic activities was demonstrated in view of industrially relevant applications.

ASSOCIATED CONTENT

Supporting Information
The Supporting Information is available free of charge at https://pubs.acs.org/doi/10.1021/acsami.1c18844.

Complete contact information is available at: https://pubs.acs.org/10.1021/acsami.1c18844

AUTHOR INFORMATION

Corresponding Authors

Francesco Basile — Department of Industrial Chemistry “Toso Montanari”, University of Bologna, 40136 Bologna, Italy; Email: ebasile@unibo.it
Domenica Tonelli — Department of Industrial Chemistry “Toso Montanari”, University of Bologna, 40136 Bologna, Italy; Email: domenica.tonelli@unibo.it

Authors

Martina Serafini — Department of Industrial Chemistry “Toso Montanari”, University of Bologna, 40136 Bologna, Italy; orcid.org/0000-0002-8344-9817;
Federica Mariani — Department of Industrial Chemistry “Toso Montanari”, University of Bologna, 40136 Bologna, Italy; orcid.org/0000-0001-6293-3920
Andrea Fasolini — Department of Industrial Chemistry “Toso Montanari”, University of Bologna, 40136 Bologna, Italy; orcid.org/0000-0001-7298-0528

Complete contact information is available at: https://pubs.acs.org/10.1021/acsami.1c18844
Author Contributions
The manuscript was written through contributions of all authors. All authors have given approval to the final version of the manuscript.

Funding
This work was supported by the Emilia-Romagna region POR FSE 2014/2020, the three-year plan “High Skills for Research, Transfer Technology and Entrepreneurship”, project “Power to fuels: electrocatalytic reduction of CO₂ to methane and alcohols”.

Notes
The authors declare no competing financial interest.

ACKNOWLEDGMENTS
The authors are grateful to Dr. Alessandra Petroli for the fruitful discussions and her valuable contribution to the analysis of liquid products using 1H NMR, to Dr. Fabrizio Tarterini for his relevant contribution to SEM investigations, and to Mr. Alessandro Magnani for his valuable technical support. The authors would also like to acknowledge support of the Emilia-Romagna region.

REFERENCES
(1) Centi, G.; Perathoner, S.; Winé, G.; Gangeri, M. Electro catalytic Conversion of CO₂ to Long Carbon-Chain Hydrocarbons. Green Chem. 2007, 9, 671.

(2) Bensaid, S.; Centi, G.; Garrone, E.; Perathoner, S.; Saracco, G. Towards Artificial Leaves for Solar Hydrogen and Fuels from Carbon Dioxide. ChemSusChem 2012, 5, 500–521.

(3) Genovese, C.; Ampelli, C.; Perathoner, S.; Centi, G. Electro catalytic Conversion of CO₂ to Liquid Fuels Using Nanocarbon-Based Electrodes. J. Energy Chem. 2013, 22, 202–213.

(4) Genovese, C.; Ampelli, C.; Perathoner, S.; Centi, G. Mechanism of C–C Bond Formation in the Electrocatalytic Reduction of CO₂ to Acetic Acid. A Challenging Reaction to Use Renewable Energy with Chemistry. Green Chem. 2017, 19, 2406–2415.

(5) Prajapati, A.; Singh, M. R. Assessment of Artificial Photosynthetic Systems for Integrated Carbon Capture and Conversion. ACS Sustainable Chem. Eng. 2019, 7, 5993–6003.

(6) Wang, Q.; Wanam, J.; Rodríguez-Jiménez, S.; Leung, J. J.; Kalathil, S. R.; Andrei, V.; Domen, K.; Reisner, E. Molecularly Engineered Photocatalyst Sheet for Scalable Solar Formate Production from Carbon Dioxide and Water. Nat. Energy 2020, 5, 703–710.

(7) Wang, X.; Gao, C.; Low, J.; Mao, K.; Duan, D.; Chen, S.; Ye, R.; Qiu, Y.; Ma, J.; Zheng, X.; Long, R.; Wu, X.; Song, L.; Zhu, J.; Xiong, Y. Efficient Photoelectrochemical CO₂ Conversion for Selective Acetic Acid Production. Sci. Bull. 2021, 66, 1296–1304.

(8) Gai, P.; Yu, W.; Zhao, H.; Qi, R.; Li, F.; Liu, L.; Lv, F.; Wang, S. Solar-Powered Organic Semiconductor—Bacteria Biohybrids for CO₂ Reduction into Acetic Acid. Angew. Chem., Int. Ed. 2020, 59, 7224–7239.

(9) Zhang, G.; Liu, B.; Wang, T.; Gong, J. Artificial Leaves for Solar Fuels. Chinese J. Chem. 2021, 39, 1450–1458.

(10) Amao, Y.; Shuto, N.; Furuno, K.; Obara, A.; Fuchino, Y.; Uemura, K.; Kajino, T.; Sekito, T.; Iwai, S.; Miyamoto, Y.; Matsuda, M. Artificial Leaf Device for Solar Fuel Production. Faraday Discuss. 2012, 155, 289–296.

(11) Fan, L.; Xia, C.; Yang, F.; Wang, J.; Wang, H.; Lu, Y. Strategies in Catalysts and Electrolizer Design for Electrochemical CO₂ Conversion toward C₂H₄. Products. Sci. Adv. 2020, 6, eaay3111.

(12) Pandel, J. E.; Ren, D.; Huang, Y.; Loo, N. W. X.; Hong, S. H. L.; Ye, B. S. Understanding the Heterogeneous Electro catalytic Reduction of Carbon Dioxide on Oxide-Derived Catalysts. ChemElectroChem 2018, 5, 219–237.

(13) Nitopi, S.; Bertheussen, E.; Scott, S. B.; Liu, X.; Engstfeld, A. K.; Horch, S.; Seger, B.; Stephens, I. E. L.; Chan, K.; Hahn, C.; Nørskov, J.; Jaramillo, T. F.; Chorkendorff, I. Progress and Perspectives of Electrochemical CO₂ Reduction on Copper in Aqueous Electrolyte. Chem. Res. 2019, 119, 7610–7672.

(14) Hori, Y.; Kikuchi, K.; Suzuki, S. Production of CO and CH₄ in Electrochemical Reduction of CO₂ at Metal Electrodes in Aqueous Hydrogen Carbonate Solution. Chem. Lett. 1985, 14, 1695–1698.

(15) Xue, Y.; Guo, Y.; Cui, H.; Zhou, Z. Catalyst Design for Electrochemical Reduction of CO₂ to Multicarbon Products. Small Methods 2021, 5, 2100736.

(16) Horii, Y. Electrochemical CO₂ Reduction on Metal Electrodes. In Modern Aspects of Electrochemistry; Springer New York: New York, NY, 2008; pp. 89–189, DOI: 10.1007/978-0-387-49489-0_3.

(17) Rahaman, M.; Dutta, A.; Zanetti, A.; Broekmann, P. Electrochemical Reduction of CO₂ into Multicarbon Alcohols on Activated Cu Mesh Catalysts: An Identical Location (IL) Study. ACS Catal. 2017, 7, 7946–7956.

(18) de Lucas-Consuegra, A.; Serrano-Ruiz, J.; Gutiérrez-Guerra, N.; Valverde, J. Low-Temperature Electrochemical Reduction of CO₂ to Liquid Fuels: Effect of the Cu Particle Size. Catalysts 2018, 8, 340.

(19) Liu, S.; Huang, S. Size Effects and Active Sites of Cu Nanoparticle Catalysts for CO₂ Electroreduction. Appl. Surf. Sci. 2019, 475, 20–27.

(20) Cui, H.; Guo, Y.; Guo, L.; Wang, L.; Zhou, Z.; Peng, Z. Heteroatom-Doped Carbon Materials and Their Composites as Electrocatalysts for CO₂ Reduction. J. Mater. Chem. A 2018, 6, 18782–18793.

(21) Duan, Y.-X.; Meng, F.-L.; Liu, K.-H.; Yi, S.-S.; Li, S.-J.; Yan, J.-M.; Jiang, Q. Amorphizing of Cu Nanoparticles toward Highly Efficient and Robust Electrocatalyst for CO₂ Reduction to Liquid Fuels with High Faradaic Efficiencies. Adv. Mater. 2018, 30, 1706194.

(22) Kas, R.; Kortlever, R.; Milbrat, A.; Koper, M. T. M.; Mul, G.; Baltrusaitis, J. Electrochemical CO₂ Reduction on CuO₄-Derived Copper Nanoparticles: Controlling the Catalytic Selectivity of Hydrocarbons. Phys. Chem. Chem. Phys. 2014, 16, 12194–12201.

(23) Arán-Ais, R. M.; Scholten, F.; Kunze, S.; Rizo, R.; Roldan Cuenya, B. The Role of in Situ Generated Morphological Motifs and Cu(I) Species in C₂H₂ Product Selectivity during CO₂ Pulsed Electroreduction. Nat. Energy 2020, 5, 317–325.

(24) Li, C. W.; Kanan, M. W. CO₂ Reduction at Low Overpotential on Cu Electrodes Resulting from the Reduction of Thick Cu₂O Films. J. Am. Chem. Soc. 2012, 134, 7231–7234.

(25) Chi, D.; Yang, H.; Du, Y.; Li, T.; Sui, G.; Wang, H.; Lu, J. Morphology-Controlled CuO Nanoparticles for Electroreduction of CO₂ to Ethanol. RSC Adv. 2014, 4, 37329–37332.

(26) Giri, S. D.; Mahajani, S. M.; Suresh, A. K.; Sarkar, A. Electrochemical Reduction of CO₂ on Activated Copper: Influence of Surface Area. Mater. Res. Bull. 2020, 123, 110702.

(27) Arán-Ais, R. M.; Rizo, R.; Grosse, P.; Algara-Siller, G.; Demblé, K.; Plodinec, M.; Lunkenbein, T.; Chee, S. W.; Cuenya, B. R. Imaging Electrochemicaly Synthesized CuO Cubes and Their Morphological Evolution under Conditions Relevant to CO₂ Electroreduction. Nat. Commun. 2020, 11, 3489.

(28) Zhu, Q.; Sun, X.; Yang, D.; Ma, J.; Kang, X.; Zheng, L.; Zhang, J.; Wu, Z.; Han, B. Carbon Dioxide Electroreduction to C₂ Products over Copper-Cuprous Oxide Derived from Electrosynthesized Copper Complex. Nat. Commun. 2019, 10, 3851.

(29) Cook, R. L.; MacDuff, R. C.; Sammells, A. F. High Rate Gas Phase CO₂ Reduction to Ethylene and Methane Using Gas Diffusion Electrodes. J. Electrochem. Soc. 1990, 137, 607–608.

(30) Ikeda, S.; Ito, T.; Azuma, K.; Ito, K.; Noda, H. Electrochemical Mass Reduction of Carbon Dioxide Using Cu-Loaded Gas Diffusion Electrodes I. Preparation of Electrode and Reduction Products. Denki Kagaku oyobi Kagyo Butsuri Kagaku 1995, 63, 303–309.

(31) Dongare, S.; Singh, N.; Bhunia, H. Electrocatalytic Reduction of CO₂ to Useful Chemicals on Copper Nanoparticles. Appl. Surf. Sci. 2021, 537, 148020.

(32) Ampelli, C.; Genovese, C.; Marepally, B. C.; Papanikolaou, G.; Perathoner, S.; Centi, G. Electrocatalytic Conversion of CO₂ to Produce Solar Fuels in Electrolyte or Electrolyte-Less Configurations ofPEC Cells. Faraday Discuss. 2015, 183, 125–145.
(33) Marepally, B. C.; Ampelli, C.; Genovese, C.; Tavella, F.; Veyre, L.; Quadrelli, E. A.; Perathoner, S.; Centi, G. Role of Small Cu Nanoparticles in the Behaviour of Nanocarbon-Based Electrodes for the Electrocatalytic Reduction of CO2. J. CO2 Util. 2017, 21, 534−542.

(34) Zamel, N.; Li, X.; Shen, J. Correlation for the Effective Gas Diffusion Coefficient in Carbon Paper Diffusion Media. Energy Fuels 2009, 23, 6070−6078.

(35) Hofmann, H. U.; Wilm, D. Über Die Kristallstruktur Von Kohlenstoff. Z. Elektrochem. 1936, 42, 504−522.

(36) Ma, X.; Li, X.; Jagadale, A. D.; Hao, X.; Abudula, A.; Guan, G. Fabrication of Cu(OH)2@NiFe-Layered Double Hydroxide Catalyst Array for Electrochemical Water Splitting. Int. J. Hydrogen Energy 2016, 41, 14553−14561.

(37) Liu, J.; Tian, Y.; Chen, Y.; Liang, J.; Zhang, L.; Fong, H. A Surface Treatment Technique of Electrochemical Oxidation to Simultaneously Improve the Interfacial Bonding Strength and the Tensile Strength of PAN-Based Carbon Fibers. Mater. Chem. Phys. 2010, 122, 548−555.

(38) Gruijić, D.; Pesić, B. Electrodeposition of Copper: The Nucleation Mechanisms. Electrochim. Acta 2002, 47, 2901−2912.

(39) Young, J. F. T. XXVIII. The Crystal Structure of Various Heusler Alloys by the Use of X-Rays. London, Edinburgh, Dublin Philos. Mag. J. Sci. 1923, 46, 291−305.

(40) Zhong, H.; Fujii, K.; Nakano, Y.; Jin, F. Effect of CO2 Bubbling into Aqueous Solutions Used for Electrochemical Reduction of CO2 for Energy Conversion and Storage. J. Phys. Chem. C 2015, 119, 55−61.

(41) Giri, S. D.; Sarkar, A. Electrochemical Study of Bulk and Monolayer Copper in Alkaline Solution. J. Electrochem. Soc. 2016, 163, H252−H259.

(42) Zav’yalova, A. A.; Imamov, R. M.; Pinsker, Z. G. Production and Investigation of Thin Films of CuO. Kristallografiya 1964, 9, 857. [Sov.Phys.Crystallog. 9, 724 (1964)]

(43) Deng, Y.; Handoko, A. D.; Du, Y.; Xi, S.; Yeo, B. S. In Situ Raman Spectroscopy of Copper and Copper Oxide Surfaces during Electrochemical Oxygen Evolution Reaction: Identification of Cu III Oxides as Catalytically Active Species. ACS Catal. 2016, 6, 2473−2481.

(44) Narsinga Rao, G.; Yao, Y. D.; Chen, J. W. Superparamagnetic Behavior of Antiferromagnetic CuO Nanoparticles. IEEE Trans. Magn. 2005, 41, 3409−3411.

(45) Rashad, M.; Rüising, M.; Berth, G.; Lischka, K.; Pawlis, A. CuO and Co3O4 Nanoparticles: Synthesis, Characterizations, and Raman Spectroscopy. J. Nanomater. 2013, 2013, 1−6.

(46) Wang, J.; Zhu, L.; Ji, L.; Chen, Z. Preparation of Nanostructured Cu(OH)2 and CuO Electrocatalysts for Water Oxidation by Electrophoresis Deposition. J. Mater. Res. 2018, 33, 581−589.

(47) Oswald, H. R.; Reller, A.; Schmalle, H. W.; Dubler, E. Structure of Copper(II) Hydroxide, Cu(OH)2. Acta Crystallogr. Sect. C Cryst. Struct. Commun. 1990, 46, 2279−2284.

(48) Zhao, X.; Du, L.; You, B.; Sun, Y. Integrated Design for Electrocatalytic Carbon Dioxide Reduction. Catal. Sci. Technol. 2020, 10, 2711−2720.

(49) Kuhl, K. P.; Cave, E. R.; Abram, D. N.; Jaramillo, T. F. New Insights into the Electrochemical Reduction of Carbon Dioxide on Metallic Copper Surfaces. Energy Environ. Sci. 2012, 5, 7050.

(50) Mistry, H.; Varela, A. S.; Bonifacio, C. S.; Zegkinoglou, I.; Sinev, I.; Choi, Y.-W.; Kissingler, K.; Stach, E. A.; Yang, J. C.; Strasser, P.; Cuenya, B. R. Highly Selective Plasma-Activated Copper Catalysts for Carbon Dioxide Reduction to Ethylene. Nat. Commun. 2016, 7, 12123.

(51) Roberts, F. S.; Kuhl, K. P.; Nilsson, A. High Selectivity for Ethylene from Carbon Dioxide Reduction over Copper Nanocube Electro catalysts. Am. Ethnol. 2015, 127, 5265−5271.

(52) Xiao, H.; Goddard, W. A.; Cheng, T.; Liu, Y. Cu Metal Embedded in Oxidized Matrix Catalyst to Promote CO2 Activation and CO Dimerization for Electrochemical Reduction of CO2. Proc. Natl. Acad. Sci. 2017, 104, 7024−7029.

(53) Kwon, Y.; Lum, Y.; Clark, E. L.; Ager, J. W.; Bell, A. T. CO2 Electroreduction with Enhanced Ethylene and Ethanol Selectivity by Nanostructuring Polycrystalline Copper. ChemElectroChem 2016, 3, 1012−1019.