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Selective electrochemical reduction of carbon dioxide to ethanol via a relay catalytic platform†

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Efficient electroreduction of carbon dioxide (CO₂) to ethanol is of great importance, but remains a challenge because it involves the transfer of multiple proton–electron pairs and carbon–carbon coupling. Herein, we report a CoO-anchored N-doped carbon material composed of mesoporous carbon (MC) and carbon nanotubes (CNT) as a catalyst for CO₂ electroreduction. The faradaic efficiencies of ethanol and current density reached 60.1% and 5.1 mA cm⁻², respectively. Moreover, the selectivity for ethanol products was extremely high among the products produced from CO₂. A proposed mechanism is discussed in which the MC–CNT/Co catalyst provides a relay catalytic platform, where CoO catalyzes the formation of CO* intermediates which spill over to MC–CNT for carbon–carbon coupling to form ethanol. The high selectivity for ethanol is attributed mainly to the highly selective carbon–carbon coupling active sites on MC–CNT.

Introduction

The excessive use of fossil fuels and deforestation result in the continuous increase in carbon dioxide (CO₂) concentration in the atmosphere and ocean, which is a crucial issue for human beings.¹ The efficient conversion of CO₂ to fuels and high-value-added chemicals can promote the development of the cyclic utilization of carbon resources and a reduction in CO₂ emissions.^{2,3} Electrocatalytic reduction of CO₂ is a promising route due to its mild operating conditions (ambient pressure and temperature) and modular reaction systems, which can be easily controlled by adjusting electrochemical parameters, making them easy to scale-up for applications.^{4–6} Ethanol is a widely used intermediate in chemical synthesis and a liquid fuel with a high energy density (26.8 MJ kg⁻¹).⁷ Direct electroreduction of CO₂ to ethanol is of great importance. However, it is challenging because the process involves not only the transfer of multiple proton–electron pairs but also carbon–carbon coupling with a high overpotential.

Cu is documented to be a key component for multicarbon chemical production. Unfortunately, most Cu-based electrodes are more favorable for ethylene over ethanol. On plasma-activated Cu nanocube electrocatalysts, the faradaic efficiencies (FE) of ethylene and ethanol were 45% and 22%, respectively.⁸ Electrodeposited

CuAg alloy films showed a good electroreduction performance with an FE of 60% for ethylene and 25% for ethanol.⁹ An oxide-derived Cu/carbon catalyst exhibited an FE of 24% for ethanol production.¹⁰ On bimetallic Ag/Cu electrocatalysts, ethanol could be formed with an FE of 41%.¹¹ Hierarchically structured Cu dendrite electrodes with a superhydrophobic surface attained an FE of 56% for ethylene and 17% for ethanol production.¹² An FE for ethanol of 41% on porphyrin-based Fe/Cu was reported and the ratio of ethanol to ethylene was 1.0 under optimized operating conditions.¹³ Copper–nitrogen-doped carbon materials achieved aqueous CO₂ electroreduction to ethanol at an FE of 55% under optimized conditions.¹⁴ On copper–cuprous oxide electrodes, ethanol and acetic acid could be produced, while the main product was acetic acid.¹⁵

In addition to Cu-based electrocatalysts, Ag-, Fe- and carbon-material-based catalysts have also been reported for the electroreduction of CO₂ to ethanol. Ag-anchored N-doped graphene/carbon foam could efficiently catalyze the conversion of CO₂ to ethanol with an FE of 82.1–85.2%, but the current density was only 0.35 mA cm⁻².¹⁶ Fe₂P₂S₆ nanosheets were used as an efficient electrocatalyst for the highly selective electroreduction of CO₂ to methanol and ethanol, with an FE of 23.1% for ethanol.¹⁷ An Ru(II) polypyridyl carbene complex immobilized on N-doped porous carbon displayed an FE of 27.5% for ethanol with very low current density.¹⁸ Carbon material electrodes can catalyze the conversion of CO₂ to ethanol with high selectivity but their current density is very low. For example, an FE of 77% for ethanol was obtained on nitrogen-doped ordered cylindrical mesoporous carbon with a current density <1 mA cm⁻². Moreover, the selectivity of ethanol among the products produced from CO₂ was nearly 100%.¹⁹ Further study showed that medium micropores embedded in the

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channel walls of nitrogen-doped ordered mesoporous carbon could promote ethanol production from CO₂ electroreduction, and the FE for ethanol reached 78%.²⁰

A literature survey shows that the electrocatalysts that have been reported exhibit high current density and low selectivity for ethanol or high selectivity with low current density. It is obvious that more efforts should be made to develop highly efficient electrocatalysts for the electroreduction of CO₂ to ethanol. CO is an important intermediate for the generation of C₂⁻ products from CO₂ electroreduction. It has been reported that high-concentration CO could be generated on the doping metal sites and spill over to the Cu sites for carbon–carbon coupling before further reduction to C₂⁻ products on Cu-based bimetallic electrodes. Rich local CO intermediates can effectively promote the next carbon–carbon coupling process in CO₂ electroreduction. Moreover, the highly selective carbon–carbon coupling active sites on a catalyst are crucial for the electroreduction of CO₂ to ethanol with high selectivity.

Cobalt-based materials can be used as catalysts for the electroreduction of CO₂ to CO. On semiconducting Co₃O₄ nanofibers, CO could be formed with an FE of 65%.²¹ Cobalt phthalocyanine (CoPc) molecules anchored on carbon nanotubes exhibit an FE of 95% for CO production.²² Inspired by the high selectivity by carbon materials for CO₂ electroreduction to ethanol and the good performance of cobalt-based materials for CO₂ electroreduction to CO, in this work, we designed a catalyst MC–CNT/Co, in which CoO was anchored in N-doped carbon material composed of mesoporous carbon (MC) and carbon nanotubes (CNT), for CO₂ electroreduction, which provided a relay catalytic platform, where CoO catalyzes the formation of *CO intermediates that spill over to MC–CNT for carbon–carbon coupling to ethanol. The selectivity for ethanol and acetaldehyde could reach nearly 100% among the CO₂ electroreduction products at –0.2 to –0.45 V and an FE of 60.1% was achieved for ethanol under a low overpotential (–0.32 V vs. RHE) with a current density of 5.1 mA cm^{–2}. The Co species, ordered mesoporous structure and CNT had an excellent synergistic effect for promoting CO₂ electroreduction to ethanol.

Experimental

Reagents

Toray carbon paper (CP, TGP-H-60, 19 × 19 cm), Nafion D-521 dispersion (5% w/w in water and 1-propanol, ≥0.92 meg per g exchange capacity) and Nafion N-117 membrane (0.180 mm thick, ≥0.90 meg per g exchange capacity) were purchased from Alfa Aesar China. 3-(Aminopropyl)trimethoxysilane (APTMS), 2-methylimidazole (purity >98%), tetraethyl orthosilicate (TEOS) (purity >99%), and cobalt nitrate hexahydrate (Co(NO₃)₂) were purchased from Beijing Innochem Company, Pluronic P123 (average Mn ~ 5800) was obtained from Sigma. Hydrochloric acid, sodium hydroxide, and toluene (purity >99%) were provided by the Beijing Chemical Company.

Synthesis of the electrode materials

Preparation of mesoporous silica (MS). MS was prepared according to work reported previously.²³ Typically, 4.0 g of

Pluronic P123 was dissolved in 50 mL of water and stirred for 5 h at room temperature. The mixture was added to 120 mL of 2 M hydrochloric acid solution and stirred for 2 h, and then TEOS (8.5 g) was added. The solution was aged at 35 and 100 °C for 24 h, respectively. Finally, the solid products were obtained by filtration. Afterwards, the material was washed with deionized water and air-dried overnight. The P123 was removed by calcining in air at 550 °C for 5 h.

Preparation of MS with amino-functional composites (MS–NH₂). 1 g of as-prepared MS was dispersed in 30 mL of toluene by ultrasonication, and then into it was dropped 2 mL of aminopropyl-methoxysilane (APTMS). The mixture was refluxed at 80 °C with continuous stirring for 24 h. The resulting functionalized MS composites were recovered by filtration followed by washing with toluene several times and then dried at 60 °C under vacuum for 12 h. The products obtained were denoted MS–NH₂.

Synthesis of MC–CNT/Co and MC–CNT. To obtain MC–CNT/Co, 2-methylimidazole was used as carbon precursor. In a typical process, a Co²⁺/MS composite template and powdered 2-methylimidazole were mixed and placed in a quartz boat inside a quartz tube. Ar was introduced as a protective and carrier gas. Then, the furnace was heated to 700 °C at a rate of 5 °C min^{–1} and kept at that temperature for 3 h to obtain the Co/MS/carbon composite. After removal of silica by 10 wt% NaOH aqueous solution, MC–CNT/Co was obtained. Moreover, MC–CNT could be obtained by etching the metal particles in MC–CNT/Co with hydrochloric acid solution (2 M) for 72 h.

Materials characterization

The morphologies of the electrodes were characterized using a S4800 scanning electron microscope (SEM) and a JEOL-2100F transmission electron microscope (TEM) operated at 200 kV. Powder X-ray diffraction (XRD) patterns were performed on an X-ray diffractometer (Model D/MAX2500, Rigaku) with Cu–K α radiation. X-ray photoelectron spectroscopy (XPS) analysis was conducted on a Thermo Scientific ESCA Lab 250Xi using 200 W monochromatic Al K α radiation. The content of Co in the catalysts was determined by inductively coupled plasma optical emission spectroscopy (ICP-OES, Vista-MPX). The N₂ adsorption/desorption isotherms were determined using a Micromeritics TriStar 3020 instrument at –196 °C. The Brunauer–Emmett–Teller (BET) method was employed to calculate the specific surface area, while the Barrett–Joyner–Halenda (BJH) method was applied to analyze the pore size distribution using the adsorption branch of the isotherm. The FTIR spectra were collected at a resolution of 4 cm^{–1} on a Bruker Vector 27 spectrophotometer in the 400–4000 cm^{–1} region. The IR spectra of the samples were measured by the conventional KBr pellet method. Thermogravimetric analysis (Pyris 1 TGA) was performed under an Ar flow from 20 to 800 °C at a heating rate of 10 °C min^{–1}.

Electrochemical measurements

An electrochemical workstation (CHI 660E, Shanghai CH Instruments Co., China) was used for all CO₂ reduction

experiments. Typically, a 10 mg sample and 20 μL of Nafion solution (5 wt%) were dispersed in 1 mL of deuterium acetonitrile by sonicating for 30 min to form a homogeneous ink. Then, the dispersion was loaded onto carbon paper with dimensions of 1 cm \times 1 cm. For CO_2 reduction experiments, cyclic voltammogram (CV) measurements were performed in an H-type cell using 0.5 M KHCO_3 (60 mL, pH = 7.8) as the electrolyte with a typical three-electrode setup, which contained a working electrode, a platinum gauze auxiliary electrode, and an Ag/AgCl reference electrode. All the measured potentials in this work are cited with respect to the RHE using the following conversion: $E_{\text{RHE}} (\text{V}) = E_{\text{Ag/AgCl}} (\text{V}) + 0.197 \text{ V} + (0.059 \text{ V} \times \text{pH})$. Before electrolysis, the electrolyte was purged with Ar or CO_2 gas for at least 30 min. The CV measurement in gas-saturated electrolyte was conducted at a sweep rate of 5 mV s^{-1} in the potential between -0.6 V and -1.4 V (vs. Ag/AgCl). The process was kept under constant magnetic stirring. The electrolysis experiments were conducted at 25°C in a typical H-type cell. It consisted of a cathode, an anode (platinum gauze auxiliary electrode) and an Ag/AgCl reference electrode. In the experiments, the cathode and anode compartments were separated by a Nafion-117 proton exchange membrane. KHCO_3 aqueous solution (0.5 M) was used as the anodic electrolyte. Then, potentiostatic electrochemical reduction of CO_2 was carried out with CO_2 bubbling (5 mL min^{-1}). The reduction of acetaldehyde was also performed over an MC-CNT/Co electrode at -0.32 V (vs. RHE) in 0.5 M KHCO_3 electrolysis solution with 0.05 M acetaldehyde. The electrolysis experiment was conducted at 25°C in an H-type cell with a Nafion-117 proton exchange membrane and lasted for 2 h. The liquid products were analyzed by ^1H NMR (Bruker Avance III 400 HD spectrometer) in acetonitrile- d_3 with dimethyl sulfoxide (DMSO) as an internal standard. The gaseous product was collected using a gas bag and analyzed by a gas chromatograph (GC, HP 4890D), which was equipped with a TCD detector using helium as the internal standard. Each data point was an average of the measurements collected from at least three separate NMR or GC analyses. Each prepared catalyst was used only once for CO_2 reduction at a chosen potential.

Calculations of faradaic efficiencies of gaseous and liquid products

Gaseous products. $\text{FE} = \text{moles of products per second} / \text{theoretical moles equivalent per second}$

Based on the GC peak areas and calibration curves, the V % of H_2 can be obtained. The amount of moles of H_2 (or CO) per second could be calculated from the flow rate of the gas and the V % of H_2 . The theoretical moles per second was obtained from the current density.

Liquid products. In NMR spectra, DMSO was used as the internal standard, and the relative peak area of $\text{CH}_3\text{CH}_2\text{OH}$ could be calculated.

The number of electrons required to produce liquid product during the entire CO_2 electroreduction reaction is:

$$N = C \times V \times N_A \times n_e$$

(V : the volume of catholyte; N_A : Avogadro constant; n_e : transfer electron number)

We can obtain the Q_{total} from the chronoamperogram and the total number of electrons by measuring:

$$N_{\text{total}} = Q_{\text{total}}/e$$

The faradaic efficiency (FE) of the liquid product is:

$$\text{FE} = N/N_{\text{total}} \times 100\%$$

Results and discussion

Ordered MS was prepared using TEOS as the precursor and P123 as the template. Amino groups were grafted onto MS using APTMS as an amination reagent, which was proved by the N-H vibration peaks at 1560 and 3120 cm^{-1} in the FTIR spectra (Fig. S1 \dagger). The mesoporous structure of MS was retained after amino group grafting (Fig. S2 \dagger).^{24–26} MC-CNT/Co was synthesized in three steps, as shown in Fig. 1a. First, the cobalt species

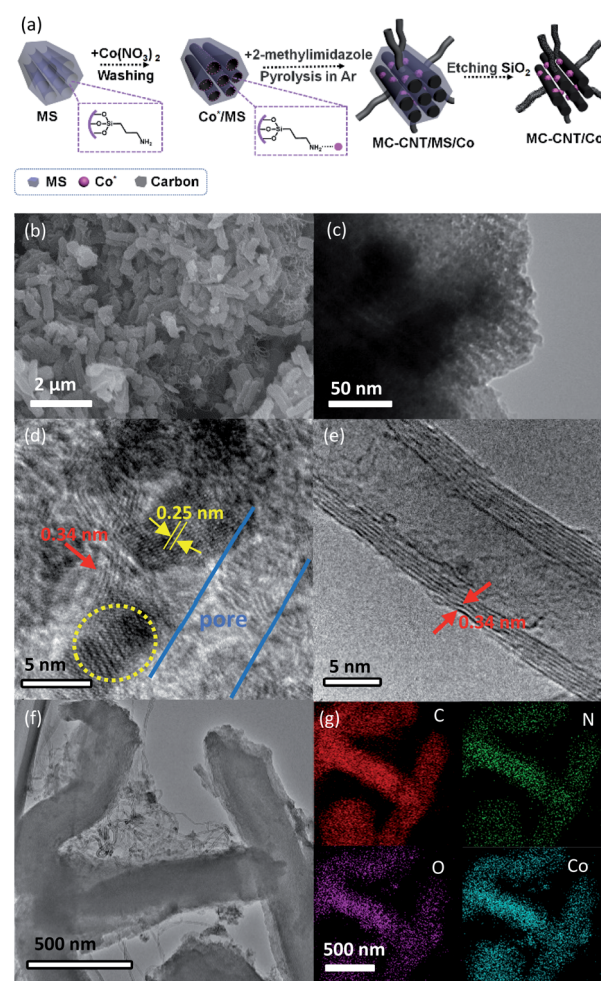


Fig. 1 (a) Schematic illustration of the synthesis process, (b) SEM, (c and f) TEM, and (d and e) high-resolution TEM images, and (g) the corresponding elemental mapping of C, O, Co and N elements of MC-CNT/Co.

was adsorbed onto MS to obtain Co²⁺/MS owing to the electrostatic force between the amino and cobalt ion. Second, MC-CNT/Co/MS was obtained by thermal treatment of a mixture composed of Co²⁺/MS composite and powdered 2-methylimidazole under an Ar atmosphere, in which 2-methylimidazole was used as the carbon and nitrogen precursor. Subsequently, MC-CNT/Co/MS was treated with 10 wt% NaOH aqueous solution to remove silica, resulting in the MC-CNT/Co. SEM and TEM images show that MC-CNT/Co has a 3D network structure formed by CNT connected to rod-like mesoporous carbon (Fig. 1b and f). As shown in Fig. 1d and e, the graphene layers with a lattice distance of 0.34 nm could be clearly seen. It has been reported that Co could catalyze the formation of CNT.²⁷ Ordered mesoporous structures observed from TEM images of MC-CNT/Co showed that the pores of the MS template were replicated (Fig. 1c). Additionally, cobalt nanoparticles (diameter: 3–5 nm) were also detected, and their lattice spacing was 0.25 nm, which was assigned to CoO (111) (Fig. 1d).^{28,29} The content of cobalt in MC-CNT/Co was 13.20 wt%, which was analyzed by inductively coupled plasma optical emission spectroscopy (ICP-OES). The corresponding elemental mapping reveals its elemental distribution, which shows a uniform N, Co distribution in the carbon framework (Fig. 1g).

Fig. 2a shows the XRD patterns of MC-CNT/Co. The broad diffraction peaks around 23.3° and 26.0° correspond to the amorphous and graphitized carbon, respectively. Three peaks at 36.8°, 43.1° and 63.4° are indexed to CoO (111), CoO (200) and CoO (220) (CoO-75-0418).^{30,31} As depicted in Fig. 2b, MC-CNT/Co exhibited a type IV adsorption-desorption isotherm and type H3 hysteresis loops, showing the existence of mesopores. The specific surface area calculated using the BET method was 303 m² g⁻¹. The BJH method was used to calculate the pore-size distribution from the adsorption branch of the nitrogen isotherm and the average pore size was 4.4 nm (Fig. 2c).

The electronic states of different elements in MC-CNT/Co were investigated by XPS. As shown in Fig. S4,† Co, N, C and O elements were observed. The high-resolution Co 2p_{3/2} XPS spectrum of MC-CNT/Co (Fig. 2d) can be deconvoluted into

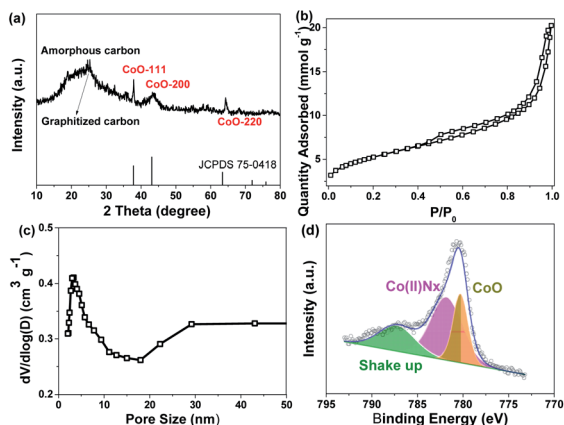


Fig. 2 (a) XRD patterns, (b) N₂ adsorption and desorption curves, (c) pore-size distribution and (d) XPS spectrum of the Co 2p of MC-CNT/Co.

three sets of peaks at 780.6 and 779.3 eV and a satellite at 785.5 eV, which are attributed to Co(II)N_x and Co(II)O.^{32–34} The O 1s signal (Fig. S5a†) was deconvoluted into three oxygen configurations, carboxyl oxygen (532 eV), hydroxyl oxygen (534 eV) in amorphous hydrogenated carbon and Co–O (529 eV).³³ Deconvolution of the N 1s peak (Fig. S5b†) revealed two peaks that can be assigned to pyridinic-N (398.5 eV) and pyrrolic-N (400.0 eV), respectively.³⁴

The CV tests over MC-CNT/Co were performed in Ar-saturated and CO₂-saturated 0.5 M KHCO₃ electrode aqueous solution (pH 7.8), respectively. In this work, the applied potential is referenced to the reversible hydrogen electrode (RHE) and the current density is calculated from the geometric surface area. As shown in Fig. 3a, an obvious reduction peak can be observed at *ca.* –0.3 V (*vs.* RHE) in a CO₂-saturated electrolyte, which shows the reduction of CO₂. Controlled-potential CO₂ electrolysis experiments were performed in 0.5 M KHCO₃ aqueous electrolyte solution using a typical H type cell. The onset potential of MC-CNT/Co was as low as –0.25 V (*vs.* RHE), which is more positive than that of many other carbon-based electrodes.^{19,35–38} More importantly, ethanol and acetaldehyde (trace) were the only liquid products detected by NMR spectroscopy. H₂ was the only gaseous product in the range of –0.20 to –0.45 V (*vs.* RHE), showing that the selectivity for C₂ liquid products from CO₂ is 100% on the MC-CNT/Co catalyst. The current density and FE for ethanol and acetaldehyde were 5.1 mA cm⁻² and 70.7% (60.1% for ethanol and 10.0% for

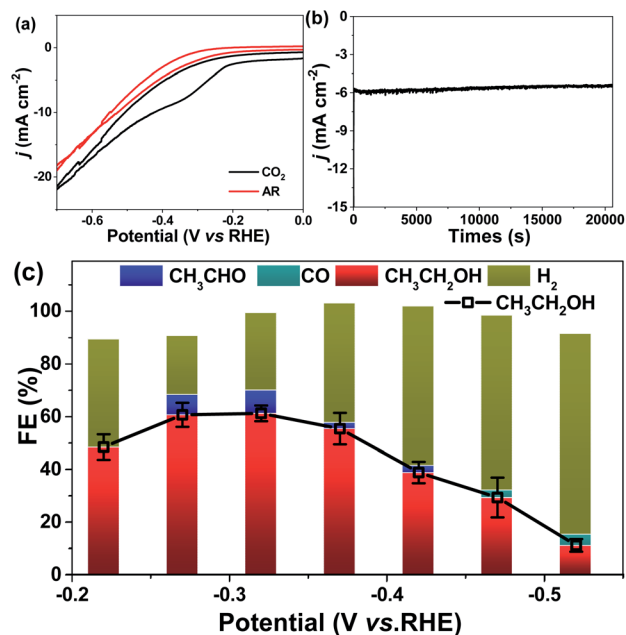


Fig. 3 (a) Cyclic voltammogram curves in the potential range from 0 V to –0.7 V (*vs.* RHE) at a sweep rate of 5 mV s⁻¹ in 0.5 M Ar-saturated and CO₂-saturated 0.5 M KHCO₃ on MC-CNT/Co electrodes, (b) total current density *versus* time in 0.5 M KHCO₃ under a CO₂ atmosphere on electrodes at –0.32 V (*vs.* RHE) and (c) FE for ethanol, aldehyde, carbon monoxide and hydrogen production obtained from the electrochemical reduction of CO₂ for 6 h on MC-CNT/Co electrode at –0.2 to –0.6 V (*vs.* RHE).

acetaldehyde), respectively (Fig. 3b and c). Apart from ethanol, a small amount of CO was detected at -0.48 V and -0.52 V (*versus* RHE), which is a very important intermediate for C_2 products, as in previously reported results.³⁹ MC-CNT/Co exhibited a higher FE for ethanol than Cu-based catalysts and a higher current density than carbon-material-based electrodes. In addition, the MC-CNT/Co catalyst showed long-term stability in electrolysis, which was known from the fact that both current density and FE did not change considerably with electrolysis time over 20 h at -0.32 V (*vs.* RHE), as shown in Fig. S6.†

From the product distribution shown in Fig. 3c, we deduced that acetaldehyde was a possible intermediate in the formation of ethanol and this can be confirmed by the electroreduction of acetaldehyde. The reduction of acetaldehyde was performed over MC-CNT/Co at -0.32 V (*vs.* RHE) in 0.5 M $KHCO_3$ electrolysis solution (Fig. S7.†). It was found that ethanol was produced with 50% conversion and an FE of 98%.

To confirm that the product was derived from CO_2 reduction, blank experiments using Ar to replace CO_2 in the electrolysis at -0.3 V *vs.* RHE (Fig. S8.†) were conducted and no product was formed after electrolysis for 6 h. The reaction in isotope-labeled $^{13}CO_2$ and $^{12}CO_2$ saturated 0.5 M $KHCO_3$ electrolyte solution at -0.32 V *vs.* RHE was also studied over MC-CNT/Co. The product was analyzed by 1H NMR after electrolysis for 1 h. In the 1H NMR spectra in ESI Fig. S9,† we can only see the H signal of the $^{13}CH_3$ group on ethanol, which is split into two peaks by coupling with the ^{13}C atom. The results indicate that carbon atoms in the product were from CO_2 rather than from other C-based chemicals in the reaction system.^{18,40}

To unveil the reasons behind the good performance of MC-CNT/Co, MC-CNT and pure CoO_x were checked as reference samples for the electroreduction of CO_2 . MC-CNT was prepared by etching CoO in MC-CNT/Co with hydrochloric acid solution (2 M) for 72 h, and characterized by TEM, XRD and XPS. Fig. S10.† shows CNT and the mesopore structure of MC-CNT, but no detectable Co or CoO nanoparticles or subnano-clusters were observed. Moreover, no signals of Co or CoO were detected in the XRD or XPS spectra (Fig. S11.†). These results indicate that Co or CoO was removed by HCl. A pure CoO_x sample was also

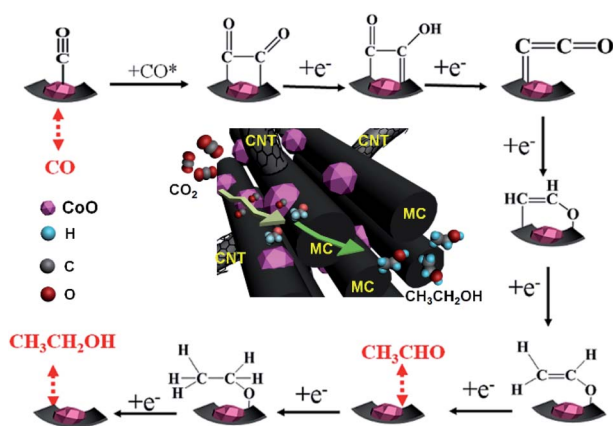
prepared by annealing $Co(NO_3)_2$ under an Ar atmosphere (Fig. S12.†). CO_2 electrolysis measurements were performed in CO_2 -saturated 0.5 M $KHCO_3$ electrolyte solution using CoO_x or MC-CNT as the catalysts and the electrolysis process was carried out in the range of -0.20 to -0.6 V (*vs.* RHE) for 2 h. On the CoO_x electrode, CO was detected with a maximum FE of 5.6% at -0.46 V (*vs.* RHE), which is an important intermediate during CO_2 electroreduction to C_2 products (Fig. S13.†). On the MC-CNT electrode, CO and ethanol as result of carbon-carbon coupling were produced with FE of 0.98% and 48.1% at -0.46 V (*vs.* RHE), respectively. The pyridinic-N and pyrrolic-N with high electron densities could facilitate the formation of ethanol, which is consistent with results reported in the literature.^{19,20} Furthermore, because of the excellent conductivity of CNT, the current density was measured to be 3.2 mA cm^{-2} , which is higher than that reported for a carbon electrode. Recent studies have also proved the vital role of CNT in facilitating the electrocatalytic reduction of CO_2 .^{19,35,41,42} These experimental results suggest that MC-CNT/Co creates a relay catalytic platform where CoO_x and MC-CNT cooperate very well to catalyze the formation of C_2 products *via* CO.

Electrochemical impedance spectroscopy (EIS) was performed at -0.56 V *vs.* RHE in CO_2 -saturated 0.5 M $KHCO_3$ solution to measure the charge transfer resistance (R_{ct}) for MC-CNT/Co and MC-CNT. As shown in Fig. S14,† the R_{ct} value of MC-CNT/Co was as low as 0.28 ohm, lower than that of MC-CNT (0.47 ohm), implying that electrons could be more easily transported on MC-CNT/Co.

According to the experimental data and reported results,^{35,43,44} a possible mechanistic pathway for the electrocatalytic production of ethanol over MC-CNT/Co is proposed and is shown in Scheme 1. One electron transfers to a CO_2 molecule to form an intermediate CO_2^{*-} anion radical. Subsequently, the obtained CO_2^{*-} reacts with a proton-electron pair and forms CO^* intermediates, which is mainly attributed to the CoO species. CO^* is the key intermediate for the dimerization³⁵ and it spills over to MC-CNT for carbon-carbon coupling to form *COCHO or *COCO intermediates. Pyrrolic-N and pyridinic-N in MC-CNT/Co and its mesoporous structure are both favorable for stabilizing CO^* intermediates and promoting the carbon-carbon coupling reaction to ethanol.^{19,20} Furthermore, CoO species and MC-CNT are intimate in MC-CNT/Co because it is synthesized by one-pot pyrolysis, which is beneficial for the spillover of CO^* intermediates to MC-CNT.

Conclusions

In summary, the MC-CNT/Co catalyst designed in this work shows excellent performance for CO_2 electroreduction to ethanol. The FE of ethanol could reach 60.1% at -0.32 V *versus* the reversible hydrogen electrode. The ordered mesoporous structure and CoO particles cooperate very well to promote the reaction. CoO catalyzes the formation of CO^* intermediates that spill over to MC-CNT for carbon-carbon coupling to ethanol. Pyrrolic-N and pyridinic-N in MC-CNT/Co and its mesoporous structure are both favorable for stabilizing CO^* intermediates and promoting the carbon-carbon coupling reaction. The high



Scheme 1 Proposed mechanism for the electrochemical reduction of CO_2 to ethanol on an MC-CNT/Co electrode.

selectivity for ethanol was attributed mainly to the highly selective carbon-carbon coupling active sites on MC-CNT. We believe that a relay catalytical platform strategy can also be used for designing highly efficient catalysts for electroreduction of CO₂ to C₂⁻ products.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

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Notes and references

- 1 X. Duan, J. Xu, Z. Wei, J. Ma, S. Guo, S. Wang, H. Liu and S. Dou, *Adv. Mater.*, 2017, **29**, 1701784.
- 2 A. Vasileff, X. Zhi, C. Xu, L. Ge, Y. Jiao, Y. Zheng and S.-Z. Qiao, *ACS Catal.*, 2019, **9**, 9411–9417.
- 3 L. Lin, H. Li, C. Yan, H. Li, R. Si, M. Li, J. Xiao, G. Wang and X. Bao, *Adv. Mater.*, 2019, **31**, 1903470.
- 4 H. Yang, Y. W. Hu, J. J. Chen, M. S. Balogun, P. P. Fang, S. Zhang, J. Chen and Y. Tong, *Adv. Energy Mater.*, 2019, **9**, 1901396.
- 5 D. Gao, I. Sinev, F. Scholten, R. M. Arán-Ais, N. J. Divins, K. Kvashnina, J. Timoshenko and B. R. Cuenya, *Angew. Chem., Int. Ed.*, 2019, **18**, 17047–17053.
- 6 H. Cui, Y. Guo, L. Guo, L. Wang, Z. Zhou and Z. Peng, *J. Mater. Chem. A*, 2018, **6**, 18782–18793.
- 7 H. Pramanik and S. Basu, *Can. J. Chem. Eng.*, 2008, **85**, 781–785.
- 8 D. Gao, I. Zegkinoglou, N. J. Divins, F. Scholten, I. Sinev, P. Grosse and B. R. Cuenya, *ACS Nano*, 2017, **11**, 4825–4831.
- 9 T. T. H. Hoang, S. Verma, S. Ma, T. T. Fister, J. Timoshenko, A. I. Frenkel, P. J. A. Kenis and A. A. Gewirth, *J. Am. Chem. Soc.*, 2018, **140**, 5791–5797.
- 10 K. Zhao, Y. Liu, X. Quan, S. Chen and H. Yu, *ACS Appl. Mater. Interfaces*, 2017, **9**, 5302–5311.
- 11 Y. C. Li, Z. Wang, T. Yuan, D.-H. Nam, M. Luo, J. Wicks, B. Chen, J. Li, F. Li, F. P. G. d. Arquer, Y. Wang, C.-T. Dinh, O. Voznyy, D. Sinton and E. H. Sargent, *J. Am. Chem. Soc.*, 2019, **141**, 8584–8591.
- 12 D. Wakerley, S. Lamaison, F. Ozanam, N. Menguy, D. Mercier, P. Marcus, M. Fontecave and V. Mougél, *Nat. Mater.*, 2019, **18**, 1222–1227.
- 13 F. Li, Y. C. Li, Z. Wang, J. Li, D.-H. Nam, Y. Lum, M. Luo, X. Wang, A. Ozden, S.-F. Hung, B. Chen, Y. Wang, J. Wicks, Y. Xu, Y. Li, C. M. Gabardo, C.-T. Dinh, Y. Wang, T.-T. Zhuang, D. Sinton and E. H. Sargent, *Nat. Catal.*, 2020, **3**, 75–82.
- 14 D. Karapinar, N. T. Huan, N. R. Sahraie, J. Li, D. Wakerley, N. Touati, S. Zanna, D. Taverna, L. H. G. Tizei, A. Zitolo, F. Jaouen, V. Mougél and M. Fontecave, *Angew. Chem., Int. Ed.*, 2019, **58**, 15098–15103.
- 15 Q. Zhu, X. Sun, D. Yang, J. Ma, X. Kang, L. Zheng, J. Zhang, Z. Wu and B. Han, *Nat. Commun.*, 2019, **10**, 3851.
- 16 K. Lv, Y. Fan, Y. Zhu, Y. Yuan, J. Wang, Y. Zhu and Q. Zhang, *J. Mater. Chem. A*, 2018, **6**, 5025–5031.
- 17 L. Ji, L. Chang, Y. Zhang, S. Mou, T. Wang, Y. Luo, Z. Wang and X. Sun, *ACS Catal.*, 2019, **9**, 9721–9725.
- 18 Y. Liu, X. Fan, A. Nayak, Y. Wang, B. Shan, X. Quan and T. J. Meyer, *Proc. Natl. Acad. Sci. U. S. A.*, 2019, **116**, 26353–26358.
- 19 Y. Song, W. Chen, C. Zhao, S. Li, W. Wei and Y. Sun, *Angew. Chem., Int. Ed.*, 2017, **56**, 10840–10844.
- 20 Y. Song, S. Wang, W. Chen, S. Li, G. Feng, W. Wei and Y. Sun, *ChemSusChem*, 2020, **13**, 293–297.
- 21 A. Aljabour, H. Coskun, D. A. Apaydin, F. Ozel, A. Hassel, P. Stadler, N. S. Sariciftci and M. Kus, *Appl. Catal., A*, 2018, **229**, 163–170.
- 22 X. Zhang, Z. Wu, X. Zhang, L. Li, Y. Li, H. Xu, X. Li, X. Yu, Z. Zhang, Y. Liang and H. Wang, *Nat. Commun.*, 2017, **8**, 14675.
- 23 A. Chen, Y. Yu, R. Wang, Y. Yu, W. Zang, P. Tang and D. Ma, *Nanoscale*, 2015, **7**, 14686–14690.
- 24 S. H. Huang and D. H. Chen, *J. Hazard. Mater.*, 2009, **163**, 174–179.
- 25 S. Wang, K. Wang, C. Dai, H. Shi and J. Li, *Chem. Eng. J.*, 2015, **262**, 897–903.
- 26 L. You, F. Yuan and F. Ma, *J. Phys. Chem. A*, 2015, **89**, 2298–2303.
- 27 Z. Li, M. Shao, Q. Yang, Y. Tang, M. Wei, D. G. Evans and X. Duan, *Nano Energy*, 2017, **37**, 98–107.
- 28 K. M. Nam, J. H. Shim, D.-W. Han, H. S. Kwon, Y.-M. Kang, Y. Li, H. Song, W. S. Seo and J. T. Park, *Mater. Chem.*, 2010, **22**, 4446–4454.
- 29 F. D. Wu and Y. Wang, *J. Mater. Chem.*, 2011, **21**, 6636.
- 30 J. Zhao, Z. Shao and H. Zhang, *Mater. Manuf. Processes*, 2020, **35**, 195–201.
- 31 F. Wu, S. Zhang, B. Xi, Z. Feng, D. Sun, X. Ma, J. Zhang, J. Feng and S. Xiong, *Adv. Energy Mater.*, 2018, **8**, 1703242.
- 32 W. Hu, Q. Wang, S. Wu and Y. Huang, *J. Mater. Chem. A*, 2016, **4**, 16920–16927.
- 33 H.-C. Huang, I. Shown, S.-T. Chang, H.-C. Hsu, H.-Y. Du, M.-C. Kuo, K.-T. Wong, S.-F. Wang, C.-H. Wang, L.-C. Chen and K.-H. Chen, *Adv. Funct. Mater.*, 2012, **22**, 3500–3508.
- 34 H. Wang, X. Bo, A. Wang and L. Guo, *Electrochem. Commun.*, 2013, **36**, 75–79.
- 35 K. Lv, Y. Fan, Y. Zhu, Y. Yuan, J. Wang, Y. Zhu and Q. Zhang, *J. Mater. Chem. A*, 2018, **6**, 5025–5031.
- 36 N. Sreekanth, M. A. Nazrulla, T. V. Vineesh, K. Sailaja and K. L. Phani, *Chem. Commun.*, 2015, **51**, 16061–16064.
- 37 J. Wu, S. Ma, J. Sun, J. I. Gold, C. Tiwary, B. Kim, L. Zhu, N. Chopra, I. N. Odeh, R. Vajtai, A. Z. Yu, R. Luo, J. Lou, G. Ding, P. J. Kenis and P. M. Ajayan, *Nat. Commun.*, 2016, **7**, 13869.

- 38 H. Wang, Y. Chen, X. Hou, C. Ma and T. Tan, *Green Chem.*, 2016, **18**, 3250–3256.
- 39 E. Bertheussen, A. Verdaguier-Casadevall, D. Ravasio, J. H. Montoya, D. B. Trimarco, C. Roy, S. Meier, J. Wendland, J. K. Norskov, I. E. Stephens and I. Chorkendorff, *Angew. Chem., Int. Ed.*, 2016, **55**, 1450–1454.
- 40 W. M. Brooks, L. N. Moxon, J. Field, M. G. Irving and D. M. Doddrell, *Biochem. Biophys. Res. Commun.*, 1985, **128**, 107–112.
- 41 Y. Liu, Y. Zhang, K. Cheng, X. Quan, X. Fan, Y. Su, S. Chen, H. Zhao, Y. Zhang, H. Yu and M. R. Hoffmann, *Angew. Chem., Int. Ed.*, 2017, **56**, 15607–15611.
- 42 Y. Liu, S. Chen, X. Quan and H. Yu, *J. Am. Chem. Soc.*, 2015, **137**, 11631–11636.
- 43 J. Shen, R. Kortlever, R. Kas, Y. Y. Birdja, O. Diaz-Morales, Y. Kwon, I. Ledezma-Yanez, K. J. Schouten, G. Mul and M. T. Koper, *Nat. Commun.*, 2015, **6**, 8177.
- 44 I. Ledezma-Yanez, E. P. Gallent, M. T. M. Koper and F. Calle-Vallejo, *Catal. Today*, 2016, **262**, 90–94.