


**CO<sub>2</sub> Reduction** Hot Paper

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# Stabilizing \*CO<sub>2</sub> Intermediates at the Acidic Interface using Molecularly Dispersed Cobalt Phthalocyanine as Catalysts for CO<sub>2</sub> Reduction

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**Abstract:** CO<sub>2</sub> electroreduction (CO<sub>2</sub>R) operating in acidic media circumvents the problems of carbonate formation and CO<sub>2</sub> crossover in neutral/alkaline electrolyzers. Alkali cations have been universally recognized as indispensable components for acidic CO<sub>2</sub>R, while they cause the inevitable issue of salt precipitation. It is therefore desirable to realize alkali-cation-free CO<sub>2</sub>R in pure acid. However, without alkali cations, stabilizing \*CO<sub>2</sub> intermediates by catalyst itself at the acidic interface poses as a challenge. Herein, we first demonstrate that a carbon nanotube-supported molecularly dispersed cobalt phthalocyanine (CoPc@CNT) catalyst provides the Co single-atom active site with energetically localized *d* states to strengthen the adsorbate-surface interactions, which stabilizes \*CO<sub>2</sub> intermediates at the acidic interface (pH=1). As a result, we realize CO<sub>2</sub> conversion to CO in pure acid with a faradaic efficiency of 60 % at pH=2 in flow cell. Furthermore, CO<sub>2</sub> is successfully converted in cation exchanged membrane-based electrode assembly with a faradaic efficiency of 73 %. For CoPc@CNT, acidic conditions also promote the intrinsic activity of CO<sub>2</sub>R compared to alkaline conditions, since the potential-limiting step, \*CO<sub>2</sub> to \*COOH, is pH-dependent. This work provides a new understanding for the stabilization of reaction intermediates and facilitates the designs of catalysts and devices for acidic CO<sub>2</sub>R.

## Introduction

Renewable electricity-powered CO<sub>2</sub> reduction (CO<sub>2</sub>R) offers a low-carbon-footprint route to realize the conversion from CO<sub>2</sub> to valuable chemicals and fuels.<sup>[1]</sup> The neutral and alkaline electrolyzers provide locally alkaline conditions to realize CO<sub>2</sub>R with high selectivity versus hydrogen evolution reaction (HER).<sup>[2]</sup> However, under the combined effects of non-faraday reaction and electric field,<sup>[3]</sup> carbonate formation and CO<sub>2</sub> crossover lead to a mixture of O<sub>2</sub> and CO<sub>2</sub> at the anode,<sup>[4]</sup> therefore subsequent energy-intensive and costly gas separation and CO<sub>2</sub> recovery processes are required.

The acidic flow cell, where the catalyst is in contact with the acidic media, provides a promising route to suppress the carbonate formation and thus circumvent CO<sub>2</sub> crossover.<sup>[5]</sup> However, CO<sub>2</sub> reduction cannot take place in pure acid where hydrogen evolution reaction (HER) dominates, since \*CO<sub>2</sub> intermediates are difficult-to-stabilize under the condition of high proton concentration.<sup>[4,6]</sup> This problem can be overcome by the addition of alkali cations,<sup>[7]</sup> which construct a locally alkaline environment at the catalyst surface to suppress HER activities<sup>[4,8]</sup> and lower the CO<sub>2</sub> adsorption energy so as to stabilize \*CO<sub>2</sub> intermediates.<sup>[6c]</sup> To match the acidic environment<sup>[9]</sup> and eliminate the resistance of catholyte<sup>[10]</sup> in acidic flow cells for large scale application, cation exchange membrane (CEM) can be employed in membrane electrode assemblies, denoted as CMEA.<sup>[2a,9,11]</sup> Although alkali cations can suppress HER and promote

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CO<sub>2</sub>R in acidic flow cells, it has recently been proven that the salt precipitation induced by alkali cations is inevitable during the electrolysis process.<sup>[12]</sup> This problem becomes even more severe in CMEA without liquid flow at the cathode, which leads to premature device death.<sup>[2a]</sup> Up to now, no CO<sub>2</sub>R activity and 100% H<sub>2</sub> selectivity has been observed in CMEA without alkali cations, which is also attributed to the unstable \*CO<sub>2</sub> intermediates at the acidic surface. To realize a high-performance CO<sub>2</sub>R in CMEA, stabilizing \*CO<sub>2</sub> intermediates in pure acid without alkali cations is necessary. Very recently, concurrent researches reported the immobilized organic cations replacing alkali cations to enable metal-free CO<sub>2</sub> reduction in pure acid.<sup>[13]</sup> It should be noted that immobilized organic cations are drawn by the long chains, making it difficult to ensure that all catalytic sites can be adequately covered by immobilized organic cations. HER will dominate on uncovered catalysts owing to the unstable \*CO<sub>2</sub> intermediates. Moreover, as the current density increases, the surface alkalinity constructed by immobilized organic cations gradually decreases, leading to an obvious decline in CO<sub>2</sub> selectivity. To reduce the dependence of CO<sub>2</sub>R on additional cations, including metal and organic cations, stabilizing \*CO<sub>2</sub> intermediates by catalyst itself at the acidic interface is crucial but still currently lacking.

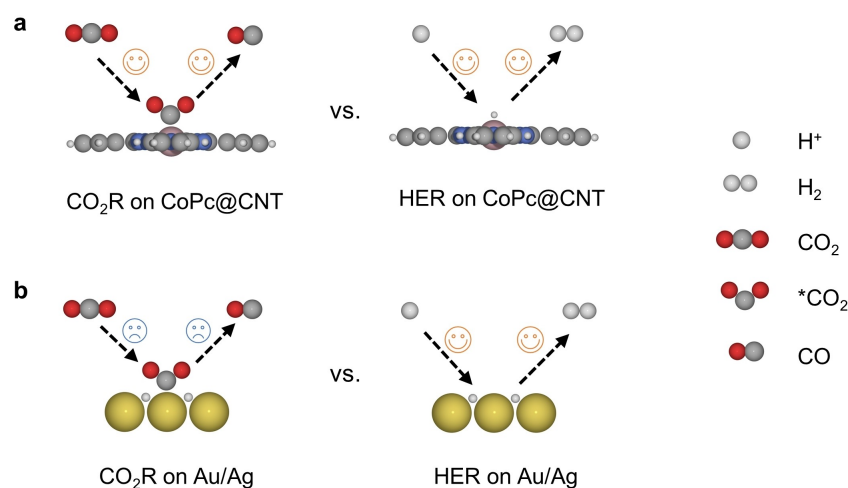
Herein, we report CO<sub>2</sub> conversion towards CO in pure acid using a carbon nanotube-supported molecularly dispersed cobalt phthalocyanine (CoPc@CNT). The Co single-atom active sites with energetically localized *d* states strengthens the adsorbate-surface interactions, which effectively stabilizes the \*CO<sub>2</sub> intermediates at the acidic interface. The key for a good CO<sub>2</sub> conversion is to render the chemisorbed \*CO<sub>2</sub> intermediate as stable, or even better more stable than H.<sup>[14]</sup> Density functional theory (DFT) calculations reveal that CO<sub>2</sub> adsorption competes with H adsorption on CoPc@CNT even at pH=1, while the CO<sub>2</sub> adsorption step is completely suppressed by the high cover-

age of H on Au and Ag (Figure 1). As a result, CoPc@CNT experimentally exhibits a faradaic efficiency (FE) of 60% for CO in acidic flow cell (pH=2) and enables CO<sub>2</sub> conversion in CMEA with a CO-FE of as high as 73%, while classic catalysts Au and Ag show no CO<sub>2</sub>R activity. In addition, the proton coupled electron transfer step, i.e., \*CO<sub>2</sub> to \*COOH, which is consistently confirmed by electrochemistry analysis with in situ Raman spectroscopy, is also proved to be the potential-limiting step on CoPc@CNT, rationalizing the lower overpotential (320 mV at 30 mA/cm<sup>2</sup>) in the acidic compared to alkaline conditions.

## Results and Discussion

Single-atom catalysts generally show stronger adsorption towards CO<sub>2</sub>R reaction intermediates compared to bulk catalysts.<sup>[15]</sup> Unlike other single-atom catalysts with complicated synthesis steps at high temperature, CoPc@CNT is easy to be prepared at mild condition and thus have a clear metal-N<sub>4</sub> coordination structure, providing an excellent model catalyst for theoretical calculations.<sup>[16]</sup> Among the metal phthalocyanines, CoPc can realize high-performance CO<sub>2</sub>R in neutral and alkaline conditions without a further structure regulation.<sup>[2c,10,17]</sup> It is anticipated that CoPc@CNT, featuring Co single-atom active sites, has the potential to enhance stabilization of CO<sub>2</sub> intermediates, enabling CO<sub>2</sub> conversion in pure acid without alkali cations. Meanwhile, Au and Ag which are known as the classic bulk metal catalysts for CO<sub>2</sub> conversion to CO in neutral and alkaline conditions, are selected as the comparison samples.<sup>[18]</sup> To predict the CO<sub>2</sub>R activity and reaction pathway of single-atom catalyst CoPc@CNT and bulk catalysts (Au and Ag) in pure acid, DFT calculations are first employed in this study.

For bulk catalysts Au and Ag, it is imperative to consider the H coverage prior to assessing the adsorption of reaction intermediates. In pure acid, since the pH are quite low, the

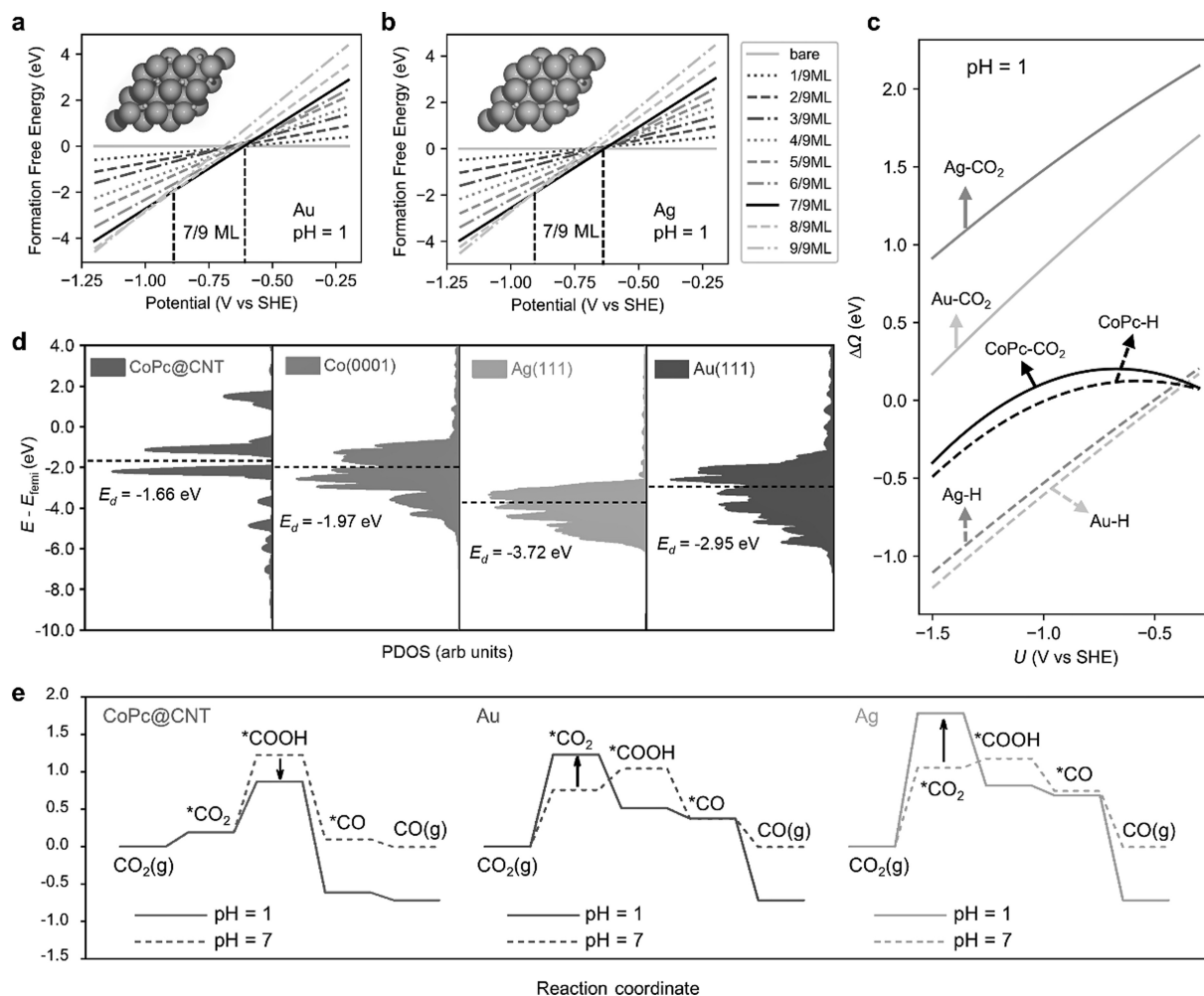


**Figure 1.** CO<sub>2</sub> adsorption competing with H adsorption on catalyst at the acidic interface. (a–b) Schematic of CO<sub>2</sub>R and HER on (a) CoPc@CNT and (b) Au/Ag in pure acid without alkali cations. The \*CO<sub>2</sub> intermediate can be stabilized at the surface of CoPc@CNT for further reduction since CO<sub>2</sub> adsorption can compete with H adsorption. As for Au and Ag, H adsorption is more facile than CO<sub>2</sub> adsorption, resulting in a destabilized \*CO<sub>2</sub> adsorption and boosted HER.

protons are much more active than in alkaline solution, which indicates that the catalyst surface will undergo quick exchange of H with solution, leading to boosted hydrogen evolution reaction (HER). The fast dynamics is usually associated with high H coverage at equilibrium. Nonetheless, previous theoretical studies mainly focus on CO<sub>2</sub>R catalytic process on the bare Au or Ag surface without H co-adsorption.<sup>[6c,15a]</sup> Our calculations, however, reveal that under acidic conditions, the catalyst surfaces are predominantly covered with H, which may strongly affect the energetics of CO<sub>2</sub>R pathway. In the case of Au(111), the H coverage will quickly reach up to 7/9 monolayer (ML) at  $-0.61$  V vs. SHE, before which the surface maintains its pristine clean state (Figure 2a). As the potential becomes more negative than  $-0.88$  V vs. SHE, the H coverage will increase consequently and reach 1 ML. A similar phenomenon can be seen on Ag(111) surface (Figure 2b). As illustrated in Figure 2a and 2b, the Au and Ag surfaces are

decorated with a high coverage of H in acidic conditions, whereas they tend to be clean at moderate potentials in alkaline media (Figure S1 and S2).

The competitive adsorption of H and CO<sub>2</sub> is crucial for the selectivity between HER and CO<sub>2</sub>R. We then evaluate the adsorption energy of H and CO<sub>2</sub> at pH=1 on H-covered Au, Ag and CoPc@CNT catalysts using grand canonical DFT, which allows explicit surface charging with an implicit electrolyte model. As illustrated in Figure 2c, for Au and Ag, the surfaces being covered with H make CO<sub>2</sub> adsorption extremely difficult, featured by high reaction energies of 1.20 eV and 1.75 eV at  $-0.7$  V vs SHE, respectively. If CO<sub>2</sub> is unable to adsorb onto the catalyst surface, the following CO<sub>2</sub>R reaction cannot take place, whereas HER will be dominating at the given condition. For CoPc@CNT, the Co atom active site can either adsorb \*CO<sub>2</sub> or \*H with roughly similar probabilities. The single Co site cannot stabilize 2\*H during optimization, and such structure quickly undergoes



**Figure 2.** Theoretical insights for CO<sub>2</sub> reduction at the acidic interface. (a–b) Formation free energy for H adsorption on (a) Au and (b) Ag as the function of potential at pH=1. (c) Adsorption free energy of \*H and \*CO<sub>2</sub> on Au, Ag and CoPc@CNT as the function of potential at pH=1. In the case of Au and Ag, the surfaces are therefore H-covered at the relevant potential for CO<sub>2</sub> reduction. Regarding the CoPc@CNT, the Co atom active site will either adsorb \*H or \*CO<sub>2</sub> in roughly similar proportion, since adsorption free energies are similar. (d) Projected density of states of *d* states of CoPc@CNT, Co(0001), Ag(111) and Au(111). Compared with metal catalysts, CoPc@CNT shows a narrower *d* state and a higher *d*-band center ( $E_d$ ). (e) The free energy diagram of CO<sub>2</sub>R pathway on CoPc@CNT and Au/Ag at  $-0.7$  V vs. SHE at pH=1 and 7.

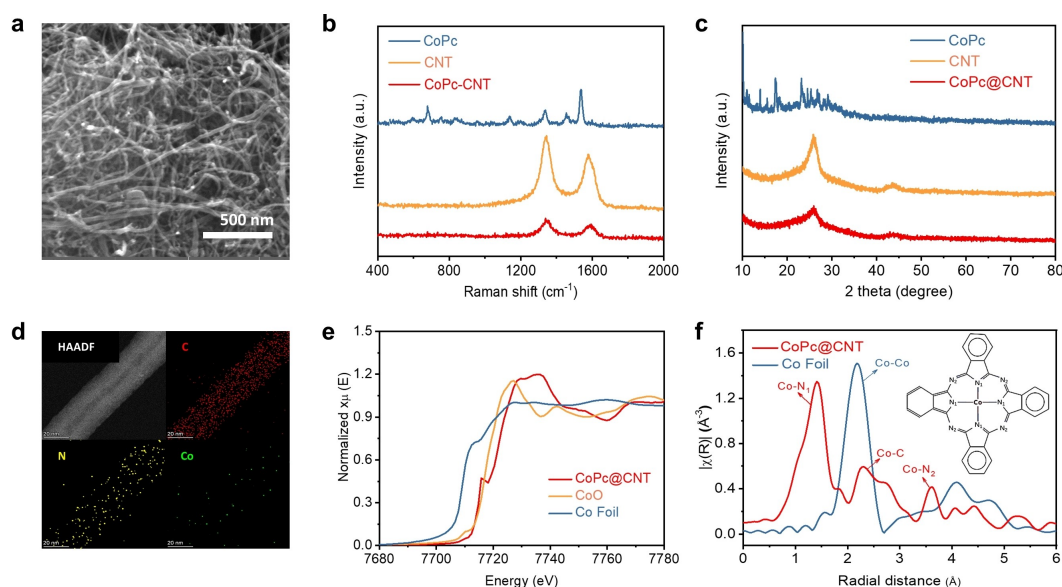
the Tafel step to form  $H_2$ . Thus, for CoPc@CNT, a fraction of the Co active sites can interact with  $CO_2$  at the potential for  $CO_2$  reduction. Based on this, on CoPc@CNT (model in Figure S3), even at  $pH=1$ ,  $CO_2$  adsorption can compete with H adsorption and exhibit a similar adsorption energy across a large potential range, indicating that CoPc@CNT is still active toward  $CO_2$  reduction, rather than being occupied by  $*H$  in acidic solution. The strong  $CO_2$  adsorption on CoPc@CNT can be attributed to the unique electronic structure of the Co single-atom site, which shows the much more energy localized  $3d$  state compared to the dispersed  $d$  band states for bulk Co (Figure 2d). According to the Newns–Anderson–Grimley model, the hybridization between the adsorbates and the  $s$  state of metal leads to a broadened adsorbate state, while such state will split into localized bonding and antibonding states when hybridizing with  $d$  state of metal. A energy localized  $d$  state will result in a stronger interaction between the adsorbate and the metal.<sup>[15a,19]</sup> In addition, since the energy separation between the metal  $d$  states and the H  $1s$  orbital is larger than that with the C  $2p$  orbital, the perturbative strength is correspondingly smaller for the H  $1s$  bonding orbital.<sup>[20]</sup> This means that more energy localized  $d$  states will have a more significant effect on  $*CO_2$  bonding compared with H bonding.

The pH effects on reaction pathway are explored by comparing the energy profile of  $CO_2R$  at  $pH=1$  with  $pH=7$  (Figure 2e). For Au and Ag, at  $pH=1$ , although the steps after  $CO_2$  adsorption are all exothermic, the high barrier of  $CO_2$  adsorption limits the  $CO_2R$  activity. In neutral condition ( $pH=7$ ), Au and Ag surfaces will not be covered with  $*H$ , and it can be found that  $CO_2$  adsorption will be much easier than that in acidic condition. As for CoPc@CNT, the energy of  $*CO_2$  protonation step (from  $*CO_2$  to  $*COOH$ ) is obviously higher than that of  $CO_2$  adsorption step (from  $CO_2$  to  $*CO_2$ ), indicating that the potential-limiting step is the protonation of  $*CO_2$  to  $*COOH$ . Previous work showed that the relative energy of  $CO_2$  adsorption and  $*CO_2$  protonation step is decisive factor in determining the pH-dependent behavior.<sup>[15a]</sup> Specifically, if  $\Delta G(CO_2 \text{ to } *CO_2) > \Delta G(*CO_2 \text{ to } *COOH)$ , a pH-independent behavior is observed because  $CO_2$  adsorption will be the potential-limiting step, whereas a converse scenario is encountered when  $\Delta G(CO_2 \text{ to } *CO_2) < \Delta G(*CO_2 \text{ to } *COOH)$ , since  $*COOH$  formation involves coupled proton and electron transfer. Accordingly, DFT calculations reveal that  $CO_2R$  is pH-dependent for CoPc@CNT, which is consistent with previous reports.<sup>[15a, 16, 21]</sup> As a result, the reaction energy in an acidic condition (0.68 eV,  $pH=1$ ) is lower than that in a neutral condition (1.03 eV,  $pH=7$ ). Based on the above results, DFT calculations suggest that Au and Ag exhibit remarkable  $CO_2R$  activity in alkaline condition but cannot compete with HER in acidic solution due to the difficult  $CO_2$  adsorption on H-covered surfaces. In contrast, on CoPc@CNT, the  $*CO_2$  intermediate can be stabilized in acidic conditions and lowering the pH will favor the potential-limiting step and decrease the overpotential of  $CO_2R$ .

To verify the above theoretical predictions, the single-atom catalyst CoPc@CNT is fabricated by mixing the multi-walled carbon nanotubes (CNT) and CoPc-saturated DMF solution with magnetic stirring, during which CoPc molecules can be anchored on CNT via strong  $\pi$ - $\pi$  interactions. It should be noted that CoPc-saturated solution instead of CoPc-supersaturated solution is used in this work, ensuring that all CoPc molecules are dispersed over CNT at the molecular level rather than forming self-accumulating clusters, which is also aligned with the computational single-atom structure model. This point is confirmed by scanning electron microscopy (SEM) images, in which only CNT is observed when using CoPc-saturated solution (Figure 3a), while obvious CoPc particles are present when CoPc-supersaturated solution is used according to previous work<sup>[22]</sup> (Figure S4). From X-ray diffraction (XRD) patterns (Figure 3b), CoPc@CNT shows only CNT diffraction peaks without the signals of CoPc, inferring that CoPc molecules are dispersed on CNT without forming CoPc crystals. Similar result is also demonstrated from the Raman spectra (Figure 3c), where only D band and G band of CNTs are probed at 1350 and 1590  $cm^{-1}$ . Moreover, the Energy-dispersive X-ray spectroscopy (EDS) elemental maps (Figure 3d) also demonstrate a high-level distribution of Co over the CNTs substrates. To identify the local electronic structure and chemical environment of Co metal in CoPc@CNT, X-ray absorption near-edge structure (XANES) (Figure 3e) and Fourier-transformed extended X-ray adsorption fine structure (EXAFS) (Figure 3f) analyses are performed. As shown in Figure 3e, the adsorption edge position of CoPc@CNT is close to that of CoO, suggesting that the oxidation state of Co atom in CoPc@CNT is similar to that in CoO (+2). In EXAFS curves (Figure 3f), the peak of around 1.41, 2.30 and 3.61 Å in CoPc@CNT are assigned to Co– $N_1$ , Co–C and Co– $N_2$  signals,<sup>[23]</sup> while no Co–Co bonding signal is observed indicating that the as-prepared CoPc@CNT in this work is indeed a single-atom catalyst. To avoid  $CO_2$  mass transfer limitations,<sup>[24]</sup> the CoPc@CNT gas diffusion electrode is fabricated by loading CoPc@CNT onto hydrophobic carbon paper with Nafion binder. In addition, bulk metal catalysts Au and Ag catalysts are fabricated by magnetron sputtering.

To confirm the successful fabrication of gas diffusion electrodes (GDE),  $CO_2R$  experiments are performed in neutral catholyte ( $CO_2$ -saturated 1 M  $KHCO_3$ ,  $pH=7.8$ ), and the faradic efficiency of CO (denoted as CO-FE) as a function of potential is shown in Figure S5–S7. The Au electrode shows a CO-FE of around 70% from  $-1.0$  to  $-1.8$  V vs. SHE, and the Ag electrode exhibits a CO-FE of 90% at  $-1.73$  V vs. SHE, which is consistent with literature.<sup>[25]</sup> Meanwhile, the CoPc@CNT electrode presents a CO-FE of nearly 100% from  $-1.0$  V to  $-1.4$  V vs. SHE. The above results indicate that the obtained electrodes enable highly efficient conversion from  $CO_2$  to CO in neutral media, and also confirm the effectiveness of the gas diffusion electrode in flow cell.

The evaluation of  $CO_2R$  performance in pure acid is carried out respectively on Au, Ag and CoPc@CNT electrodes. In flow cells, both catholyte and anolyte are sulfuric



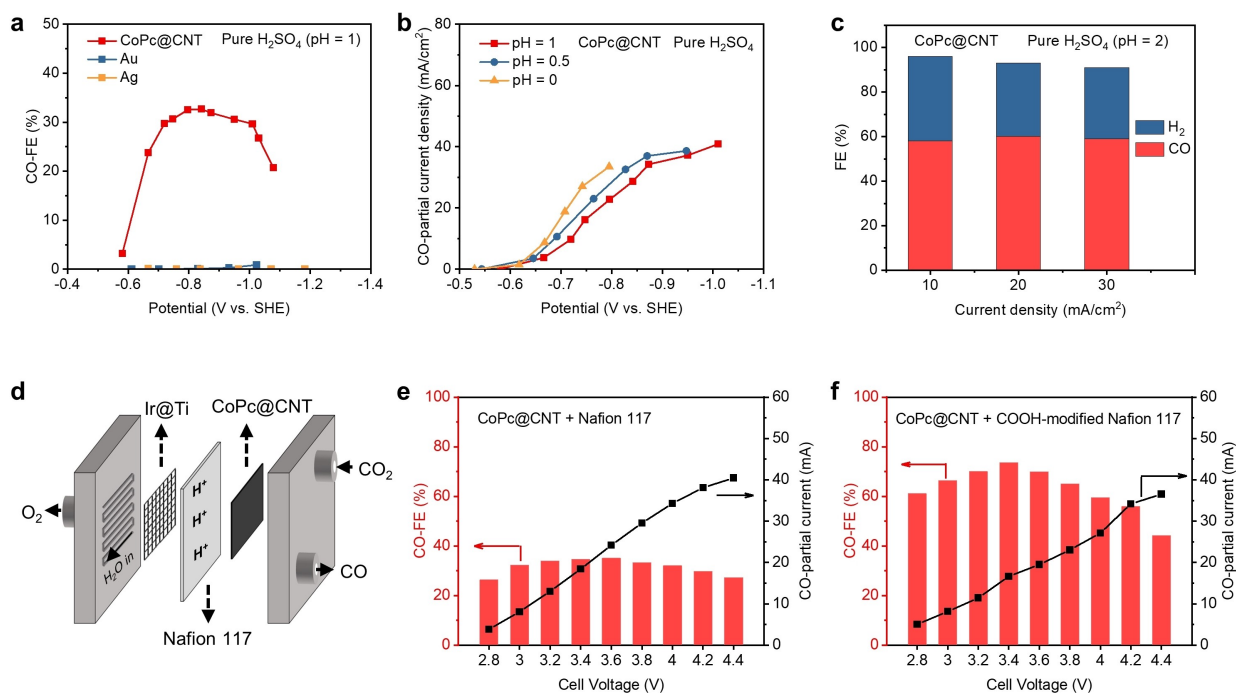
**Figure 3.** Synthesis and Characterizations of single-atom catalyst CoPc@CNT. (a) SEM images of CoPc@CNT. (b) XRD patterns and (c) Raman spectra of CoPc particle, CNT and CoPc@CNT. (d) EDS elemental maps of CoPc@CNT. (e) Co K-edge XANES spectra of CoPc@CNT, Co Foil and CoO. (f) Co K-edge EXAFS spectra of CoPc@CNT and Co Foil.

acid aqueous solution (pH=1), furthermore, the electrolyte differs from previous works in that no alkali cations are added.<sup>[4,7a,8,18b]</sup> Although alkali cations can suppress HER and enhance CO<sub>2</sub>R in flow cell, it is not suitable for CMEA due to the salt precipitation.<sup>[2a]</sup> Nafion 117 is employed to separate catholyte and anolyte, as well as for facilitating the transport of protons. Ir@Ti mesh is used as the counter electrode to realize the water oxidation reaction in acidic media. Since the catalyst can maintain stability during short-term testing, a complete polarization curve can be obtained by a single catalyst (Figure S8). As shown in Figure 4a and S9, a CO-FE of  $\approx 32\%$  is obtained at  $-0.8\text{ V}$  vs. SHE for CoPc@CNT. However, no CO<sub>2</sub> activities are observed on both Au and Ag electrodes (Figure S10–S11). These results are consistent with the above theoretical predictions that CoPc@CNT instead of Au and Ag can drive CO<sub>2</sub> conversion in pure acid due to the strong CO<sub>2</sub> adsorption (Figure 2). In addition, both excessive (FePc@CNT) and insufficient (NiPc@CNT and Ni@N-doped carbon) interactions with the reaction intermediate are unfavorable for pure acidic CO<sub>2</sub>R (Figure S12), and further explorations are needed to understand the selectivity differences among single-atom catalysts. Regardless, the stability of \*CO<sub>2</sub> intermediate is necessary for acidic CO<sub>2</sub>R in the absence of alkali cations.

The CO<sub>2</sub>R activities of CoPc@CNT are further explored in different acidic conditions. Even under more acidic conditions (pH=0.5 and 0), CoPc@CNT stabilizes the reaction intermediates to enable CO<sub>2</sub> conversion (Figure 4b). The reaction potential of CO<sub>2</sub>R shifts positively with a decrease in acidity, aligning with our theoretical calculations, indicating that CO<sub>2</sub>R is pH-dependent for CoPc@CNT. However, HER is enhanced dramatically as the acidity increase, resulting in obvious decreases of CO-FE (Figure S13). On the contrary, HER is effectively sup-

pressed by reducing the proton concentration thus a notable CO-FE of 60% is achieved at pH=2 (Figure 4c and S14). Due to the low ionic conductivity and the large ohm resistance, CO<sub>2</sub>R is difficult to be operated under a higher current density and a weaker acidic condition (Figure S15) in flow cells.

CMEA is an easily scale-up device which can efficiently minimize the ohmic loss of electrolyte in flow cells (Figure S16).<sup>[2a,11]</sup> For CMEA device (Figure 4d), cation exchange membrane is placed between cathode and anode as solid electrolyte, through which the protons generated at the anode transport to the cathode to participate in CO<sub>2</sub>R. In addition, ultra-pure water as the anolyte flows into the anode chamber, which can avoid the interference of impurity cations on cathodic CO<sub>2</sub>R reaction. As shown in Figure 4e, a CO-FE of around 35% is obtained at 3.6 V, which is similar to that of CO<sub>2</sub>R performed in flow cells at pH=1 (Figure 4c). The stability test of MEA is performed at 3.4 V, and it still maintains 80% of the initial CO-FE after 13-hour operation (Figure S17). The decrease in CO<sub>2</sub> selectivity is primarily attributed to catalyst deactivation, and this issue is expected to be improved through substituent group modifications according to the previous report.<sup>[26]</sup> To improve the CO<sub>2</sub> selectivity by weakening the surface acidity of Nafion membrane, polyethylene microporous separator (thickness of 32  $\mu\text{m}$ ) with traces of acetic acid (CH<sub>3</sub>COOH) aqueous solution (pH=3) is introduced between CoPc@CNT and Nafion membrane. The CO-FE has shown a significant improvement, increasing from 35% to 52%, as depicted in Figure S18. To overcome the issue of acetic acid volatilization while retaining its function in reducing the surface acidity of the Nafion membrane, carboxyl groups (–COOH) are modified onto the surface of the Nafion membrane through a combination of electrostatic



**Figure 4.** Catalytic activity of CoPc@CNT at the acidic interface. (a) CO-FE comparison of CoPc@CNT, Au and Ag in pure H<sub>2</sub>SO<sub>4</sub> electrolyte (pH = 1) under different applied potentials using flow cell. (b) CO-partial current densities of CoPc@CNT in pure acid (pure H<sub>2</sub>SO<sub>4</sub> electrolyte, pH = 0, 0.5 and 1) under different applied potential using flow cell. (c) FE of CO and H<sub>2</sub> on CoPc@CNT in pure H<sub>2</sub>SO<sub>4</sub> electrolyte (pH = 2) under different current densities using flow cell. (d) Schematic of CMEA device for CO<sub>2</sub> reduction. (e) CO-FE and CO-partial current of CoPc@CNT under different cell voltages using CMEA. (f) CO-FE and CO-partial current of CoPc@CNT at varied cell voltages in CMEA when Nafion membrane is modified by carboxyl groups (–COOH).

assembly and chemical reactions, as illustrated in Figure S19. The increased electrochemical impedance suggests that the COOH groups have been successfully modified onto the surface of the Nafion membrane (Figure S20), achieving the desired effect of reducing surface acidity. After the modification, the CO-FE exceeds 60% across a wide voltage range (2.8–4.0 V), with the highest value reaching up to 73% at 3.4 V (Figure 4f and S21). Comparing the changes in partial currents and FE of H<sub>2</sub> and CO before and after COOH modification (Figure S22), the increase in CO-FE is due to the COOH group inhibiting HER rather than enhancing CO<sub>2</sub>R. Meanwhile, the protonation process is the potential-limiting step for CO<sub>2</sub>R on CoPc@CNT, leading to a decrease in CO partial current density after COOH modification. From the above, we successfully realize CO<sub>2</sub> conversion in CMEA using CoPc@CNT.

Obviously, there are still some unresolved issues, particularly concerning the issue of CO-partial current density. This issue presents two main challenges. Firstly, at high applied potentials, the higher proton concentration at the catalyst surface enhances the adsorption of H, thereby inhibiting CO<sub>2</sub> adsorption and leading to decreased CO<sub>2</sub>R selectivity. Secondly, the multi-step nature of the CO<sub>2</sub>R process may be slower than the competing HER, even under similar adsorption energy, favoring HER selectivity. We believe that addressing this issue solely from the catalyst's perspective is challenging at present, but there is hope for resolving it by combining control of the surface micro-

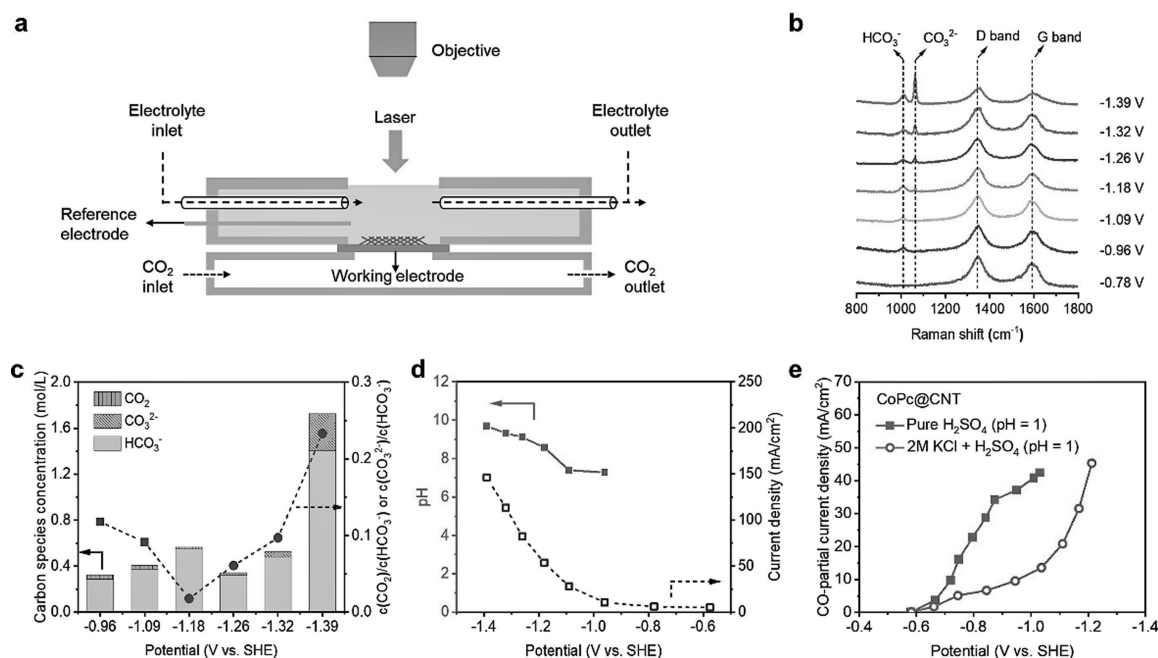
environment. In this work, the introduction of organic cations onto the catalyst's surface is proven to be an effective strategy for enhancing the CO-partial current density (Figure S23). Relative to the non-additive condition, the addition of 0.2 M tetramethylammonium chloride significantly increases the CO-partial current density from 40 mA/cm<sup>2</sup> to 71 mA/cm<sup>2</sup> in flow cell. Under high current conditions, the organic cations of tetramethylammonium will encounter the same issue as alkali metal ions, which is the formation of bicarbonate precipitates, resulting in a decrease in CO selectivity. This issue can be overcome by cross-linking for organic cations, which has already been verified by concurrent research.<sup>[13b]</sup> It should be noted that the immobilized organic cations are drawn by the long chains, making it difficult to ensure that all catalytic sites can be adequately covered by immobilized cations. HER will dominate on uncovered catalysts owing to the unstable CO<sub>2</sub> intermediates. Thus, it is expected that the catalyst itself can stabilize CO<sub>2</sub> intermediates and reduce its dependence on the surface microenvironment, which is exactly what the article aims to emphasize. From the above, to achieve a more significant improvement in selectivity and current density for metal-free acidic CO<sub>2</sub> reduction, it is necessary to integrate catalyst design with microenvironment control.

DFT calculations reveal that the protonation of \*CO<sub>2</sub> to \*COOH step is the potential-limiting step for CoPc@CNT (Figure 2e), meaning that reducing the proton concentration (i.e., increasing the local pH) at the catalyst surface will

increase the reaction barrier. Although pH dependence of the electrochemical reaction can be explored by changing the bulk pH,<sup>[15a]</sup> this method neglects the variation in local pH under the electrolysis, so the conclusions obtained may not be accurate. To avoid this problem, we combine electrochemistry analysis with in situ Raman spectroscopy to investigate the relationship between overpotential and local pH. If the protonation process is the potential-limiting step, a higher overpotential is expected when local pH increases. Herein, the overpotential can be obtained by electrochemistry analysis, and local alkalinity is constructed by adding highly concentrated  $K^+$ ,<sup>[4,27]</sup> which can be quantitatively probed by in situ Raman spectroscopy.

The schematic representation of in situ electrochemical Raman spectroscopy is depicted in Figure 5a and S24. In this study,  $HCO_3^-$  and  $CO_3^{2-}$  are selected as pH probes because they are the products of the  $CO_2-OH^-$  neutralization reaction,<sup>[28]</sup> which avoids any interference from incorporating additional pH-sensitive species. Figure 5b shows the Raman spectra with different cathodic potential (from  $-0.78$  to  $-1.39$  V vs. SHE) in K-containing acidic media (2.0 M KCl in HCl aqueous solution,  $pH=1$ ). The pronounced peaks of around  $1350$  and  $1590$   $cm^{-1}$  are attributed to D band and G band of the carbon nanotube, which indicates successful focusing of the light spot on the catalyst. In the spectra, the Raman signals at the wavenumbers of  $1012$  and  $1064$   $cm^{-1}$  are assigned to  $HCO_3^-$  and  $CO_3^{2-}$  vibrational modes, respectively. Since the reaction is conducted under the condition of  $pH=1$ ,  $HCO_3^-$  and  $CO_3^{2-}$  are not present in the initial reaction solution even when a low potential is

applied ( $-0.78$  V). They are only produced in the regions of catalyst surface and diffusion layer when the potential reaches  $-0.96$  V or higher. Although surface enhancement effects were not employed, combining the testing principle with the actual conditions reveals that the Raman signals of  $HCO_3^-$  and  $CO_3^{2-}$  originate from the region near the catalyst. The distinct Raman features of  $HCO_3^-$  and  $CO_3^{2-}$  ensure that they can be independently quantified using calibration curves (Figure S25), and their corresponding concentration as a function of potential are illustrated in Figure 5c. The  $CO_2$  concentration is originated from the reasonable assumption, according to previous report.<sup>[29]</sup>  $CO_2$  and  $CO_3^{2-}$  cannot coexist in water, and  $pH=8.37$  is a threshold value, as shown in Figure S26. When the  $CO_3^{2-}$  signal is not observed in Raman spectra, and considering the continuous supply of gaseous  $CO_2$  (1 bar), the dissolved  $CO_2$  is assumed to be a constant saturation concentration of  $0.033$  M. Conversely, when the  $CO_3^{2-}$  signal is observed in Raman spectra, the concentration of dissolved  $CO_2$  is regarded as constant at  $0$  M. Based on the acid-base equilibrium reactions, local pH can be further derived (Figure 5d). It is worth noting that the local pH is up to  $\approx 7.3$  when the cathodic potential is  $-0.96$  V vs. SHE with the low current density of  $10.53$   $mA/cm^2$ . This high pH value can be attributed to  $K^+$  cations, which modulate the distribution of electric field in the double layer and thus suppress the migration of hydronium ions.<sup>[4]</sup> Moreover, a higher local pH is also observed at a more negative potential. The calculated pH values based on Raman tests and assumption align perfectly with the theoretical range



**Figure 5.** Experimental verification of the potential-limiting step. (a) Schematic diagram of in situ electrochemical Raman test. (b) Raman spectra of CoPc@CNT in 2.0 M KCl aqueous solution ( $pH=1$ ) under different applied potentials. (c) Carbon species concentrations derived from the spectra in b and the ratio of  $CO_2/HCO_3^-$  (square) and  $CO_3^{2-}/HCO_3^-$  (circle) under different applied potentials. (d) pH profile derived from carbon species ratio in c and current density under different applied potentials. (e) CO-partial current densities of CoPc@CNT in acidic catholyte ( $pH=1$ ) with or without highly concentrated K cations under the different density using flow cell.

(Table S1), further validating the rationality of the assumed  $\text{CO}_2$  values. Meanwhile, no signals of  $\text{HCO}_3^-$  or  $\text{CO}_3^{2-}$  are observed in alkali-cation-free acidic  $\text{CO}_2\text{R}$  process (Figure S27). From above, in situ electrochemical Raman spectroscopy proves that highly concentrated  $\text{K}^+$  can induce an obvious increase of local pH. Similar trends are also observed in COMSOL simulations (Figure S28 and S29).

As shown in Figure 5e (details in Figure S30), overpotential shows a dramatically increase after adding highly concentration of  $\text{K}^+$ . At the same  $\text{CO}$ -partial current density of  $30 \text{ mA/cm}^2$ , the applied potential of the  $\text{K}$ -containing system with high local pH ( $-1.16 \text{ V}$  vs. SHE) is  $320 \text{ mV}$  higher than that of the pure acidic system with low local pH ( $-0.84 \text{ V}$  vs. SHE). Combined with results from in situ Raman spectroscopy, a higher overpotential is observed experimentally as the local pH increases, proving that the protonation of  $^*\text{CO}_2$  to  $^*\text{COOH}$  step is the potential-limiting step for  $\text{CoPc@CNT}$  (Figure 2e). This result is also confirmed by potential analysis with a wide pH range<sup>[15a]</sup> and kinetic isotope experiments<sup>[30]</sup> (Figure S31 and S32).

## Conclusion

In summary, we demonstrate  $\text{CO}_2$  conversion in pure acid without alkali cations using the single-atom catalyst  $\text{CoPc@CNT}$ . In this work,  $\text{CoPc@CNT}$  provides the  $\text{Co}$  single-atom active sites with energetically localized  $d$  states to stabilize the chemisorbed  $^*\text{CO}_2$  intermediates at the acidic interface, exhibits a faradaic efficiency of 60 % in flow cells and realizes  $\text{CO}_2$  conversion in CMEA with a  $\text{CO-FE}$  of 73 %. In addition, the pH-dependent protonation of  $^*\text{CO}_2$  to  $^*\text{COOH}$  is confirmed as the potential-limiting step for  $\text{CoPc@CNT}$ , thus lowering the pH will make the reaction energy of such step more favorable and drive  $\text{CO}_2\text{R}$  with a lower overpotential compared to alkaline conditions. This work not only provides a new understanding for the stabilization of  $^*\text{CO}_2$  intermediates in acidic media, but also facilitates the design of catalysts and devices for  $\text{CO}_2\text{R}$  in CMEA.

## Data Availability

The data that support the findings of this study are available from the corresponding author upon reasonable request.

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## Conflict of Interest

The authors declare no conflict of interest.

## Data Availability Statement

The data that support the findings of this study are available from the corresponding author upon reasonable request.

**Keywords:** Acidic Interface ·  $\text{CO}_2$  Adsorption, Stabilizing  $^*\text{CO}_2$  Intermediates ·  $\text{CO}_2$  Reduction · Molecularly Dispersed Catalyst

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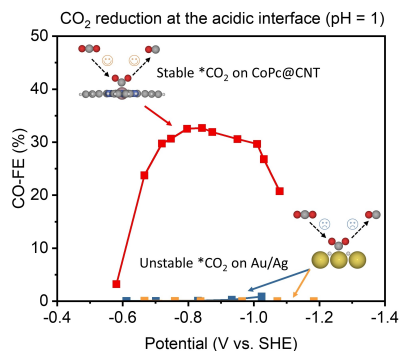
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## Research Articles

CO<sub>2</sub> Reduction

S. Feng, X. Wang, D. Cheng, Y. Luo,  
M. Shen, J. Wang, W. Zhao, S. Fang,  
H. Zheng, L. Ji, X. Zhang, W. Xu, Y. Liang,  
P. Sautet, J. Zhu\* — e202317942

Stabilizing \*CO<sub>2</sub> Intermediates at the Acidic Interface using Molecularly Dispersed Cobalt Phthalocyanine as Catalysts for CO<sub>2</sub> Reduction



Without the assistance of metal cations, CoPc@CNT itself stabilizes \*CO<sub>2</sub> intermediates at the acidic interface, and thus enables CO<sub>2</sub> conversion in pure acid, even at pH = 1. As for Au/Ag, CO<sub>2</sub> adsorption step is completely suppressed by the high coverage of H, causing only hydrogen evolution reaction to occur.