High-performance electrocatalytic reduction of CO₂ to CO by ultrathin PdCu alloy nanosheets

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ABSTRACT

Electrochemical reduction of carbon dioxide (CO₂RR) into fuels and chemicals is a significant step to balance the carbon cycle. For CO₂ reduction to CO, the binding energy of intermediates *COOH and *CO is crucial for the electrocatalytic activity and selectivity. Herein, we prepare a series of ultrathin and tunable PdCu alloy nanosheets by a one-pot wet chemistry process and demonstrate that the bimetallic nanosheets can improve the electronic configurations and break the inherent scaling relationship of intermediate binding energy, which leads to a high performance of CO₂RR. Specifically, Pd₁Cu₁ exhibits the best CO₂RR performance among the series, with a faraday efficiency (FE) of 97% CO at ~0.80 V vs reversible hydrogen electrode (RHE) in 0.1 M KHCO₃ and 96% at ~0.88 V in 1 M KOH with a current density of 521 mA cm⁻². This is ascribed to moderate *COOH binding and weak binding energy of *CO on Pd₁Cu₁, as manifested in both experimental and theoretical studies. Results from this work highlight the unique potentials of bimetallic alloy nanosheets as high-performance catalysts for electrochemical reduction of CO₂.

1. Introduction

The combustion of fossil fuels leads to the increase of carbon dioxide (CO₂) in the atmosphere, breaking the carbon cycle and causing environmental problems, such as greenhouse effect, acid rain, etc. [1,2]. Electrochemical CO₂ reduction reaction (CO₂RR) to value-added chemicals and fuels is an effective strategy to alleviate the greenhouse issues and promote a circular economy [3–7]. Among the products of CO₂RR, CO is considered as the most promising one, since it is a critical feedstock for the production of renewable fuels and chemicals, including acids, esters, and alcohols, which can replace conventional fossil-based fuels and chemicals [8–10]. However, owing to the chemical inertness of CO₂, the wide range of reduction products and competitive hydrogen evolution reaction (HER) [11,12], effective catalysts are needed to reduce the high energy barrier and improve CO selectivity for CO₂RR, which depend on the binding energy and chemisorption configuration of reaction intermediates [13]. An inherent challenge is that the intermediate bindings follow the conventional scaling relationships due to the similar surface – adsorbate bonds [14,15]. For example, the C-binding energy of the intermediates, such as *COOH and *CO, shows the same trend when the center of the d band is shifted [13], resulting in a reduced activity. For pure Pd electrodes, the strong binding energy of Pd-C bond is beneficial to reduce the Gibbs free energy of the *COOH intermediate, while the strong binding energy of *CO is not conducive to the formation of CO, resulting in a low Faraday efficiency of CO (FE₀₀) [16–18]. This scaling relationship can be broken by the rational design and engineering of ideal catalysts that feature a moderate *COOH binding energy and weak *CO binding energy [19–21].

In recent years, bimetallic Pd-based alloy nanostructures have attracted great attention due to their tunable electronic/surface configurations, which can impact the binding energy of key reaction intermediates by manipulation of the atomic ratio in the alloy [22–25]. For instance, Pd–Au alloy catalysts have been found to exhibit a weakened Pd–C bond and balanced bindings of *COOH and *CO, leading to a high efficiency and mass activity toward CO₂RR to CO [26]. Pd₀.₇₅Au₀.₂₅ catalysts have also been observed to display a significant

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decrease in the binding of \(^*\)CO and \(^*\)H, while retain a strong binding of \(^*\)COOH, which facilitates the electron-transfer kinetics toward CO production (\(\text{FE}_{\text{CO}} = 95.3\%\)) \([27]\]. With a low d-orbital vacancy, Cu has also been used to engineer Pd-based alloys and modulate the electronic effect toward well-coordinated \(^*\)COOH and \(^*\)CO. For instance, Cu-Pd alloys with a 1:1 atomic ratio show a high \(\text{FE}_{\text{CO}}\) of 87\% at \(-0.90\) V due to the increase of the energy barrier of the \(^*\)CO protonation step, as compared to the monometallic Pd \([28]\).

In addition, the catalytic performance can also be enhanced by structural engineering of the metallic catalysts at the nanoscale. Towards this end, two-dimensional (2D) nanostructures have been drawing extensive attention due to the relatively high surface area-to-volume ratio and high density of exposed atoms on their surface \([29]\). Herein, PdCu alloy nanosheets are prepared by a one-pot wet chemistry process at various Pd:Cu atomic ratios. Alloying Cu to Pd is beneficial for breaking the scaling relation between the binding energy of specific intermediates, and thus facilitates the formation of CO. Electrochemically, the PdCu alloy nanosheets exhibits an \(\text{FE}_{\text{CO}}\) over 95\% with a current density of 521 mA cm\(^{-2}\) in 1 M KOH at \(0.88\) V. Density functional theory (DFT) calculations show that Pd\(_1\)Cu\(_1\) exhibits the best selective production of CO among the series, due to a reduced energy barrier of CO formation.

### 2. Experimental

#### 2.1. Materials

Palladium(II) acetylacetonate (Pd(acac)\(_2\)), copper(II) acetylacetonate (Cu(acac)\(_2\)), molybdenum carbonyl (Mo(CO)\(_6\)), trietylphosphine oxide (TOPO), oleic acid (OA), isopropyl alcohol, N, N-dimethylformamide (DMF) and Nafion (5\%) were purchased from Sigma-Aldrich. Ethanol (99.9\%) was purchased from Damao. CO\(_2\) (purity 99.995\%), Ar (purity 99.99\%), N\(_2\) (purity 99.99\%) and H\(_2\) (purity 99.999\%) were obtained from Zibo Baiyan Gases. DI water was obtained from a Barnstead Nanopure water purification system (18.3 M\(\Omega\) cm). All materials were used without further purification.

#### 2.2. Synthesis of PdCu alloy nanosheets

PdCu alloy nanosheets was synthesized via a modified one-pot wet chemistry process. Firstly, 5 mg (16.4 mmol) of Pd(acac)\(_2\), 4.3 mg (16.4 mmol) of Cu(acac)\(_2\), 6 mg (22.7 mmol) of Mo(CO)\(_6\) and 63 mg (162.9 mmol) of TOPO were added into 4 mL of OA in a 50 mL round-bottom flask under sonication for 10 min, before the addition of 0.75 mL of DMF under vigorous stirring in an Ar atmosphere for 30 min. The flask

![Fig. 1](image_url)
was then heated into an oil bath at 58 °C for 18 h. The resulting solution was centrifuged at 8,000 rpm for 5 min, and the precipitate was collected and rinsed 3 times with DI water and ethanol. Finally, the product was desiccated in a vacuum oven (40 °C), which denoted as Pd_1Cu_y. PdCu alloy nanosheets at other Pd/Cu atomic ratios were prepared by using the same synthetic method but with a different amount of Cu(acac)_2 and denoted as Pd_xCu_y. Monometallic Pd and Cu were also prepared by following the same procedure by using only Pd(acac)_2 or Cu(acac)_2 precursors.

3. Results and discussion

3.1. Characterization of the samples

The PdCu alloy nanosheets at different Pd:Cu atomic ratios were prepared by a one-pot wet chemistry process, where palladium(II) acetylacetonate (Pd(acac)_2) and copper(II) acetylacetone (Cu(acac)_2) were reduced thermally in oleic acid and N,N-dimethylformamide, as illustrated in Fig. 1a (details in the Supporting Information). Five samples were prepared, Pd, Pd_1Cu_y, Pd_2Cu_y, Pd_3Cu_y and Cu, and the crystalline structure was first examined by X-ray diffraction (XRD) measurements. As shown in Fig. 1b-c, one can see that the monometallic Pd sample exhibited four major diffraction peaks at 2θ = 40.1°, 46.6°, 68.1°, and 82.1° that can be assigned to the (1 1 1), (2 0 0), (2 2 0) and (3 1 1) crystalline planes of fcc Pd (PDF No. 46-1043). For the PdCu bimetallic samples, these diffraction peaks can be seen to shift to a higher angle with an increasing Cu content, suggesting the formation of Pd-Cu alloys, due to a reduced lattice spacing with Pd being replaced with smaller Cu atoms [30,31]. For instance, the four diffraction peaks of Pd_1Cu_2 appeared at 40.8°, 47.4°, 69.6°, and 83.9°, and the diffraction profile was markedly different from that of the monometallic Cu sample, which featured three diffraction peaks at 2θ = 43.3°, 50.4°, and 74.1°, consistent with fcc Cu (PDF No. 04-0836).

The surface morphology of the samples was characterized by transmission electron microscopy (TEM) measurements. As shown in Fig. 1d and Fig. S1, the samples all exhibited an ultrathin nanosheet structure. In high-resolution TEM (HRTEM) measurements (Fig. 1e), Pd_1Cu_y can be seen to display well-defined lattice fringes with an interplanar spacing of 0.224 nm (Fig. 1j), which is in line with the (1 1 1) facets of PdCu alloy. The selected area electron diffraction (SAED) patterns of Pd_1Cu_y were shown in the inset to Fig. 1e, which can be ascribed to the (1 1 1) crystal planes of PdCu alloys, in good agreement with results from the XRD measurements (Fig. 1b-c). This is further confirmed in elemental mapping analysis based on energy dispersive spectroscopy (EDS) where the Pd and Cu elements were evenly distributed over the entire nanosheets (Fig. 1f-i).

The PdCu alloy samples, the binding energy of these two doublets can be seen to exhibit a marked red-shift with increasing Cu loading, from 335.3/340.5 eV and 336.4/341.5 eV for Pd_1Cu_2 to 335.0/340.2 eV and 336.1/341.3 eV for Pd_2Cu_y and 334.8/340.0 eV and 335.9/341.1 eV for Pd_3Cu_y. The high-resolution Cu 2p spectra are depicted in Fig. 2b, which can also be deconvoluted into two doublets. For the monometallic Cu sample, the two doublets appeared at 934.2/954.0 eV and 931.9/951.7 eV, due to 2p_1/2 and 2p_3/2 electrons of metallic Cu and Cu^{2+} species, respectively [32]. Yet, for the PdCu alloy samples, the binding energy of these two doublets can be seen to exhibit a marked red-shift with increasing Cu loading, from 335.3/340.5 eV and 336.4/341.5 eV for Pd_1Cu_2 to 335.0/340.2 eV and 336.1/341.3 eV for Pd_2Cu_y and 334.8/340.0 eV and 335.9/341.1 eV for Pd_3Cu_y. The corresponding high-resolution Pd 3d scans are shown in Fig. 2a, which can be deconvoluted into two doublets. For the monometallic Pd sample, the two doublets can be identified at 335.7/341.0 eV and 336.9/342.2 eV, due to the 3d_3/2 and 3d_5/2 electrons of metallic Pd^{0} and Pd^{2+} species, respectively [32]. Yet, for the PdCu alloy samples, the binding energy of these two doublets can be seen to exhibit a marked red-shift with increasing Cu loading, from 335.3/340.5 eV and 336.4/341.5 eV for Pd_1Cu_2 to 335.0/340.2 eV and 336.1/341.3 eV for Pd_2Cu_y and 334.8/340.0 eV and 335.9/341.1 eV for Pd_3Cu_y. The corresponding high-resolution Cu 2p spectra are depicted in Fig. 2b, which can also be deconvoluted into two doublets. For the monometallic Cu sample, the two doublets appeared at 934.2/954.0 eV and 931.9/951.7 eV, due to 2p_1/2 and 2p_3/2 electrons of metallic Cu and Cu^{2+} species, respectively [33,34]. The oxidation state of metals was probably caused by the surface partial oxidation upon exposure to air. Notably, the binding energies of these doublets were markedly higher in the PdCu samples, and increased with increasing Pd contents, 931.9/951.7 eV and 934.2/954.0 eV for Pd_1Cu_2 to 932.4/952.2 eV and 934.6/954.4 eV for Pd_2Cu_y and 932.9/952.7 eV and 935.1/955.0 eV for Pd_3Cu_y. Taken together, these results suggest effective electron transfer from Cu to Pd, due to the formation of PdCu alloys and a lower electron negativity of Cu than that of Pd [29].

3.2. Electrocatalytic performance for CO_2RR

The electrocatalytic activity of the PdCu alloy nanosheets towards CO_2RR was then investigated in an H-type cell in a typical three-electrode setup by linear sweep voltammetry (LSV) in Ar- or CO_2-saturated 0.1 M KHCO_3. From Fig. S3a it can be seen that Pd_1Cu_y displayed a markedly higher current density in CO_2-saturated 0.1 M KHCO_3 than in Ar-saturated 0.1 M KHCO_3, indicating an apparent CO_2RR activity. The corresponding product selectivity is depicted in Fig. 3b, where CO can be identified as the major product whereas H_2 and CH_4 are the minor ones. Specifically, the FE_{CO} on Pd_1Cu_y was over 80% within a wide potential range of −0.6 to −1 V, reaching a maximum of 97% at −0.9 V. The performance was markedly lower with other samples in the series, where the maximum FE_{CO} was only about 80%, 78% and 62% for Pd_2Cu_y, Pd_3Cu_y and Pd, respectively (Fig. S2a,b and Fig. S3a), whereas...
for the monometallic Cu sample, the maximum FE\textsubscript{CO} was only 27% at 0.8 V and H\textsubscript{2} became the major product, along with others like CH\textsubscript{4}, C\textsubscript{2}H\textsubscript{4} and HCOOH (Fig. S3d). The total current density and partial geometric current density of the products at varied applied potentials are shown in Fig. S2c-d, Fig. S3b-c and S3e-f. It can be clearly seen that Pd\textsubscript{1}Cu\textsubscript{1} possessed the highest geometric current density of CO (j\textsubscript{co}), reaching a maximum of 33 mA cm\textsuperscript{-2} at 1.1 V (Fig. 3c). This suggests that Pd\textsubscript{1}Cu\textsubscript{1} stood out as the best catalyst for CO\textsubscript{2}RR to CO among the sample series within the context of FE and current density.

The intrinsic activity of the PdCu alloy nanosheets was evaluated by normalizing the current density to the electrochemical surface area (ECSA), which was determined by double layer capacitance (Fig. S4). Pd\textsubscript{1}Cu\textsubscript{1} can be seen to possess a better activity than other samples in the series, and the trend of the current density normalized to ECSA is the same as that of the geometric current density (Fig. S5). Furthermore, the Pd\textsubscript{1}Cu\textsubscript{1} also shows the highest turnover frequency (TOF) value (Fig. 3d, 2064.7 h\textsuperscript{-1}) and the highest formation rate of CO (Fig. S6), among the catalysts. Notably, the cathodic energy efficiency (EE) of the Pd\textsubscript{1}Cu\textsubscript{1} sample for CO formation was also better than results of relevant catalysts reported previously in the literature (Fig. 3e and Table S2).

The corresponding Tafel plots are shown in Fig. S7a, from which Pd\textsubscript{1}Cu\textsubscript{1} can be seen to exhibit a Tafel slope (183 mV dec\textsuperscript{-1}) lower than those of Pd\textsubscript{2}Cu\textsubscript{3} (243 mV dec\textsuperscript{-1}) and Pd\textsubscript{3}Cu\textsubscript{1} (196 mV dec\textsuperscript{-1}), suggesting that the rate determining step (RDS) of CO\textsubscript{2}RR was the generation of CO\textsubscript{2}* and the electron-transfer kinetics was much faster on Pd\textsubscript{1}Cu\textsubscript{1} than on other samples [2,35]. Indeed, in electrochemical impedance spectroscopy (EIS, Fig. S7b) measurements, Pd\textsubscript{1}Cu\textsubscript{1} possesses a lower charge-transfer resistance (R\textsubscript{CT}) than others. Furthermore, the current density

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Fig. 3. (a) LSV curves of Pd\textsubscript{1}Cu\textsubscript{1} in N\textsubscript{2}- and CO\textsubscript{2}-saturated 0.1 M NaHCO\textsubscript{3} at a scan rate of 10 mV s\textsuperscript{-1}, (b) FE of CH\textsubscript{4}, CO and H\textsubscript{2} on Pd\textsubscript{1}Cu\textsubscript{1}, (c) Partial current densities and (d) TOF of CO on the PdCu alloy nanosheets, (e) Energy efficiency of CO\textsubscript{2} to CO of the catalysts reported in the literature, (f) Stability test of Pd\textsubscript{1}Cu\textsubscript{1} at -0.8 V for 42 h. Inset shows the corresponding LSV profiles before and after the stability test.
and Faraday efficiency of the Pd$_1$Cu$_1$ catalyst remained virtually invariant at $-0.8$ V for 42 h in 0.1 M KHCO$_3$ (Fig. 3f), and there was no significant change of the LSV profiles (insert Fig. 3f) and TEM image (Fig. S8), indicating excellent stability of the Pd$_1$Cu$_1$ nanosheets.

Due to the low solubility of CO$_2$ in aqueous solution, only a relatively low current density can be produced in an H-type cell, which limits their practical application that requires a CO partial current density of at least 100 mA cm$^{-2}$ [36]. Thus, the catalytic activity of the Pd$_{1}$Cu$_{1}$ catalyst was also examined in a flow cell reactor in 1 M KOH electrolyte (Fig. 4a), where a small onset potential and the high current density were obtained (Fig. 4b), due to hydroxide ions can suppress the competing HER and lower the CO$_2$ activation energy barrier [11]. Fig. 4c shows the partial current densities and FE$_{CO}$ at various electrode potentials, where $J_{CO}$ can be seen to increase to 521 mA cm$^{-2}$ with an FE$_{CO}$ of 96% at $-0.88$ V. The Pd$_{1}$Cu$_{1}$ catalysts also exhibited excellent stability at high current density, which remained almost unchanged at 230 mA cm$^{-2}$, with FE$_{CO}$ of 97% at $-0.68$ V, in 1 M KOH for 17 h (Fig. 4d).

### 3.3. DFT theoretical calculation

Density functional theory (DFT) calculations were then performed to unravel the correlation between the catalyst structure and adsorption energetics of key reaction intermediates. The structural models are shown in Fig. S9-S10. The different electronic structures of the alloys at different compositions are related to the change of the d-band center, shown in Fig. S9-S10. The different electronic structures of the alloys at different compositions are related to the change of the d-band center.

The binding energy of *COOH and *CO first decreased with the increasing content of Cu, reaching the valley at Pd$_{1}$Cu$_{1}$; yet with a further increase of the Cu content (i.e., Pd$_{1}$Cu$_{3}$ and Cu), the binding of *COOH was slightly enhanced, while the binding of *CO stayed virtually unchanged. In a conventional scaling relationship, the variation of the adsorption energy of *CO and *COOH is correlated to each other, and *COOH binding cannot be optimized without affecting the *CO adsorption [21]. Yet, the results presented above suggest that manipulation of the elemental composition of the PdCu alloys can retain the moderate binding of *COOH and concurrently weakened binding of *CO at appropriate Cu contents, thus breaking the scaling relationship. The weakened adsorption of *CO is conducive to the desorption of CO, and moderate adsorption of *COOH is beneficial to the subsequent reaction. This coordination between *CO and *COOH is of great importance to boost the kinetics toward CO production. In addition, the binding of *H is gradually weakened with the increase of the Cu content, thus suppressing the competition of HER with the appropriate Cu contents, another feature that is conducive to CO$_2$RR electrocatalysis (Fig. S11a).

Analysis of the molecular orbital overlap can be used to further explain the interaction between reaction intermediates and catalytic active sites. Fig. 5d-e depicts the projected density of states (PDOS) of the C in chemisorbed *COOH and *CO and metal atoms on catalysts, and Fig. S11b shows that of *H. As shown in Fig. 5d, there is a large overlap between the C 2p and Pd d orbitals, indicating a strong interaction between C and Pd [39,40]. With the introduction of Cu, the overlap decreased significantly, indicating weakened binding of *COOH; yet with an excessive Cu content, the overlap actually became slightly enhanced, indicating that the binding of *COOH was somewhat strengthened. In the case of *CO, the overlap area decreased first and then remained basically unchanged with the increase of Cu content, indicating that the binding of *CO decayed first and then stayed virtually unchanged. The different variations further illustrate the breaking of the conventional scaling relationship in PdCu alloys. In addition, as the Cu content increased, a decreasing overlap can be observed between

![Fig. 4. (a) Schematic illustration and (b) polarization curves of Pd$_1$Cu$_1$ in 1 M KOH, (c) FE and partial current density of CO$_2$RR on Pd$_1$Cu$_1$ in 1 M KOH solution, (d) Long-term i-t profiles in 1 M KOH electrolyte.](image-url)
the H s and Pd d orbitals over the PdCu surface (Fig. S11b).

To unravel the energetics and reaction pathways of CO$_2$RR to CO and formate, the free energy diagrams on Pd$_1$Cu$_1$ are constructed and shown in Fig. 5c. It can be seen that the entire reaction pathway of CO generation is spontaneous and exothermic on Pd$_1$Cu$_1$, indicating that CO formation is energetically preferred. For other catalysts, certain energy barriers need to be overcome in the process of CO generation (Fig. S12a).

The free energy of HER on PdCu is shown in Fig. S12b. Too strong adsorption of hydrogen may lead to occupation of the active site, whereas too weak adsorption will be conducive to HER and compromise the CO$_2$RR activity [35]. However, the Pd$_1$Cu$_1$ shows only a moderate adsorption of hydrogen (0.18 eV), which not only suppresses the HER process, but also provides sufficient adsorbed hydrogen needed for the reduction of CO$_2$. In summary, results from the DFT calculations suggest that the Pd$_1$Cu$_1$ exhibits the optimal electronic structure for the selective reduction of CO$_2$ to CO, among the present sample series.

4. Conclusion

In summary, a series of ultrathin and tunable PdCu alloy nanosheets were prepare by a one-pot wet-chemistry process. Results from spectroscopic and theoretical studies demonstrated that the resulting nanosheets improved the electronic configurations and broke the inherent scaling relationship of the binding energy of key reaction intermediates. Specifically, the Pd$_1$Cu$_1$ sample exhibited a high FE$_{CO}$ of 97% at −0.8 V and a current density of 33 mA cm$^{-2}$ at −1.1 V. In a flow cell and 1 M KOH electrolyte, a current density of 521 mA cm$^{-2}$ and an FE$_{CO}$ of 96% were obtained at −0.88 V. Furthermore, results from DFT calculations showed that the conventional scaling relationship of intermediates was broken at high Cu contents in the PdCu alloys, where a moderate *COOH binding and negligible *CO binding led to a high performance in the selective production of CO. Such fundamental insights may be exploited for the rational design and engineering of high-performance catalysts for electrochemical energy conversion.

Data availability

Data will be made available on request.

CRediT authorship contribution statement


Declaration of Competing Interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

Data availability

Data will be made available on request.

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Appendix A. Supplementary material

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