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Bridge Sites of Au Surfaces Are Active for Electrocatalytic CO₂ Reduction

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Abstract

Prior in situ attenuated total reflectance Fourier transform infrared (ATR–FTIR) studies of electrochemical CO₂ reduction catalyzed by Au, one of the most selective and active electrocatalysts to produce CO from CO₂, suggest that the reaction proceeds solely on the top sites of the Au surface. This finding is worth updating with an improved spectroelectrochemical system where in situ IR measurements can be performed under real reaction conditions that yield high CO selectivity. Herein, we report the preparation of an Au-coated Si ATR crystal electrode with both high catalytic activity for CO₂ reduction and strong surface enhancement of IR signals validated in the same spectroelectrochemical cell, which allows us to probe the adsorption and desorption behavior of bridge-bonded *CO species (*CO_B). We find that the Au surface restructures irreversibly to give an increased number of bridge sites for CO adsorption within the initial tens of seconds of CO₂ reduction. By studying the potential-dependent desorption kinetics of *CO_B and quantifying the steady-state surface concentration of *CO_B under reaction conditions, we further show that *CO_B are active reaction intermediates for CO₂ reduction to CO on this Au electrode. At medium overpotential, as high as 38% of the reaction occurs on the bridge sites.

Graphical Abstract

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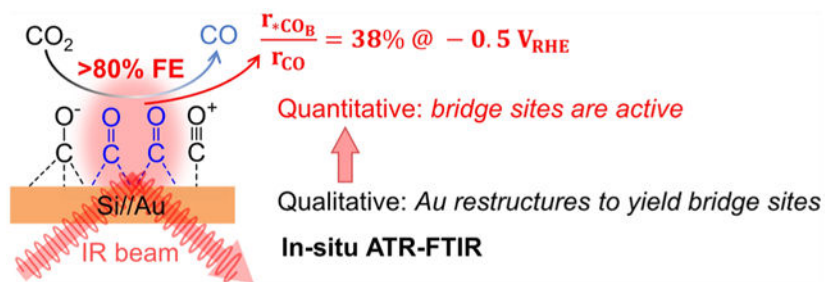
Supporting Information

The Supporting Information is available free of charge at <https://pubs.acs.org/doi/10.1021/jacs.2c01098>.

Experimental details, electrocatalytic performance, SEM image of Si//Au surface, structures and photographs of the spectroelectrochemical and H-type cells, XPS spectra, and additional ATR–FTIR results (PDF)

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1. INTRODUCTION

Electrocatalytic CO_2 reduction reactions are explored as a viable means to produce carbon-neutral chemicals.^{1–7} The rich chemistry of carbon enables CO_2 electroreduction to generate a large variety of products including single-carbon (C_1) and multi-carbon (C_{2+}) species.^{8–18} As the simplest two-electron reduction product, CO is perhaps the most accessible species from CO_2 reduction and finds wide application in industrial processes such as the Fischer–Tropsch reaction¹⁹ and hydroformylation.²⁰ In addition, CO can be electrochemically upgraded to C_{2+} products.^{21,22} Different kinds of materials including metals,^{23–27} metal coordination compounds/materials,^{28–33} and metal-free materials^{34–36} have been found to be active for CO_2 electroreduction to CO. Among them, Au metal is arguably the most well-known for its outstanding selectivity and activity. Polycrystalline Au without any modification or special treatment can achieve high Faradaic efficiency (FE) for CO_2 -to-CO conversion.^{37,38} Catalyst structure optimization and reaction condition engineering can further improve the selectivity and significantly lower the overpotential.^{27,39–42}

Interestingly, rough Au surfaces can enhance vibrational absorption via surface plasmon resonance (known as surface enhanced IR absorption spectroscopy, SEIRAS). This enables the application of in situ attenuated total reflectance (ATR) Fourier transform infrared (FTIR) spectroscopy to detect sub-monolayer coverages of adsorbates for mechanistic understanding under reaction-relevant conditions.^{43–53} Surendranath et al. were among the first to study Au-catalyzed CO_2 electroreduction with in situ ATR–FTIR, and linearly bonded or atop $^*\text{CO}$ ($^*\text{CO}_\text{L}$) and bridge-bonded $^*\text{CO}$ ($^*\text{CO}_\text{B}$) were observed at CO_2 reduction potentials.⁴⁵ However, the use of a Pt counter electrode led to some controversy.^{46,54} Xu and coworkers performed similar measurements in a Pt-free setting and observed only $^*\text{CO}_\text{L}$ species at positive electrode potentials.⁴⁶ More recently, Surendranath et al. reported observation of irreversibly bonded $^*\text{CO}_\text{B}$ species in experiments that followed recommended practices to mitigate contamination,⁵⁵ which reflects the dependence of Au surface structures/properties on electrode preparation. Other studies of this kind probed reactants, reaction intermediates, and/or electrolyte species on Au surfaces.^{41,50–53,55–57} Almost all these studies agree that $^*\text{CO}_\text{L}$ is the only active form of $^*\text{CO}$ on Au.^{45,46,51,52,55,57} However, they share one notable limitation that the catalytic performance, that is, selectivity, activity, and stability, of the Au material deposited on the ATR crystal has never been validated in the spectroelectrochemical cell,^{45,46,51,52,55} which makes it difficult to tie spectroscopic observations directly to catalytic reactivity and to exclude interference from side reaction or contamination. While some latest work started to

validate catalytic performance of ATR electrodes,⁵⁷ the sensitivity of electrode properties to preparation conditions^{47,48,55} entails such validation to be a standard practice. Meanwhile, new electrode structures should be investigated to enrich the prevalent understanding of the reaction mechanism(s).

In this work, we successfully develop a new procedure to deposit a both catalytically and spectroscopically active Au layer on the surface of a Si ATR crystal. The resulting Si//Au electrode, as measured in the spectroelectrochemical cell, exhibits high FEs of 80–90% for CO₂ reduction to CO in the potential range from –0.5 to –0.8 V versus RHE (RHE = reversible hydrogen electrode; all potentials are with respect to RHE unless otherwise stated) together with reasonable current densities and stability. This catalytic performance is on par with that of a typical polycrystalline Au catalytic electrode measured in a standard H-type electrochemical cell, allowing us to carry out an in situ ATR–FTIR study of an Au electrode with benchmarked CO₂ reduction reactivity. We find that the Au surface restructures within the initial tens of seconds of CO₂ reduction to give an increased number of bridge sites for CO adsorption, which has not been reported previously. Using time-resolved scan (TRS), the kinetics of *CO_B desorption from the Au surface is for the first time measured as a function of the applied electrode potential. *CO_B surface concentration is quantified by electrochemical titration and correlated to the IR peak area. On the basis of these results, we derive the contribution of the *CO_B pathway to the total current of CO₂ reduction, which turns out to be a notable portion in the medium overpotential range. This amends the conventional knowledge that *CO_B is a spectator in Au-catalyzed CO₂ electroreduction.

2. RESULTS AND DISCUSSION

Our Si//Au electrode is prepared following the procedure illustrated in Figure 1a (experimental details are provided in the Supporting Information). First, a Cr/Au layer is deposited on the total reflection surface of the Si ATR crystal, followed by a spin-coated thin poly(tetrafluoroethylene) (PTFE) layer. The crystal is then annealed at 350 °C in an Ar atmosphere. The annealed Cr/Au layer renders desirable electrical conductivity, mechanical strength, and adhesion with the Si substrate.⁵⁸ PTFE serves as a binder for the Au catalyst layer to be deposited and facilitates delivery of the gas reactant CO₂.^{59–61} Finally, another Au layer is deposited to render the electrocatalytic activity for CO₂ reduction (Figure S1a,b). X-ray photoelectron spectroscopy (XPS) depth profiling reveals the Si/Cr/Au/PTFE/Au multi-layer structure as expected (Figure 1b). We note that some Cr has diffused through the bottom Au layer after annealing.^{58,62} The topmost Au layer manifests a rough surface consisted of 20 nm particles (Figure S2). The Si//Au electrode is assembled into a home-modified ATR–IR spectroelectrochemical cell (Figure S3a,b) and evaluated for its catalytic performance for CO₂ reduction in purified 0.1 M KHCO₃ aqueous electrolyte. CO is detected by gas chromatography at –0.5 V, and FE(CO) stays over 80% at more negative potentials (Figure 1c). Both FE(CO) and current density are stable for at least 30 min of continuous operation at –0.6 V (Figure 1d), making it sufficient for ATR–FTIR measurements. This performance is on par with 20 nm Au sputter-coated on a carbon fiber paper electrode (Figure S1c) which is tested in a standard H-cell (Figure S3c,d), representing the typical reactivity of polycrystalline Au electrodes.^{38,63}

Successful validation of the catalytic performance of our Si//Au electrode in the spectroelectrochemical cell enables us to perform ATR–FTIR studies of Au under real CO₂ reduction conditions. A potential step program featuring three electrode potentials is adopted (Figure 2a). Constant potential electrolysis (CPE) is first performed at –0.6 V for CO₂ reduction to take place. Then, the potential is changed to 0.4 V for the Au surface to adsorb some of the CO generated earlier and to be probed by IR.⁴⁶ After that the potential is adjusted to –0.2 V to allow for CO desorption and regeneration of the Au sites (Figure S4).⁴⁶ This sequence is repeated with a varied duration of the –0.6 V step. After the first 5 s of CO₂ reduction at –0.6 V, a band at 2108 cm^{–1} is observed at 0.4 V, which is assigned to *CO_L adsorbed on singlefold top sites (Figure 2b), consistent with previous reports.^{45,46,52,55,57} A weaker band centered at around 2000 cm^{–1} is also observed and assigned to be *CO_B adsorbed on twofold bridge sites.^{47,64} Note that this vibrational frequency is considerably higher than that of *CO previously reported to be induced by impurities/contamination.^{46,54} As the potential step sequence continues with increasingly longer CO₂ reduction, the area of the *CO_L peak remains largely unchanged, whereas the *CO_B peak grows evidently with a slight blue shift of 9 cm^{–1} and becomes the dominant *CO species (Figure 2c). Concomitantly, an additional band with much weaker intensity emerges at around 1940 cm^{–1}, which we assign as CO adsorbed on multifold Au sites (*CO_M).⁴⁷ Note that both *CO_B and *CO_M have previously been observed on Au surfaces.⁴⁷ As the potential step sequence further proceeds with gradually reduced CO₂ reduction durations, both *CO_B and *CO_L bands remain largely unchanged, which suggests that the increased amount of *CO_B is not dependent on the CO₂ reduction duration but is a direct result of some surface restructuring induced by the reaction conditions that forms more bridge sites. When the CO₂ reduction potential in this sequence is changed to –0.7 V, similar enrichment of *CO_B can be observed in a smaller amount of CO₂ reduction time (Figure S5), indicating a larger driving force for restructuring under a more reducing condition. Consistently, no *CO_B can be detected at 0.4 V even after 12 cycles of 5 s CO₂ reduction at –0.5 V (Figure S6). When the sequence starts with a fresh electrode under a N₂ atmosphere and is switched to CO₂ from the second cycle (Figure S7a,d–h), the *CO_B band is stronger than *CO_L almost immediately after the switch and appears to further grow with longer CO₂ reduction (Figure S7b,c), suggesting that the Au surface restructuring is mainly induced by the electrochemical condition but may be augmented by the presence of CO₂.

The restructuring to expose/activate more bridge sites within the first tens of seconds of electrochemical CO₂ reduction has not been reported before for Au. Because this phenomenon was not observed in previous SEIRAS studies with Au ATR electrodes prepared by the conventional wet method,^{45,46,50,55,57} we suspect it may be related to differences in electrode preparation which can influence surface roughness, crystallinity, and nanostructures as well as catalytic performance. While the restructuring of our Au electrodes may not be universal, it provides us a model system to answer the fundamental question whether *CO_B could be an active intermediate of the CO₂ reduction reaction. Considering the relatively weak adsorption strength of *CO_B on our Au electrode indicated by its higher vibrational frequency compared to *CO_B on other noble metals,^{27,52,65–70} we hypothesize that *CO_B from CO₂ reduction could desorb to form CO. To test this hypothesis, the desorption kinetics of *CO_B is first measured (Scheme S1). CO is generated from CO₂

reduction at -0.7 V and then captured at 0.4 V, after which the electrode potential is held at $V_{\text{desorption}}$ to measure $^*\text{CO}_B$ desorption using TRS ATR-FTIR with a temporal resolution of 1.648 s (Figure 3a). From the recorded spectra plotted in Figure 3b–e, the desorption of $^*\text{CO}_B$ is clearly visible at ~ 1985 cm^{-1} . Two broad bands centered at 1870 and 1928 cm^{-1} become prominent in the later minutes at $V_{\text{desorption}}$, which are possibly $^*\text{COOH}$ ⁷¹ and $^*\text{CO}_M$, respectively. Therefore, the desorption profile of $^*\text{CO}_B$ at each $V_{\text{desorption}}$, that is, IR peak area versus time, is acquired from the initial 60 s at $V_{\text{desorption}}$ to ensure correct quantification of $^*\text{CO}_B$ desorption (Figure 3f). Here, we assume the surface concentration ($\Gamma_{^*\text{CO}_B}$, the number of CO molecules adsorbed on bridge sites normalized to the geometric area of the Si//Au electrode) is proportional to the IR peak area ($A_{^*\text{CO}_B}$).⁷² The desorption profiles can be simply fitted by the first-order kinetics equation⁷³

$$-\ln \frac{\Gamma_{^*\text{CO}_B}}{\Gamma_0} = -\ln \frac{A_{^*\text{CO}_B}}{A_0} = k_d(t - t_0) \quad (1)$$

where k_d is the rate constant, and Γ_0 (A_0) refers to $^*\text{CO}_B$ surface concentration (IR peak area) at t_0 moment. After acquiring the k_d values for four different desorption potentials with reasonable linearity (Figure 3g), we analyze the potential dependence of the desorption rate constant. We find the relationship between k_d and $V_{\text{desorption}}$ follows a simple exponential equation (Figure 3h)^{74,75}

$$\ln k_d = a \cdot V_{\text{desorption}} + b \quad (2)$$

where a and b are both constants. Extrapolating this equation allows us to estimate k_d values at CO_2 reduction potentials where the generation of CO from CO_2 reduction prevents a direct measurement of the desorption kinetics.

To determine the value of $\Gamma_{^*\text{CO}_B}$, which is needed to calculate the CO production rate via the $^*\text{CO}_B$ pathway, we quantify $\Gamma_{^*\text{CO}}$ at 0.4 V with electrochemical titration and correlate it with the IR peak area (Scheme S1).^{43,44} A potential step program is adopted (Figure 4a) and monitored with TRS ATR-FTIR. CO is generated at -0.7 V, captured at 0.4 V, and then electrochemically stripped at 1.0 V (oxidized to CO_2). By holding the potential at 0.4 V for 30 s, unstable reductive intermediates and free CO in the local electrolyte are removed under continuous CO_2 bubbling. The corresponding IR spectra well reflect the capture and oxidative removal of $^*\text{CO}_L$, $^*\text{CO}_B$, and $^*\text{CO}_M$ (Figure 4b). The 0.4 and 1.0 V steps are repeated once to measure the background (e.g., contribution from double layer capacitance), during which no $^*\text{CO}$ is observed by IR (Figure 4c). Based on the net anodic charge, the total $\Gamma_{^*\text{CO}}$ is determined to be 22.0 nmol cm^{-2} (geometric surface area) at the last moment of 0.4 V (corresponding to the IR peak area of 0.0322 , Figure S8). We note that this value is substantially higher than the atomic density of the single-crystal Au(111) surface (2.01 nmol cm^{-2}) as a result of the high surface roughness of the deposited Au.^{47,48}

Assuming the ratio between the IR peak area and surface concentration of each type of $^*\text{CO}$ is the same constant regardless of the electrode potential and surface concentration within

the IR intensity range of this study, we can then determine Γ_{*CO_B} under CO₂ reduction conditions using this relationship

$$\Gamma_{*CO_B} = \frac{A_{*CO_B}}{0.0322} \cdot 22.0 \text{ nmol cm}^{-2} \quad (3)$$

We are able to acquire steady-state A_{*CO_B} under CO₂ reduction reaction conditions by directly holding the electrode potential at -0.7 and -0.6 V (Figure S9a,b). We note that the accumulation of $*CO_B$ at -0.6 V is considerably slower than that at -0.7 V (Figure S9d,e) likely because of the slower restructuring to generate bridge sites (Figures 2 and S5). At -0.5 V where the restructuring is expected to be even slower, $*CO_B$ is not detected by IR within 5 min (Figure S9c). Therefore, we first perform CO₂ reduction at -0.7 V and then switch the electrode potential stepwise from 0.2 to -0.8 V to measure steady state Γ_{*CO_B} at each CO₂ reduction potential (-0.5 to -0.8 V) using the same Si//Au electrode (Figure S10a). In situ ATR-FTIR reveals gradual $*CO$ desorption and decline of A_{*CO_B} from 0.2 to -0.3 V (Figure S10b,c), agreeing with the previous observation of faster desorption at more negative potential in this region (Figure 3). As the potential becomes more negative, A_{*CO_B} evidently increases due to CO production from CO₂ reduction. The A_{*CO_B} values measured this way for -0.6 and -0.7 V match well with those obtained from directly setting the electrode at CO₂ reduction potentials (Figure S9e). The steady-state IR spectra at -0.5 , -0.6 , and -0.7 V are shown in Figure 4d and the corresponding Γ_{*CO_B} values are determined using eq 3 (Figure 4e). We then calculate the desorption rate of $*CO_B$, r_{*CO_B} , which is equal to the reaction rate of CO₂ reduction to CO via the $*CO_B$ pathway at steady state (CO re-adsorption at these potentials is unlikely on Au surfaces⁴⁶) using eq 4 (Figure 4e)

$$r_{*CO_B} = \Gamma_{*CO_B} \times k_d \quad (4)$$

The total CO production rate r_{CO} is given by eq 5

$$r_{CO} = j_{CO} \div F \div 2 \quad (5)$$

where j_{CO} is the CO partial current density (Figure 1c) and F is the Faraday constant. Comparing r_{*CO_B} to r_{CO} evidently shows that the $*CO_B$ pathway accounts for a significant proportion of CO₂ reduction to CO (Figure 4f). For example, at -0.5 V, approximately 38% of the reaction occurs on the bridge sites. We note that r_{*CO_B} increases with overpotential (Figure 4e), which is reasonable because of faster $*CO_B$ desorption at more negative potentials (Figure 3). It also appears that the contribution of $*CO_B$ to total CO production decreases at larger overpotential (Figure 4f), which may be related to the competition between $*CO_B$ and $*CO_L$ pathways.

Finally, we note that our aforementioned results are, as best we can tell, free from any significant influence of possible contamination. The spectroelectrochemical cell is

configured with an anionic exchange membrane and a high-purity graphite counter electrode, and the electrolyte is purified before use, following a recommended protocol to avoid contamination from metal impurities.⁵⁴ The electrochemically purified KHCO_3 electrolyte has been used in our lab for years to benchmark many different electrocatalysts for different electrocatalytic CO_2 reduction products with no sign of interference or contamination.^{12–18,76–83} To further confirm that our IR results are not caused by contamination from the electrolyte, we show that we can reliably reproduce the spectroelectrochemical results in Figure 2 with an ultrahigh-purity KHCO_3 electrolyte (Figure S11, see the Supporting Information for electrolyte preparation and purification details). Our Si//Au electrode manifests selective and stable CO_2 reduction activity characteristic of clean Au surfaces. Consistently, no metal impurities are detected on the surface by XPS either before or after electrocatalysis (Figure S12). More specifically, a control electrode without the topmost 20 nm Au layer has essentially no catalytic activity for CO_2 reduction to CO (Figure S1b), and no CO re-adsorption can be observed in the in situ ATR–FTIR measurement (Figure S13). This result adds additional support to the conclusion that the use of the Cr adhesion layer (or any contamination from it) is not responsible for the catalytic and spectroscopic features that we observe on the Si//Au electrode. In addition, the above-mentioned lack of $^*\text{CO}_B$ re-adsorption at 0.4 V (only $^*\text{CO}_L$ is observed) while we limit the prior CO_2 reduction potential to -0.5 V (Figure S6) is another piece of evidence against possible contamination in our system.

3. CONCLUSIONS

In summary, we have presented a time-resolved in situ ATR–FTIR study of Au-catalyzed electrochemical CO_2 reduction. From quantitatively probing bridge-bonded $^*\text{CO}$ species across a wide electrode potential range including the CO_2 reduction region, we have for the first time revealed that the Au surface can restructure during the initial phase of CO_2 reduction, and that the resulting bridge sites are catalytically active for CO production. Future studies employing advanced characterization techniques that are currently under rapid development, for example, high-resolution scanning probe microscopy imaging under electrochemical conditions, could help visualize the atomic-scale restructuring.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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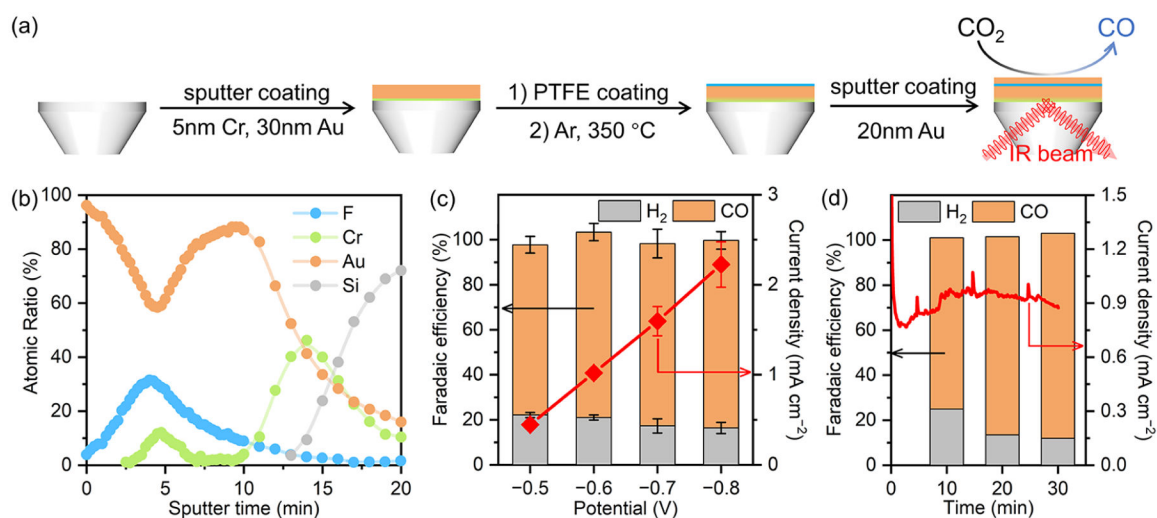


Figure 1.

(a) Preparation of the Si//Au electrode. (b) XPS depth profile of Si//Au. (c) CO₂ reduction performance of Si//Au in 0.1 M KHCO₃. Error bars represent standard deviations from measurements of three independently prepared electrodes. (d) FE and current density of 30 min CO₂ reduction electrolysis by Si//Au at -0.6 V.

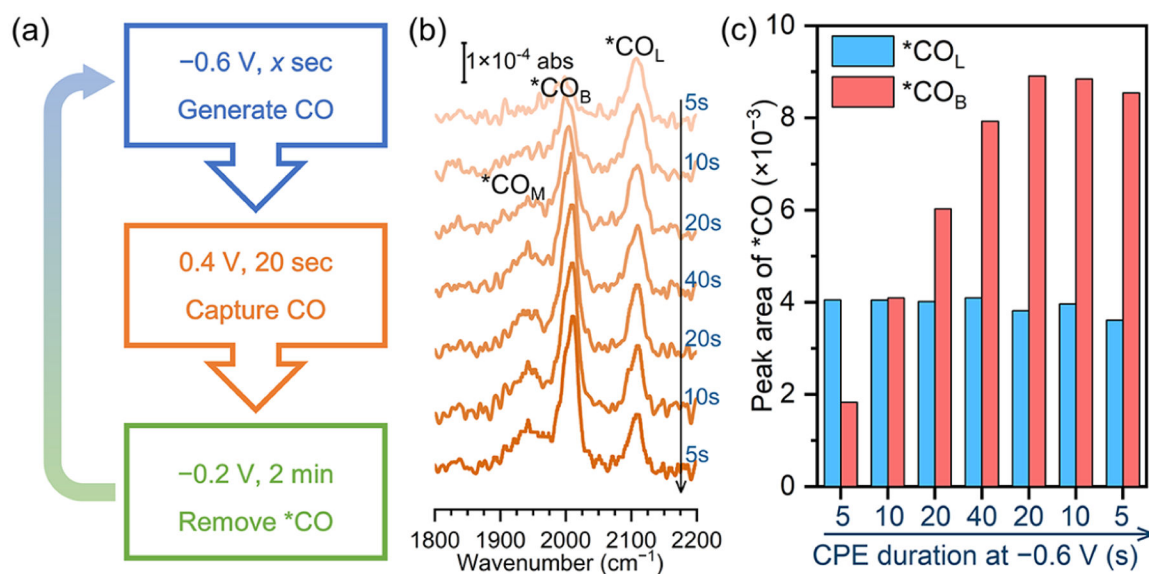


Figure 2.

(a) Potential step program for studying CO adsorption on Au. A total of seven cycles are performed in which the time at -0.6 V is varied in the sequence of 5, 10, 20, 40, 20, 10, and 5 s. (b) ATR-FTIR spectra recorded at 0.4 V of each potential cycle. Each spectrum is labeled with the time of CO_2 reduction at -0.6 V prior to the 0.4 V step. Each spectrum is averaged from 14 scans. (c) Peak areas of $^*\text{CO}_L$ and $^*\text{CO}_B$ observed at 0.4 V.

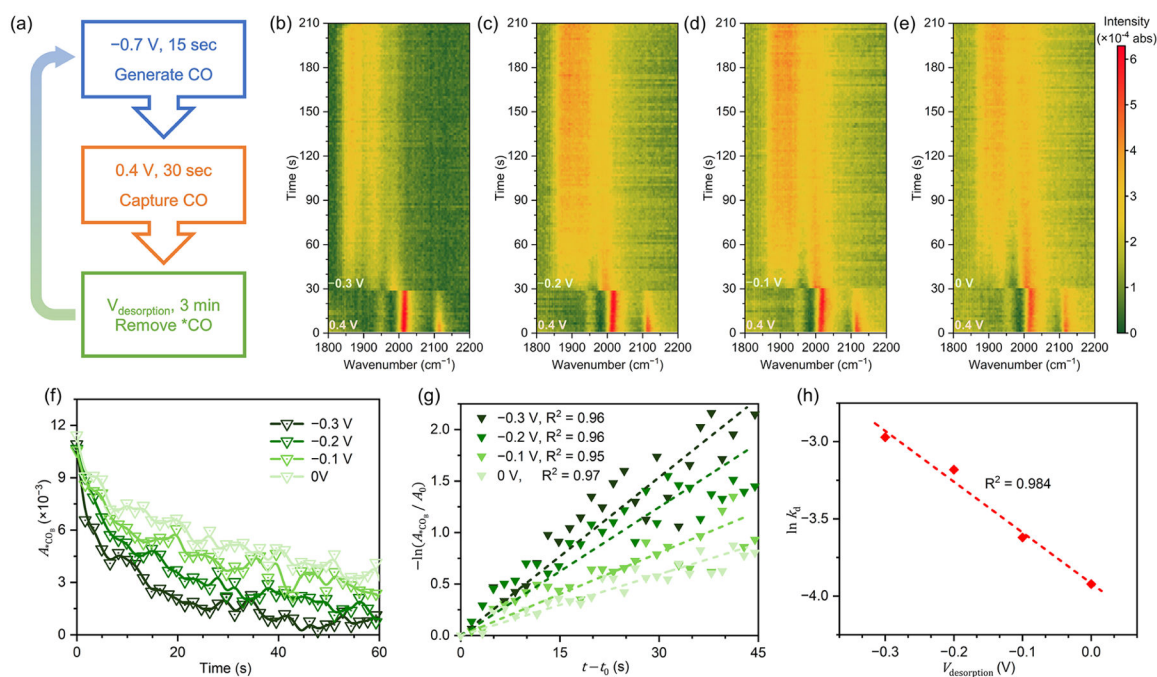


Figure 3.

(a) Potential step program for studying the desorption kinetics of *CO_B . A total of four cycles are performed in which $V_{\text{desorption}}$ is varied in the sequence of -0.3 , -0.2 , -0.1 , and 0 V. (b–e) Heatmaps of TRS ATR-FTIR spectra recorded at 0.4 V and $V_{\text{desorption}}$ for each cycle. (f) Desorption profiles of *CO_B at varied $V_{\text{desorption}}$. (g) Fitting desorption profiles to first-order kinetics. (h) Potential dependence of the desorption rate constants.

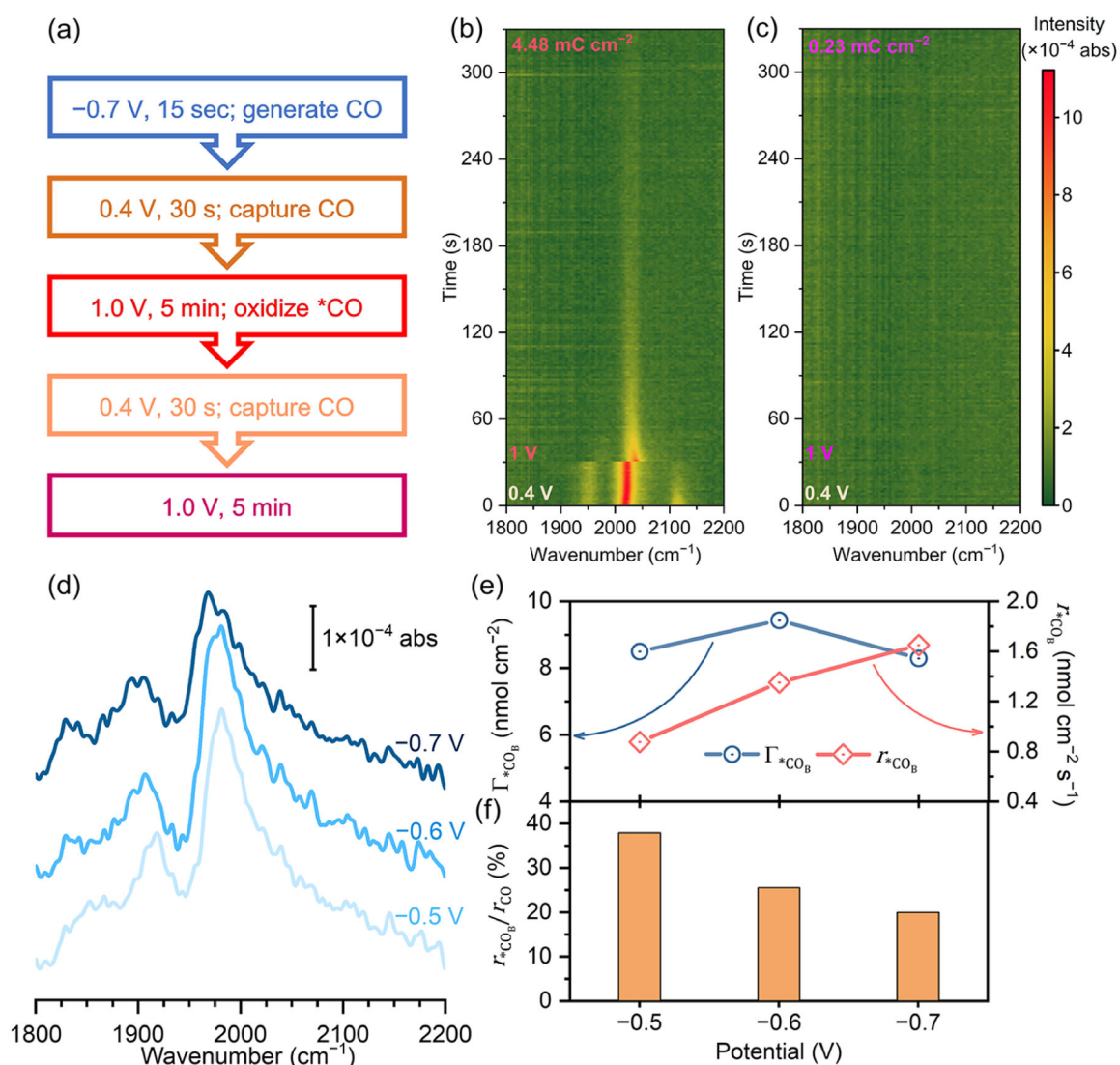


Figure 4.

(a) Potential step program for quantifying Γ^*_{CO} . After CO is generated at -0.7 V , the electrode potential is cycled twice between 0.4 and 1.0 V . (b-c) Heatmaps of TRS ATR-FTIR spectra recorded at 0.4 and 1.0 V for the first (b) and second (c) potential cycles. Amount of charge recorded at 1.0 V is labeled on each graph. (d) Representative ATR-FTIR spectra recorded at CO_2 reduction potentials. (e) Steady-state $\Gamma^*_{\text{CO}_B}$ and $^*\text{CO}_B$ desorption rate at CO_2 reduction potentials. (f) Potential-dependent share of the $^*\text{CO}_B$ pathway in total CO production.