Breaking Scaling Relationships in CO₂ Reduction on Copper Alloys with Organic Additives

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INTRODUCTION

The development of high-performing catalysts for sustainable and economically viable transformations remains a central goal of the chemical industry. Chemical transformations are controlled by thermodynamic and kinetic rate laws that manifest as linear scaling relationships. Such relationships relating structure, activity, and reaction conditions are established for a range of reactions, including H₂O oxidation and N₂, O₂, CO₂, and H₂O reduction performed on both heterogeneous and homogeneous catalysts. Because they provide theoretical or empirical trends for a particular chemical process, these scaling relationships not only help explain chemical reactivity but also guide the rational design of new and improved catalysts. Determining the underlying connections in chemical processes is particularly desirable toward deconvoluting fundamental selectivity limitations and targeting specific products. Typically, the experimental establishment and breaking of scaling relationships, including mapping of volcano plots, deals with a small set of data points constrained by inherent experimental limitations. Herein, high-throughput experimentation on 14 bulk copper bimetallic alloys allowed for data-driven identification of a scaling relationship between the partial current densities of methane and C₂+ products. This strict dependence represents an intrinsic limit to the Faradaic efficiency for C−C coupling. We have furthermore demonstrated that coating the electrodes with a molecular film breaks the scaling relationship to promote C₂+ product formation.
conduct studies that systematically and broadly vary select parameters. We have identified bulk alloying of Cu as an underdeveloped, though promising, strategy for catalyst optimization, with a large parameter space available based on the metal identity and composition, well suited for investigation using our high-throughput screening system. Additionally, organic additives represent an attractive orthogonal parameter of catalyst design. They can impact performance in a manner that has seldom been achieved by tailoring inorganic electrocatalyst composition or morphology alone.

Inspired by recent success using molecular films to enhance the selectivity of catalysts for CO₂R, herein, we describe the generation of a uniquely broad and systematic CO₂R catalyst database by combining a Cu bimetallic alloying strategy with the use of organic additives. Selectivity analysis highlights the impact of integrating high-throughput experimentation and data science to discover a power-law scaling relationship between partial current densities of CH₄ and C₂⁺, that is broken upon coating with an organic additive, demonstrating a fundamental limitation of CO₂R on Cu and a strategy to overcome it through hybrid inorganic–organic interfaces.

RESULTS AND DISCUSSION

To elucidate correlations in CO₂R, experiments were designed to observe a large dynamic range of catalyst properties while mitigating conflation with experiment parameters such as electrolyte composition and mass transport conditions. For the present work, we varied catalyst composition, applied potential, and molecular additive presence. The choice of Cu alloys was guided by our previous discovery that the alloying elements In, Co, Mn, and Zn alter the activity and selectivity of Cu in different ways, although that study was limited to the detection of H₂, CH₄, and C₂H₆. Studying Cu alloys with each of these elements and with different concentrations that span face-centered cubic (fcc) alloys and intermetallic phases (XRD of homogeneous alloys shown in Figures S12−S16), we sought to obtain a more comprehensive map of the reactivity of Cu-based alloy catalysts and to identify any systematic trends. The molecular additive, N,N′-ethylene-phenanthroline dibromide (1-Br₂), was selected based on its ability to enhance Faradaic efficiency (FE) and geometric partial current densities for C₂⁺ products upon forming a well-defined film on polycrystalline Cu, primarily composed of a para-para isomer of the one-electron reduced and dimerized phenanthroline (Figure 1a).

The catalyst performance with or without the additive was evaluated by chronoamperometry (CA) at a series of up to 6 potentials with subsequent product analysis using the batch reactor flow system illustrated in Figure 1. This system uses rapid electrolyte flow, as opposed to vigorous CO₂ bubbling, to generate suitable and reproducible mass transport conditions. The rapid concentration of reaction products enhances measurement throughput by enabling shorter electrolysis and faster chromatography compared to traditional methods. Hybrid metal–organic electrodes were prepared via electrodeposition of organic films on the polycrystalline metal electrode from an aqueous 0.1 M KHCO₃ buffered electrolyte containing 0.1 mM 1-Br₂. In total, experiments with 14 alloy catalysts and pure Cu provide electrochemical and partial current densities for 137 unique combinations of catalyst composition, additive presence, and applied potential, as shown for select Mn-doped catalysts in Figure 2a,b and for all catalysts in Figures S1 and S2.

Pairwise relationships of the geometric partial current density and the FE for representative products (Figure S5) highlight the effect of the combined strategy of alloying and organic films. The intrinsic modification of catalyst selectivity can be detected through analysis of the Pearson correlation coefficient of the logarithm of partial current densities. A close-to-unity positive correlation indicates that selectivity between the two products cannot be tuned with the parameters under consideration, which is indicative of a free-energy scaling relationship. A substantially negative correlation indicates a trade-off in selectivity, wherein enhanced formation of one product occurs at the expense of the other, which is indicative of kinetic competition for a shared reaction intermediate.

The large data set provided here via high-throughput experimentation enables the study of correlation coefficients and their modification (Figure 2). Previous work on polycrystalline Cu indicates that the kinetic regimes that govern the CO₂R product distribution differ with applied potential due to modulation of the energy landscape as a function of overpotential as well as second-order effects such as CO₂ mass transport and changes to the pH at the catalyst surface. To facilitate the observation of how the catalyst itself affects selectivity, we aim to mitigate the influences from the extrinsic effects by limiting the overpotential range (−0.84 to −1.1 V vs reversible hydrogen electrode, RHE) and using rapid electrolyte flow over flat catalyst films with a maximum current density of 15 mA cm⁻², which promotes uniform mass.

Figure 1. (a) The electrochemical reductive coupling of two 1-Br₂ molecules results in a mixture of two products; (b) The high throughput catalyst screening system where a select catalyst is positioned under a recirculating electrochemical batch reactor. After electrocatalyst operation, a robot sample handler (RSH) uses a syringe (orange) to extract aliquots from the headspace and then catholyte, with each aliquot injected into the respective analytical instrument (green, syringe positions in translucent orange) for gas or high-pressure liquid chromatography (GC, HPLC). The reference electrode (RE) is placed in the electrolyte inlet to the working electrode (WE) chamber, which is separated from the counter electrode (CE) chamber by a bipolar membrane (BPM).
transport and limits pH gradients in the electrochemical reactor. This potential range includes the onset of substantial partial current density for highly reduced products, making alteration of correlation coefficients in this range a prime target for controlling product selectivity with catalyst modification. We first demonstrate a Pearson correlation analysis to ascertain the extent by which high correlation coefficients can be lowered via variation in catalyst composition. For example, the box in Figure 2d with A = CH₄ and B = C₂⁺ shows a high correlation coefficient of 0.99 for these products when considering a series of 7 electrolysis experiments with a polycrystalline Cu catalyst in which partial current densities for both CH₄ and C₂⁺ products varied from approximately 1 μA cm⁻² to 3 mA cm⁻². The analogous analysis for polycrystalline Cu₉₈Mn₀₂ catalysts, both without additive and with 1-Br₂.
Cu−M alloys is summarized by the boxes with A = C₂+, and B = CH₄, where the correlation coefficient was calculated for each alloying element using various combinations of alloy composition and applied potential. The source data and their utilization of a correlation analysis are illustrated for the Cu−Mn system in Figure 2a-c.

The total number of electrolysis conditions and range of alloy compositions (x in Cu₁₋ₓMₓ) are as follows: 7 conditions with M = Co and x = 0.02 or 0.16; 10 conditions with M = In and x = 0.02 or 0.17; 16 conditions with M = Zn and x = 0.04, 0.13, 0.21, or 0.51; and 8 conditions with M = Mn and x = 0.02 or 0.16. Despite the variation in composition and potential within each of these Cu−M systems, each correlation coefficient remains in excess of 0.98, and in total, the correlation coefficient for all Cu−M alloys is not meaningfully changed from that observed with pure Cu. Figure 2f shows the aggregation of data for Cu and its alloys, demonstrating that a power-law relationship is closely followed over a broad range of composition and applied potential. This striking relationship, over 3 orders of magnitude, in partial current densities for CO, CH₄, and C₂+ products are converted to molar flux of CO* required to produce the respective products, whose normalization provides the ternary composition for inclusion in this figure. Each electrolysis experiment produces 1 data point that indicates the catalyst's selectivity with respect to the three reaction pathways highlighted in part a that start from the common CO* intermediate.

Figure 3. (a) Possible reaction mechanisms (selected from a multitude of variations previously proposed)¹¹ where pathways are highlighted in color with respect to their products in part b. There are two branching points between CH₄ and C₂+, products that could be responsible for the relationship observed in Figure 2. The strong relationship between the gray and green pathways is broken with the addition of molecular additives, implying a potential change in mechanism. (b) Summary of molar selectivity for reduction of the CO* intermediate. Measured partial current densities for CO, CH₄, and C₂+ products are converted to molar flux of CO* required to produce the respective products, whose normalization provides the ternary composition for inclusion in this figure. Each electrolysis experiment produces 1 data point that indicates the catalyst's selectivity with respect to the three reaction pathways highlighted in part a that start from the common CO* intermediate.

Experimental conditions used herein. Also, the alloying elements substantially alter other aspects of the product distribution, making this collection of catalyst electrodes particularly well-suited for inferring intrinsic reactivity trends; the catalyst morphology is kept relatively constant with respect to the compendium of results in the literature.¹¹,¹⁴,¹⁶,¹⁷,²⁰−²²,²⁴−²⁶,³⁰,⁴¹ For example, through study of well-defined Cu surfaces, Hori and others identified that the relative production of CH₄ and C₂H₄ is highly facet-dependent.¹⁵ The distribution of exposed facets of a polycrystalline fcc-phase metal electrode could be altered via alloying due to changes in growth kinetics and/or relative surface energies upon addition of the alloying element, which would in principle provide a method to break the CH₄/C₂+ scaling relationship by tuning catalyst composition. However, the observation that the scaling relationship holds over a broad range of alloy compositions with distinct product distributions indicates that facet selectivity is not the main contributor in the observed product selectivity.

CO₂R to highly reduced products such as CH₄ and C₂+, products proceeds via a common *CO intermediate.⁴²,⁴³ Methane synthesis is proposed to proceed via a Langmuir−Hinshelwood pathway, where a surface *H couples with *CO to form a *CHO or *COH intermediate that is further hydrogenated toward methane.⁴²,⁴⁴,⁴⁵ Meanwhile, the production of C₂+, products occurs via the coupling of two precursor *CO molecules, potentially involving intermediate *CHO or *COH adsorbrates.¹²,⁴³ The absence of CO−CH₄ or CO−C₂H₄ power-law relationships without the additive and the observation of a negative correlation coefficient between CO−ethylene in the presence of the additive are consistent with kinetic competition for a common intermediate (Figure S5). Despite variation in free CO produced (Figure S19), for the conditions tested, the assumed variation of CO concentration near the surface of the electrode is inconsequential with respect to the CH₄–C₂+ relationship. Therefore, although the specific mechanism or mechanisms remain debated and may involve multiple pathways depending on the

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morphology or crystal facet, the observed scaling relationship between CH\textsubscript{4} and C\textsubscript{2}\textsubscript{a} indicates that the relative kinetics at the branching point(s), remarkably, remain rigorously locked at the same ratio over the many catalysts and applied potentials tested herein. Breaking this dependence is highly desirable for improved selectivity for C\textsubscript{2}\textsubscript{a} products.

The Pearson correlation analysis was extended to the impact of the 1-Br\textsubscript{2} additive (Figure 2e), where the correlation coefficient for each set of conditions includes the aggregation of all catalyst compositions and potentials. Coating the catalysts using 1-Br\textsubscript{2} lowers the correlation coefficient for CH\textsubscript{4} and C\textsubscript{2}a from 0.99 to 0.74, a striking alteration whose implication is that, within the range of catalyst compositions considered in the present work, tuning the selectivity between CH\textsubscript{4} and C\textsubscript{2}a is only achieved in the presence of the additive, underscoring the importance of multimodal catalyst development.\textsuperscript{46}

The basis by which the additive disrupts the scaling relationship between CH\textsubscript{4}/C\textsubscript{2}a (Figure 2f) by increasing C\textsubscript{2}a production and suppressing CH\textsubscript{4} reveals a new frontier. Figure 3a illustrates the portion of the CO\textsubscript{2}R reaction network wherein branching ratios dictate whether the common CO* intermediate results in the generation of CO, CH\textsubscript{4}, or C\textsubscript{2}a products. Figure 3b highlights how catalyst modification with the organic additive moves product distribution almost completely away from CH\textsubscript{4} to the CO–C\textsubscript{2}a vector of the graph. While accessing the CO-rich portion of the graph is commonplace in CO\textsubscript{2}R electrocatalysis, the C\textsubscript{2}a-rich portion of the graph is only accessed in the presence of 1-Br\textsubscript{2}.\textsuperscript{11} The maximal selectivity was obtained with a Cu\textsubscript{0.32}Zn\textsubscript{0.15} alloy where 96% of the CO* intermediate was reduced to carbon-coupled products.

The breaking of the scaling relationship in the presence of 1-Br\textsubscript{2} cannot be explained by morphological changes or alloy segregation, as no nanostructuring was observed (Figures S7–S10, XRD in Figures S12–S16), therefore suggesting that the molecular additive improves selectivity via changes in the microkinetic pathway(s) in this system. The organic additive may affect CH\textsubscript{4}–C\textsubscript{2}a branching point(s) by (i) alleviating a rate limitation of the formation of the bound *CHO/*COH intermediate and lowering the barrier toward C–C coupling or (ii) promoting dimerization of the bound *CO relative to hydrogenation toward CH\textsubscript{4}. In either case, kinetic competition for the *CO would be enhanced in the presence of the additive, which is consistent with the observation of a large and negative Pearson correlation between C\textsubscript{2}a and CO (Figure 2e).

We additionally note that neither of these explanations for the mechanism underlyng the scaling law disruption has implications for the selectivity within the set of C\textsubscript{2}a products. As shown in Figure S6, additional correlations among these products are observed both in the absence and in the presence of the molecular additive, motivating future tuning of the catalyst system to tackle other branching points in the reaction network for enhanced control over product selectivity.

## CONCLUSION

High-throughput screening of the CO\textsubscript{2}R activity and selectivity of Cu alloys with Co, In, Mn, and Zn revealed the propensity of organic additive 1-Br\textsubscript{2} to enable the development of hybrid electrocatalysts that can reduce CO\textsubscript{2} to high-order products with improved activity and selectivity. The large data set led to the observation of a CH\textsubscript{4}–C\textsubscript{2}a scaling relationship that demonstrates a particularly robust link between these products over a large range of conditions. The CH\textsubscript{4}–C\textsubscript{2}a relationship represents an intrinsic limitation of selectivity tuning through alloying. However, it can be disrupted to favor C\textsubscript{2}a products by the presence of the organic additive, highlighting the potential of hybrid organic–inorganic catalysts to tune branching ratios in the CO\textsubscript{2}R reaction network. These observations highlight the importance of data-driven identification of relationships that provide mechanistic insights to guide the study of complex reactions and catalyst development. Disentangling the possible explanations of the combined mechanistic influence of the additive and alloying elements will require substantial further investigation that will be guided by the observed data relationships elucidated in this study.
Author Contributions
Y.L. and N.B.W. contributed equally to this work.

Notes
The authors declare no competing financial interest.

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