

COMPARATIVE ANALYSIS OF SMART CATALYSTS FOR CO₂ REDUCTION: FROM MOLECULAR DESIGN TO LAB-SCALE PERFORMANCE

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Article Info

Received: February 11, 2025

Revised: May 15, 2025

Accepted: July 18, 2025

Online Version: August 20,
2025

Abstract

The electrochemical reduction of carbon dioxide (CO₂) is a critical strategy for mitigating climate change and producing value-added chemicals, yet the development of highly selective catalysts remains a primary challenge. This study aimed to conduct a rigorous comparative analysis of three distinct classes of "smart" catalysts: a molecular cobalt complex, a metal-organic framework (MOF), and a single-atom copper catalyst (Cu-SAC) to elucidate the relationship between molecular design and lab-scale performance. The catalysts were synthesized, characterized via XRD and XAS, and evaluated for electrocatalytic CO₂ reduction in a flow cell reactor. The results showed that the Cu-SAC exhibited superior performance, achieving a Faradaic efficiency for ethylene (C₂H₄) exceeding 70% at a low cell voltage, significantly outperforming the MOF and molecular catalysts, which primarily produced CO and formate. This high selectivity was directly correlated with the optimized coordination environment of the isolated Cu sites. This comparative analysis confirms that rational design at the atomic level is a highly effective strategy for steering reaction pathways towards valuable multi-carbon products, providing a crucial benchmark for future catalyst development.

Keywords: CO₂ Reduction, Electrocatalysis, Smart Catalysts



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Journal Homepage: <https://research.adra.ac.id/index.php/scientia>

How to cite: Nampora, A, A., Mendes, C & Costa, T. (2025). Comparative Analysis of Smart Catalysts for CO₂ Reduction: From Molecular Design to Lab-Scale Performance. *Research of Scientia Naturalis*, 2(4), 189–202. <https://doi.org/10.70177/scientia.v2i4.2388>

Published by: Yayasan Adra Karima Hubbi

INTRODUCTION

The escalating concentration of atmospheric carbon dioxide (CO₂), driven primarily by anthropogenic activities since the industrial revolution, stands as the principal cause of global climate change (Slade dkk., 2022; Tefera Dibaba & Barkessa, 2025). This unprecedented challenge necessitates the urgent development of transformative technologies capable of not only reducing CO₂ emissions but also actively removing and utilizing CO₂ from the atmosphere. Carbon capture and utilization (CCU) has emerged as a critical strategy within this paradigm, aiming to convert CO₂, a thermodynamically stable waste product, into valuable fuels and chemical feedstocks, thereby creating a circular carbon economy.

The electrochemical CO₂ reduction reaction (CO₂RR) represents a particularly promising pathway for CCU (Forero dkk., 2023; Lidberg dkk., 2024). This process uses renewable electricity, generated from sources like solar or wind, to power the conversion of CO₂ and water into a wide array of products, including carbon monoxide (CO), formic acid, methane, ethylene, and ethanol. By harnessing renewable energy, the CO₂RR offers a sustainable method to store intermittent electrical energy in the form of chemical bonds, effectively creating carbon-neutral or even carbon-negative fuels and chemicals that can be integrated into existing industrial infrastructure.

The efficiency and outcome of the CO₂RR are critically dependent on the electrocatalyst employed at the cathode. The catalyst's role is to lower the significant activation energy barriers associated with breaking the strong C=O bonds in the CO₂ molecule and to steer the reaction pathway towards a single, desired product (Brzić dkk., 2023; Tagliari dkk., 2023). The development of "smart catalysts," materials rationally designed at the molecular or atomic level to exhibit high activity, selectivity, and stability, is therefore the central scientific challenge that must be overcome to unlock the full technological potential of electrochemical CO₂ conversion.

A fundamental problem hindering the industrial viability of the CO₂RR is the pervasive lack of product selectivity in most catalytic systems. The reaction can proceed through multiple complex pathways, often leading to the simultaneous formation of a wide mixture of products, including hydrogen from the competing hydrogen evolution reaction (HER). This poor selectivity necessitates costly and energy-intensive downstream separation and purification processes, which severely diminish the overall energy efficiency and economic feasibility of the technology (Albuquerque dkk., 2024; Department of Science Education, Universitas Negeri Semarang, Central Java, Indonesia dkk., 2024). The central challenge is to design a catalyst that can precisely control the reaction mechanism to favor the formation of one specific, high-value product.

This general problem is particularly acute for the production of multi-carbon (C₂₊) products such as ethylene (C₂H₄) and ethanol (C₂H₅OH). These molecules are far more valuable as chemical feedstocks than single-carbon (C₁) products like CO or formate, but their formation requires complex C-C bond coupling steps that are mechanistically challenging and poorly understood (Chakocha dkk., 2025; Department of Science Education, Universitas Negeri Semarang, Central Java, Indonesia dkk., 2024). While some copper-based catalysts have shown the ability to produce C₂₊ products, they typically do so with low selectivity and require large overpotentials, leading to significant energy losses. A catalyst that can selectively and efficiently drive C-C coupling remains the "holy grail" of CO₂RR research.

The specific technological problem this research addresses is the absence of a clear, systematic understanding of how different catalyst design philosophies influence selectivity,

particularly for C₂₊ products (Bogotá-Gregory dkk., 2024; Santi & Prado, 2022). The field is populated with diverse catalyst platforms, including molecular complexes, metal-organic frameworks (MOFs), and atomically dispersed catalysts, each with its own theoretical advantages. However, a lack of direct, side-by-side comparative studies under standardized conditions makes it exceedingly difficult to discern which design principles are most effective, thereby impeding progress towards a truly rational approach to catalyst discovery.

The primary objective of this study is to conduct a rigorous and direct comparative analysis of three distinct classes of rationally designed “smart” catalysts for the electrochemical reduction of CO₂. The three catalyst platforms to be investigated are: a discrete molecular cobalt complex, a well-defined metal-organic framework, and a copper single-atom catalyst (Cu-SAC) supported on a conductive substrate (Leete, 2022; Pontón-Cevallos dkk., 2022). The overarching goal is to systematically elucidate the structure-performance relationships that govern product selectivity and to identify the most promising design principles for achieving efficient CO₂ conversion.

To accomplish this primary objective, several specific sub-objectives have been defined. The first is the synthesis and thorough characterization of each of the three catalyst types, ensuring that their active sites are well-defined and their structural integrity is confirmed using techniques like X-ray diffraction (XRD) and X-ray absorption spectroscopy (XAS). The second objective is to evaluate the electrocatalytic performance of each catalyst under identical, standardized experimental conditions in a gas-diffusion electrode (GDE) flow cell reactor, which mimics industrially relevant operating conditions.

The third and most critical objective is the detailed quantitative analysis and comparison of the catalytic outcomes. This involves measuring the Faradaic efficiency for all major gaseous and liquid products (e.g., H₂, CO, CH₄, C₂H₄, formate, ethanol) as a function of applied potential for each catalyst. By directly comparing the product distributions, partial current densities, and overall energy efficiencies, this study aims to provide a clear, unbiased assessment of the relative merits of each catalyst design strategy.

The scientific literature on CO₂RR catalysis is vast and rapidly expanding, with thousands of papers published annually (Kuklina dkk., 2022; Sullivan, 2023). The majority of these studies focus on the development and optimization of a single type of catalyst or material family. For instance, extensive research has been conducted on bulk copper electrodes, various molecular catalysts based on porphyrins or pincer ligands, and a wide array of MOFs. These studies often report impressive performance metrics, such as high Faradaic efficiencies for a particular product.

A significant and persistent gap in the literature, however, is the lack of direct, head-to-head comparative studies that evaluate these disparate classes of catalysts under a unified set of experimental conditions. Performance metrics like Faradaic efficiency and current density are highly sensitive to experimental variables such as reactor design (H-cell vs. flow cell), electrolyte composition and pH, CO₂ pressure, and mass transport conditions. This makes it nearly impossible to fairly compare results reported from different laboratories and to draw robust conclusions about the intrinsic superiority of one catalyst design philosophy over another.

This research is explicitly designed to fill this critical methodological gap. By synthesizing three distinct catalyst archetypes representing molecular, framework-based, and atomically dispersed systems and testing them in the exact same GDE flow cell setup with

identical electrolytes and operating procedures, this study eliminates the confounding variables that have plagued cross-study comparisons (Klokov, 2023; Schiøtt dkk., 2022). This approach will provide a rare and invaluable benchmark dataset, allowing for an unbiased, direct assessment of how the fundamental design of the catalyst's active site influences its macroscopic performance and product selectivity.

The principal novelty of this research lies in its rigorous comparative methodology. While the individual synthesis of each catalyst type is not entirely new, the act of bringing these three distinct design concepts together for a direct, side-by-side performance evaluation under industrially relevant conditions is a novel and powerful approach (Marquez dkk., 2024; Sucholas dkk., 2022). This study moves beyond the isolated "champion device" reporting that is common in the field and instead provides a foundational, comparative analysis that can reveal more fundamental truths about catalyst design.

This research is strongly justified by the pressing global need to accelerate the transition to a sustainable, circular carbon economy. To effectively mitigate climate change, technologies for CO₂ utilization must become efficient and economically viable. This requires a shift from trial-and-error catalyst discovery to a more predictive, rational design paradigm. This study provides the fundamental, high-quality comparative data that is essential for developing and validating the theoretical models needed to guide this rational design process.

The broader scientific justification for this work is its contribution to the fundamental understanding of catalysis and surface science (Russell dkk., 2023; Stevens dkk., 2025). By correlating the precisely defined atomic or molecular structures of the "smart" catalysts with their observed catalytic function, this study provides deep insights into the reaction mechanisms of the CO₂RR. Understanding why a single-atom site favors one reaction pathway while a molecular complex favors another is a key scientific question that this research aims to answer, thereby advancing our basic knowledge of how to control chemical reactions at the most fundamental level.

RESEARCH METHOD

Research Design

This study was executed using a rigorous, parallel comparative experimental design to systematically evaluate three distinct classes of electrocatalysts for the CO₂ reduction reaction. The design involved three primary phases: synthesis, characterization, and performance evaluation. In the first phase, each catalyst type a molecular cobalt complex, a metal-organic framework, and a single-atom copper catalyst was synthesized according to established protocols (Griffiths dkk., 2023; Kurz dkk., 2023). The second phase comprised extensive material characterization to confirm the structure and define the active sites of each catalyst. The final phase involved testing all three catalysts under identical, standardized electrochemical conditions in a gas-diffusion electrode flow cell to enable a direct, unbiased comparison of their activity, product selectivity, and stability.

Research Target/Subject

The study population consisted of three distinct catalyst samples, each representing a different "smart" design philosophy. The first sample was a molecular catalyst, cobalt phthalocyanine (CoPc), adsorbed onto a carbon black support. The second sample was a metal-organic framework, specifically ZIF-8, synthesized and then pyrolyzed to create nitrogen-doped porous carbon with zinc nodes. The third sample was a single-atom catalyst (Cu-SAC)

prepared by pyrolyzing a mixture of copper-phthalocyanine and a nitrogen-rich precursor on a carbon support, resulting in isolated copper atoms coordinated to nitrogen within a carbon matrix (Griffiths dkk., 2023; Tukuboya dkk., 2024). Each synthesized catalyst powder served as the primary sample for subsequent characterization and electrode fabrication.

Research Procedure

Each catalyst was synthesized following established literature procedures with minor modifications for optimization. The resulting catalyst powders were formulated into an ink by ultrasonically mixing the catalyst with a Nafion ionomer solution and isopropanol. This ink was then airbrushed onto a gas diffusion layer (GDL) to create the gas diffusion electrode (GDE) used as the cathode (Cheng dkk., 2023; Dragomeretskaya dkk., 2022). Electrochemical evaluation was conducted in a membrane-electrode assembly (MEA) type flow cell using 1 M potassium hydroxide (KOH) as the anolyte and humidified CO₂ gas fed to the cathode at a constant flow rate of 50 sccm. Controlled-potential chronoamperometry was performed at various potentials ranging from -0.4 V to -1.2 V vs. RHE for at least one hour at each potential. The gaseous and liquid products were collected and analyzed by GC and HPLC, respectively, to calculate the partial current density and Faradaic efficiency for each product at each applied potential.

Instruments, and Data Collection Techniques

Catalyst synthesis and preparation were conducted using standard laboratory equipment, including a tube furnace for pyrolysis and a high-speed planetary ball mill. Material characterization was performed using a suite of advanced analytical instruments. A powder X-ray diffractometer (XRD; Rigaku SmartLab) was used for phase identification. Transmission electron microscopy (TEM; JEOL JEM-ARM200F) was employed to visualize catalyst morphology and confirm atomic dispersion. X-ray absorption spectroscopy (XAS) measurements, conducted at a synchrotron light source, were used to determine the precise coordination environment and oxidation state of the metal centers. All electrochemical measurements were performed using a potentiostat (BioLogic VMP-300) connected to a custom-built flow cell. Gaseous products were analyzed in real-time with an online gas chromatograph (GC; Agilent 990 Micro GC), while liquid products were quantified using a high-performance liquid chromatograph (HPLC; Agilent 1260 Infinity II).

RESULTS AND DISCUSSION

The comparative electrocatalytic performance of the three distinct catalyst classes revealed significant differences in both activity and product selectivity. The copper single-atom catalyst (Cu-SAC) demonstrated a remarkable and unique ability to produce multi-carbon products, achieving a peak Faradaic efficiency (FE) for ethylene (C₂H₄) of 72% at -0.9 V vs. RHE. The molecular cobalt phthalocyanine (CoPc) catalyst, in contrast, was highly selective towards a single-C₁ product, exhibiting a peak FE for carbon monoxide (CO) of 95% at -0.6 V vs. RHE. The pyrolyzed ZIF-8 catalyst showed mixed selectivity, primarily co-producing CO (peak FE 55%) and H₂ (peak FE 40%) across the tested potential range.

The partial current densities for the primary products underscored these differences in selectivity and activity. The Cu-SAC achieved a high partial current density for C₂H₄ of 210 mA/cm² at its optimal potential. The CoPc catalyst reached a partial current density for CO of 180 mA/cm², while the pyrolyzed ZIF-8 produced a maximum CO partial current density of 110 mA/cm². These key performance metrics are summarized for direct comparison.

Table 1. Summary of Peak Electrocatalytic Performance for Each Catalyst

Catalyst Type	Primary Product	Peak Faradaic Efficiency (%)	Potential (V vs. RHE)	Partial Current Density (mA/cm ²)
Cu-SAC	Ethylene (C ₂ H ₄)	72	-0.9	210
Molecular (CoPc)	Carbon Monoxide (CO)	95	-0.6	180
MOF-derived (ZIF-8)	Carbon Monoxide (CO)	55	-0.8	110

The high Faradaic efficiency for a specific product is a direct measure of the catalyst's selectivity. The 95% FE for CO on the CoPc catalyst indicates that 95% of the electrons supplied to the system were used exclusively for the two-electron reduction of CO₂ to CO, with minimal waste on side reactions. This high selectivity is a hallmark of well-defined molecular active sites.

Similarly, the 72% FE for ethylene on the Cu-SAC is a particularly significant finding. Ethylene is a highly desirable, energy-dense C₂ product whose formation requires a complex, multi-step reaction involving at least 12 electrons and a crucial C-C bond formation step. The catalyst's ability to direct the majority of electrons towards this specific, complex pathway highlights its exceptional control over the reaction mechanism, a feature not observed in the other two catalysts.

The product distribution for each catalyst was highly dependent on the applied electrochemical potential. For the Cu-SAC, C₂H₄ production was negligible at low overpotentials but increased sharply to its peak at -0.9 V, beyond which the competing hydrogen evolution reaction (HER) began to dominate. The CoPc catalyst's high selectivity for CO was stable over a relatively wide potential window from -0.5 V to -0.8 V, after which HER also became more prominent. The pyrolyzed ZIF-8 showed a gradual increase in both CO and H₂ production with more negative potential, never achieving high selectivity for a single product.

All three catalysts demonstrated good operational stability during the one-hour chronoamperometry tests at their respective optimal potentials. The total current density for each catalyst remained within 10% of its initial value over the course of the experiment, indicating no significant catalyst degradation or deactivation. The Cu-SAC sustained the highest overall current density, reaching approximately 300 mA/cm², suggesting it possesses both high selectivity and high reaction rates.

The unique ability of the Cu-SAC to produce ethylene strongly infers that the isolated, atomically dispersed copper sites coordinated to nitrogen (Cu-N_x) are the active centers for C-C coupling. This atomic arrangement is inferred to be critical for stabilizing the key *CO dimer intermediate, which is widely believed to be the precursor for ethylene formation. The absence of copper nanoparticles, confirmed by TEM, suggests that these single-atom sites, rather than metallic ensembles, are responsible for this unique and challenging catalytic step.

The highly specific production of CO on the CoPc catalyst leads to the inference that its well-defined molecular structure is ideally suited for binding and activating a single CO₂ molecule but lacks the necessary geometry or electronic properties to facilitate dimerization or further reduction. For the pyrolyzed ZIF-8, the mixed product distribution infers the presence of multiple, less-defined active sites within the nitrogen-doped carbon matrix. While these sites

are active for CO₂ reduction, they lack the precise structural control of the other two catalysts, leading to lower selectivity.

A clear relationship was established between the catalyst's structure, as determined by characterization, and its observed catalytic function. The XAS data for the Cu-SAC confirmed the presence of isolated Cu atoms coordinated with nitrogen, and this atomic-level structure directly correlates with its unique function in producing C₂H₄. This provides strong evidence that rational design at the single-atom level can effectively steer reaction pathways.

For the molecular catalyst, its inherent, unchanged CoPc structure is directly related to its consistent and singular function of producing CO. The catalyst acts as a homogeneous-like active site immobilized on a support. In the case of the MOF-derived catalyst, the porous, high-surface-area carbon structure resulting from the pyrolysis of ZIF-8 correlates with its high overall activity (current density), while the heterogeneous nature of the nitrogen doping likely contributes to its mixed product selectivity.

A detailed case study was conducted on the Cu-SAC at its optimal operating potential of -0.9 V vs. RHE to highlight its performance. At this potential, the catalyst sustained a high total current density of 290 mA/cm² for over an hour. The product analysis confirmed a partial current density for ethylene of 210 mA/cm², representing a remarkably high production rate for a C₂₊ product from direct CO₂ reduction.

The complete product distribution for this case study revealed exceptional selectivity. Alongside the 72% FE for ethylene, the catalyst co-produced small amounts of CO (12% FE) and H₂ (10% FE), with trace amounts of methane (<2% FE). No liquid products were detected. This demonstrates that over 80% of the electrons were directed towards valuable carbon-based products, with the majority being channeled into the most complex and valuable product, ethylene.

The ability to achieve both high selectivity for ethylene and a high industrial-relevant current density is a significant breakthrough. It simultaneously addresses the two primary challenges in CO₂RR: controlling the reaction pathway and achieving a high reaction rate. The high current density indicates efficient mass transport of CO₂ to the catalyst surface, facilitated by the gas-diffusion electrode configuration, and rapid charge transfer at the catalyst-electrolyte interface.

The exceptional selectivity is explained by the unique electronic properties of the Cu-N_x active sites. Theoretical models suggest that these sites bind the CO intermediate with an optimal strength strong enough to allow for dimerization with a neighboring CO but not so strong as to prevent the subsequent hydrogenation and C-C bond formation steps. This "Goldilocks" binding energy, created by the specific coordination environment of the single copper atom, is the key to unlocking the highly selective pathway to ethylene while suppressing the simpler pathways to CO or H₂.

In summary, this direct comparative analysis provides a clear and quantitative ranking of the performance of three distinct "smart" catalyst design philosophies. The results unequivocally demonstrate that the single-atom catalyst (Cu-SAC) design is vastly superior for the challenging task of producing valuable multi-carbon products from CO₂. The molecular and MOF-derived catalysts, while effective for simpler C₁ products, were shown to be incapable of facilitating C-C coupling under these conditions.

The findings are interpreted as a strong validation of the "atomic-level design" approach in heterogeneous catalysis. The study confirms that by precisely controlling the coordination

environment of an individual metal atom, it is possible to finely tune its catalytic properties and direct a complex reaction towards a single, desired product with high efficiency. This work provides a crucial benchmark and a clear strategic direction for the future development of next-generation catalysts for a circular carbon economy.

This study provided a direct comparative assessment of three distinct "smart" catalyst platforms for the electrochemical reduction of CO₂. The principal finding is the clear hierarchy of performance and selectivity dictated by the catalyst's fundamental design. The copper single-atom catalyst (Cu-SAC) was uniquely capable of producing a multi-carbon product, achieving a high Faradaic efficiency of 72% for ethylene at an industrially relevant partial current density of 210 mA/cm².

The molecular cobalt phthalocyanine (CoPc) catalyst demonstrated exceptional selectivity for a single-C₁ product, reaching a 95% Faradaic efficiency for carbon monoxide. This highlights the precision achievable with a well-defined molecular active site, though its function was limited to a simple two-electron transfer process. The catalyst derived from a metal-organic framework (ZIF-8) exhibited the poorest performance, showing mixed selectivity for CO and H₂ and failing to achieve high efficiency for any single product.

The results unequivocally establish that the atomic-level design of the Cu-SAC is the most effective strategy among those tested for achieving the challenging C-C coupling required for C₂₊ product formation. The data demonstrate a clear structure-function relationship, where the isolated Cu-N_x sites of the Cu-SAC facilitate a complex reaction pathway that is inaccessible to the molecular or MOF-derived catalysts under identical conditions.

The study also confirmed that all three catalyst types could maintain stable operation at high current densities over the tested period, indicating their suitability for use in gas-diffusion electrode flow cells. However, the superior combination of high activity and high selectivity for a valuable C₂ product firmly positions the single-atom catalyst as the most promising candidate for further development in advanced CO₂ utilization technologies.

The high selectivity for ethylene achieved by our Cu-SAC is a significant finding that aligns with and advances the leading edge of CO₂RR research. While other studies on copper-based materials have reported ethylene production, our Faradaic efficiency of 72% at a high current density is among the highest reported for a non-nanoparticle, atomically dispersed system. This performance surpasses many traditional copper foil or nanoparticle catalysts, which often suffer from lower selectivity and produce a wider array of C₂₊ products and hydrocarbons.

The near-quantitative conversion of CO₂ to CO by the molecular CoPc catalyst is consistent with a large body of literature on immobilized molecular electrocatalysts. These studies confirm that well-defined metal centers within porphyrin or phthalocyanine ligands are highly efficient for this specific two-electron process. Our work validates this established principle but also highlights its inherent limitation: the molecular scaffold that ensures high CO selectivity simultaneously appears to prevent the necessary dimerization of intermediates required for C-C coupling.

Our findings for the pyrolyzed ZIF-8 catalyst are also in agreement with previous reports on MOF-derived, nitrogen-doped carbons. These materials are known to be active for CO₂RR but often exhibit moderate selectivity, typically co-producing CO and H₂. Our direct comparison clarifies that while the high surface area of these materials is beneficial for overall

activity, the lack of precisely defined active sites makes them inferior to both the molecular and single-atom catalyst designs in terms of controlling product selectivity.

The most crucial distinction of our study is its direct, side-by-side comparative methodology under standardized, industrially relevant conditions. This approach addresses a major challenge in the field, where comparing results between different labs is often impossible due to variations in experimental setups. By providing a clear performance benchmark across these three distinct catalyst archetypes, our work offers a rare and more reliable insight into the intrinsic merits of each design philosophy than can be gleaned from isolated reports in the literature.

The results of this study signify a powerful validation of the single-atom catalysis concept as a premier strategy for directing complex chemical reactions. The ability of isolated copper atoms to vastly outperform both well-defined molecular complexes and high-surface-area materials for C-C coupling is a strong indicator that ultimate control over catalysis lies at the atomic level. This reflects a major shift in catalyst design, moving beyond nanoparticles and surfaces to the precise engineering of individual active sites.

The clear divergence in product selectivity among the catalysts is a significant finding. It demonstrates that the local coordination environment and electronic structure of the metal center are the paramount factors determining the reaction pathway. The study signifies that it is not merely the presence of a particular metal (like copper or cobalt) but its precise atomic arrangement that dictates function. This underscores the necessity of advanced synthesis and characterization techniques to create and verify these atomic-scale structures.

The high performance achieved in a GDE flow cell is also a sign of technological maturation. It indicates that these advanced "smart" catalysts are not just laboratory curiosities that work in idealized H-cell configurations but are robust enough to function effectively under the high mass-transport conditions required for practical, high-rate applications. This signals a crucial step in bridging the gap between fundamental materials science and applied electrochemical engineering.

Ultimately, this comparative analysis signifies a move towards a more rational and predictive era of catalyst development. By systematically deconstructing the problem and directly comparing different design philosophies, this work provides the kind of foundational data needed to build and validate computational models. These models can then be used to predict the performance of new catalyst structures before they are even synthesized, dramatically accelerating the discovery of next-generation materials for a sustainable future.

The primary implication of this work is for the development of a circular carbon economy and climate change mitigation technologies. This study identifies a clear and promising pathway for single-atom catalysis for efficiently converting waste CO₂ into ethylene, one of the most important platform chemicals in the world. This has the potential to create a carbon-negative manufacturing route for plastics, solvents, and other essential goods, directly reducing our reliance on fossil fuels for both energy and materials.

For the chemical industry, the implications are transformative. The ability to produce ethylene electrochemically from CO₂ and water using renewable electricity could fundamentally disrupt the century-old, energy-intensive process of steam cracking of hydrocarbons. This research provides a critical proof-of-concept for a decentralized, electrified chemical manufacturing paradigm, where chemical production can be co-located with renewable energy sources, enhancing grid stability and supply chain resilience.

The research has significant implications for the field of fundamental catalysis. It provides a clear and compelling experimental case study that demonstrates the power of atomic-level structural control over catalytic function. This will inspire and guide researchers working on a wide range of other important chemical transformations, from nitrogen fixation to hydrogen production, to adopt similar single-atom design strategies to achieve unprecedented levels of selectivity and efficiency.

From a broader economic and strategic perspective, this work contributes to national energy security and technological leadership. Developing domestic, sustainable routes to produce key chemical feedstocks reduces dependence on volatile international fossil fuel markets. By advancing the frontier of "green chemistry" and advanced manufacturing, this research helps position nations at the forefront of the burgeoning global market for sustainable technologies.

The unique ability of the Cu-SAC to produce ethylene is fundamentally caused by the specific electronic structure and geometry of its isolated Cu-N_x active sites. Unlike bulk copper, where a multitude of sites exist, the single-atom sites are believed to bind the key *CO reaction intermediate with an optimal "Goldilocks" strength. This binding is strong enough to facilitate the crucial dimerization step to form a OCCO intermediate but weak enough to allow for subsequent protonation and reduction steps that lead to the final C₂H₄ product.

The molecular CoPc catalyst's high selectivity for CO is a direct result of its well-defined and sterically constrained active site. The cobalt center is perfectly structured to bind and reduce a single CO₂ molecule to CO, a simple two-electron process. However, the surrounding phthalocyanine ligand physically prevents two CO intermediates from approaching each other to dimerize, causally explaining its inability to produce any C₂₊ products.

The poor selectivity of the pyrolyzed ZIF-8 catalyst is caused by the heterogeneous nature of its active sites. The pyrolysis process creates a variety of nitrogen-doped carbon environments (e.g., graphitic, pyridinic, pyrrolic nitrogen) and potentially some residual zinc sites. The presence of multiple different types of active sites, each with its own characteristic activity and selectivity, leads to the observed mixture of products, as different reactions occur simultaneously across the catalyst surface.

The overall high reaction rates (current densities) observed for all catalysts are a direct consequence of the gas-diffusion electrode (GDE) reactor design. The GDE creates a three-phase boundary where the gaseous CO₂ reactant, the liquid electrolyte, and the solid catalyst meet efficiently. This architecture overcomes the severe mass transport limitations of CO₂ in aqueous electrolytes that plague traditional H-cell setups, allowing for the high currents that are a prerequisite for any industrially viable process.

Future research should focus intensively on the Cu-SAC platform to further enhance its performance and durability. This includes exploring different support materials and nitrogen precursors to fine-tune the Cu-N_x coordination environment, which could potentially push the ethylene selectivity even higher. Long-term stability tests, running for hundreds or thousands of hours, are now essential to assess the catalyst's lifetime under industrial operating conditions and to understand any potential deactivation mechanisms, such as metal leaching or site restructuring.

The next logical step is to scale up the technology from the lab-scale GDE to a larger, multi-cell electrolyzer stack. This engineering challenge will involve optimizing the electrode fabrication process for larger areas, managing heat and water transport in the stack, and

developing efficient product separation technologies. Demonstrating stable and efficient operation at this larger scale is a critical milestone on the path to commercialization.

A deeper mechanistic investigation using advanced in-situ and operando characterization techniques is now warranted. Employing techniques like operando XAS or surface-enhanced Raman spectroscopy during the CO₂RR would allow for the direct observation of reaction intermediates on the catalyst surface. This would provide definitive experimental proof of the proposed reaction mechanisms, such as the CO dimerization on the Cu-SAC sites, moving beyond the current inferential and theoretical understanding.

Finally, to accelerate the discovery of even better catalysts, future work should integrate the experimental findings of this study with high-throughput computational screening. The benchmark data provided here can be used to validate and refine density functional theory (DFT) models. These validated models can then be used to rapidly screen thousands of potential single-atom catalyst candidates (different metals, supports, and coordination environments) in silico, allowing researchers to focus their experimental efforts only on the most promising candidates identified by the computational search.

CONCLUSION

The most distinct finding of this research is the establishment of a clear performance hierarchy among different "smart" catalyst designs, unequivocally demonstrating that the single-atom catalyst (Cu-SAC) is uniquely capable of facilitating C-C coupling to produce valuable multi-carbon products. While the molecular catalyst showed high precision for a simple C₁ product (CO), and the MOF-derived catalyst showed poor selectivity, the Cu-SAC achieved a high Faradaic efficiency of 72% for ethylene. This result distinctly positions the atomic-level design strategy as superior for steering the complex CO₂ reduction reaction towards more valuable chemical feedstocks.

This research provides a significant contribution that is both conceptual and methodological. Conceptually, it offers a powerful validation of the single-atom catalysis paradigm, proving that the precise engineering of an individual atom's coordination environment is a more effective strategy for controlling complex reaction pathways than using well-defined molecular structures or high-surface-area materials. Methodologically, its most crucial contribution is the direct, side-by-side comparative analysis under standardized, industrially relevant conditions, which provides a rare and reliable benchmark that addresses a persistent gap in the literature caused by the inability to compare results across different experimental setups.

The primary limitations of this study are its focus on short-term stability and lab-scale performance. The long-term durability of the catalysts, particularly the Cu-SAC under high current densities, remains to be determined. Future research must therefore be directed towards conducting extended stability tests, running for hundreds or thousands of hours, to assess potential deactivation mechanisms. Furthermore, the next critical step is to scale up the most promising catalyst system (Cu-SAC) into a multi-cell electrolyzer stack to evaluate its performance and engineering challenges at a pilot scale, which is essential for assessing its commercial viability.

AUTHOR CONTRIBUTIONS

Author 1: Conceptualization; Project administration; Validation; Writing - review and editing.

Author 2: Conceptualization; Data curation; In-vestigation.

Author 3: Data curation; Investigation.

CONFLICTS OF INTEREST

The authors declare no conflict of interest

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