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Electrocatalyst Microenvironment Engineering for Enhanced Product Selectivity in Carbon Dioxide and Nitrogen Reduction Reactions

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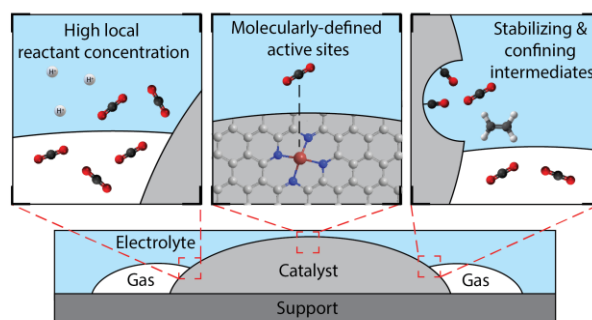
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TOC graphic



Abstract

Carbon and nitrogen fixation strategies are regarded as alternative routes to produce valuable chemicals used as energy carriers and fertilizers that are traditionally obtained from unsustainable and energy-intensive coal gasification (CO and CH₄) Fischer-Tropsch (C₂H₄) and Haber-Bosch (NH₃) processes. Recently, the electrocatalytic CO₂ reduction reaction (CO₂RR) and N₂ reduction reaction (NRR) have received tremendous attention with the merits of being both efficient strategies to store renewable electricity while providing alternative preparation routes to fossil-fuel driven reactions. To date, the development of the CO₂RR and NRR processes is primarily hindered by the competitive hydrogen evolution reaction (HER), however the corresponding strategies for inhibiting this undesired side reaction are still quite limited. Considering such complex reactions involving three gas-liquid-solid phases and successive proton-coupled electron transfers, it appears meaningful to review the current strategies for improving product selectivity in light of their respective reaction mechanisms, kinetics, and thermodynamics. By examining the developments and understanding in catalyst design, electrolyte engineering and three phase interface modulation, we discuss three key strategies in improving product selectivity for the CO₂RR and NRR: i) targeting molecularly defined active sites, ii) increasing the local reactant concentration at the active sites, and iii) stabilizing and confining product intermediates.

Keywords Carbon Dioxide, Nitrogen, Electrochemical Reduction, Microenvironment, Selectivity, Electrocatalyst, Electrolyte, Three-phase Interface

1. Introduction

Many of today's environmental, economic, and societal issues are related to the transformation of two inert gases, N_2 and CO_2 . The transformation of N_2 *via* the Haber-Bosch process accounts for ~2% of the world energy consumption, providing nitrogen fertilizers required to sustain the current global food production. Meanwhile, the amount of CO_2 released into the atmosphere from the combustion of fossil fuels has reached unprecedented levels, further accelerating climate change.¹⁻⁵ Both CO_2 and nitrogen undergo complex environmental cycles (**Figure 1a-b**), increasing the challenges associated with their capture and conversion. Implementing CO_2 and N_2 sustainable cycles and minimizing their environmental impact is critical, as recently highlighted in the latest IPCC report or in Europe in the EU green deal and Fit for 55 package.⁶⁻⁸

The electrochemical conversion of CO_2 and N_2 into value-added products or net zero commodities such as materials, renewable fuels and energy vectors, appears as an appealing solution in this context, as it can utilize sustainable sources of electricity powered by solar, wind, wave, and hydro energy to promote reactions currently carried out using fossil fuels. This approach would provide a carbon-neutral route to C- and N- containing products whilst enabling the efficient storage of intermittent renewable sources of electricity as chemical bonds, largely overperforming battery storage energy efficiency.⁹

The main hurdle to developing energy efficient processes for converting nitrogen to ammonia and carbon dioxide to energy dense products such as hydrocarbons is selectivity. The chemical inertness of these reactants disadvantage their transformation compared to more kinetically facile reactions such as the hydrogen evolution reaction (HER). Furthermore, selectivity is one of the most challenging aspects to address when developing electrocatalysts to mediate the CO_2 reduction reaction (CO_2RR) and the nitrogen reduction reaction (NRR). In both cases multiple reaction products are typically observed, resulting from the reduction of CO_2 and N_2 themselves as well as from the proton sources used to mediate these reduction reactions, which involve successive coupled electron-proton transfers. In this regard, the electrocatalyst microenvironment plays a vital role and can be engineered to improve selectivity through three key strategies: i) targeting a narrow distribution of molecularly defined active sites, ii) increasing the reactant/proton ratio at the three-phase interface where the reaction takes place to lower the undesired formation of H_2 and iii), the stabilisation and confinement of reaction intermediates in the electrode vicinity to favour the formation of multi-electron reduction products.

While there exists an extensive amount of literature in both the CO₂RR and NRR fields, including several recent reviews of specific subtopics,¹⁰⁻¹³ we target in this review to illustrate through a handful of selected examples the key strategies for increasing selectivity towards value-added products.

After a brief explanation of the kinetic and thermodynamic origins of multiple product generation in the CO₂RR and NRR, we discuss the key factors in catalyst design in steering product selectivity, namely nano-structuring, surface functionalisation, control of crystal size and facets, and single site engineering. We then explore the impact of the electrolyte on activity at the electrode surface, including aspects such as pH, the alkali metal cation, and the use of novel electrolytes. The final section focuses on the implementation and optimisation of triple-phase interfaces to improve local reactant concentration and mass transport. We conclude with our perspectives on this rapidly growing topic and where we envisage future challenges and opportunities to lie.

It is important to note that the NRR field has been strongly affected by a series of false positives, and a standardized set of experiments have been outlined to identify, quantify and eliminate experimental artefacts.¹⁴ Ammonia contamination may arise from sources such as the air, chemicals and the experimental set-up, which is particularly significant when the quantity of ammonia produced in NRR is very low. Additionally, labile nitrogen-containing compounds such as nitrates, nitrites, nitrogen oxides and amines are often present in the N₂ gas stream, the air, and the catalyst itself. To reliably attribute ammonia production to the NRR, quantitative isotope measurements with ¹⁵N₂ gas and the removal of impurities from the gas stream are imperative. To preserve a fair comparison of performance between catalytic materials, in this review we present only examples that follow the guidelines provided in the above reference 14, unless clearly stated otherwise.

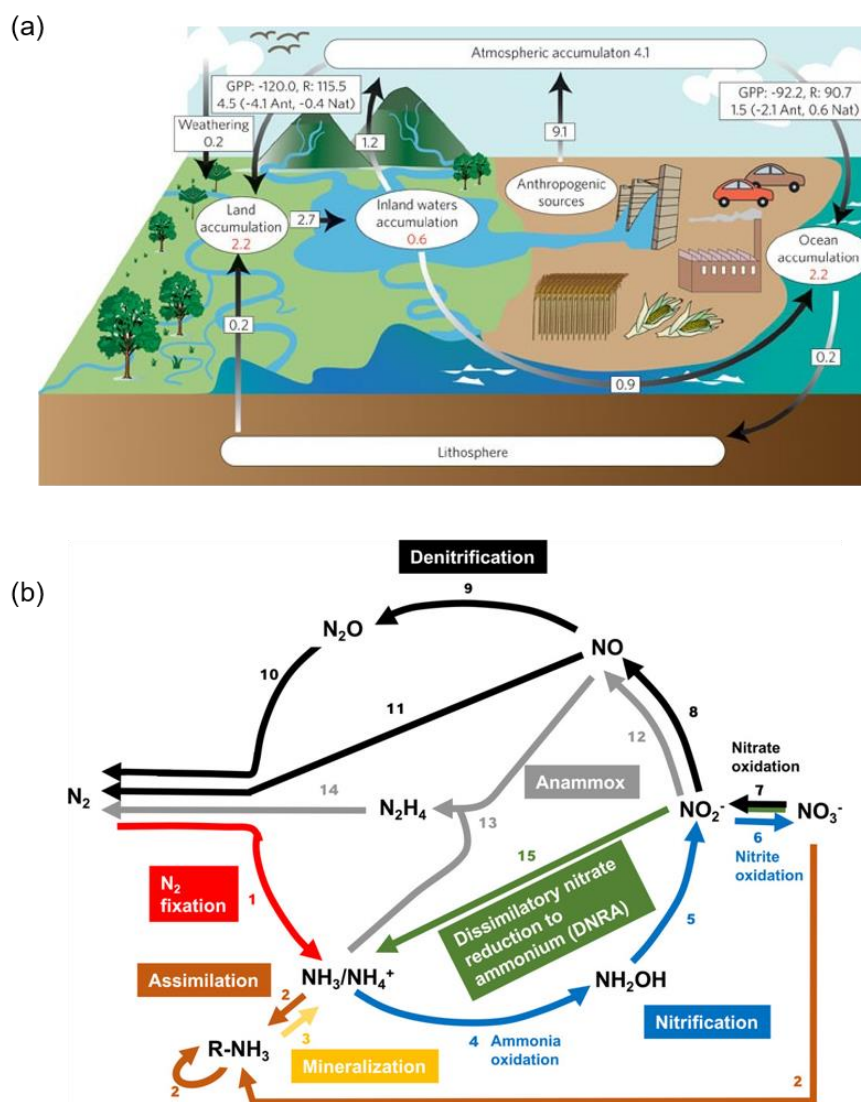


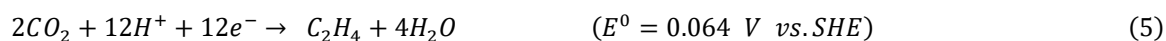
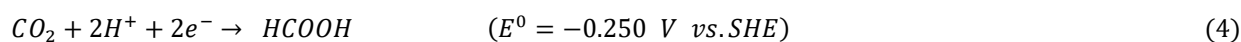
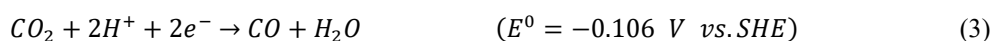
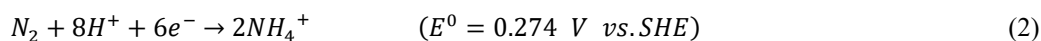
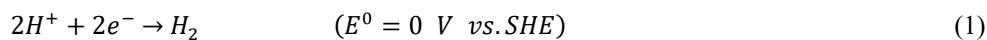
Figure 1. (a) Scheme of the carbon cycle. Reproduced from *Nat. Geosci.* **2009**, *2* (9), 598-600. Copyright 2009 Springer Nature. (b) Cycle of biologically driven N-transformations that occur in natural and human-influenced terrestrial and marine environments. Reproduced from *Chem. Rev.* **2020**, *120* (12), 5308-5351. Copyright 2020 American Chemical Society.

2. Mechanistic and thermodynamic origin of multiple product generation in CO₂RR and NRR

Both the CO₂RR and NRR to value-added products involve multiple successive proton-coupled electron transfers (Table 1), which represent a significant kinetic challenge to overcome to achieve high selectivity, particularly compared to the more kinetically facile two-electron hydrogen generation reaction.¹⁵⁻¹⁷ This kinetic challenge is further complexified by the low availability of the reactants, as both CO₂ (~33 mM at P_{CO₂}=1atm) and N₂ (~0.7 mM at P_{N₂}=1 atm) have typically poor solubility in water.¹⁸ In the context of the CO₂RR to multi-carbon products, the low solubility of the primary reaction products such as CO also decreases the overall catalyst selectivity for multi-carbon products

which result from the subsequent reduction of these primary products.

Table 1. Selected standard potentials of CO₂ and N₂ in aqueous solutions (V vs. SHE) at 1.0 atm and 25 °C, calculated according to the standard Gibbs energies of the reactants in reactions. Reproduced from *Chem. Soc. Rev.* **2014**, *43* (2), 631-675, copyright 2014 Royal Society of Chemistry, and *Chem. Soc. Rev.* **2019**, *48* (24), 5658-5716, copyright 2019 Royal Society of Chemistry.



In addition, a thermodynamic challenge is associated with the CO₂RR, since proton reduction (HER) is more thermodynamically favourable than the reduction of CO₂ to most products (**Figure 2a** and **Equations 3-6**).¹⁹⁻²¹ Although less critical in the case of NRR, the standard electrochemical potential for the proton reduction reaction is yet close to that of the nitrogen reduction reaction (NRR) at 0.057 V vs. SHE (**Equation 2**).²² The intrinsic stronger binding of H atoms over N₂ on most metal surfaces, highlighted in **Figure 2b**, further illustrates the challenge to increase NRR selectivity vs. HER.²³

This illustrates the three main challenges (thermodynamic, kinetic, or related to the mass-transport of the reactants and primary reaction products) that must be overcome to reach high selectivity in the CO₂RR and NRR. We will review in the next sections the three main axes currently explored toward that goal, focusing on catalyst design, electrolyte engineering and three-phase interface modulation.

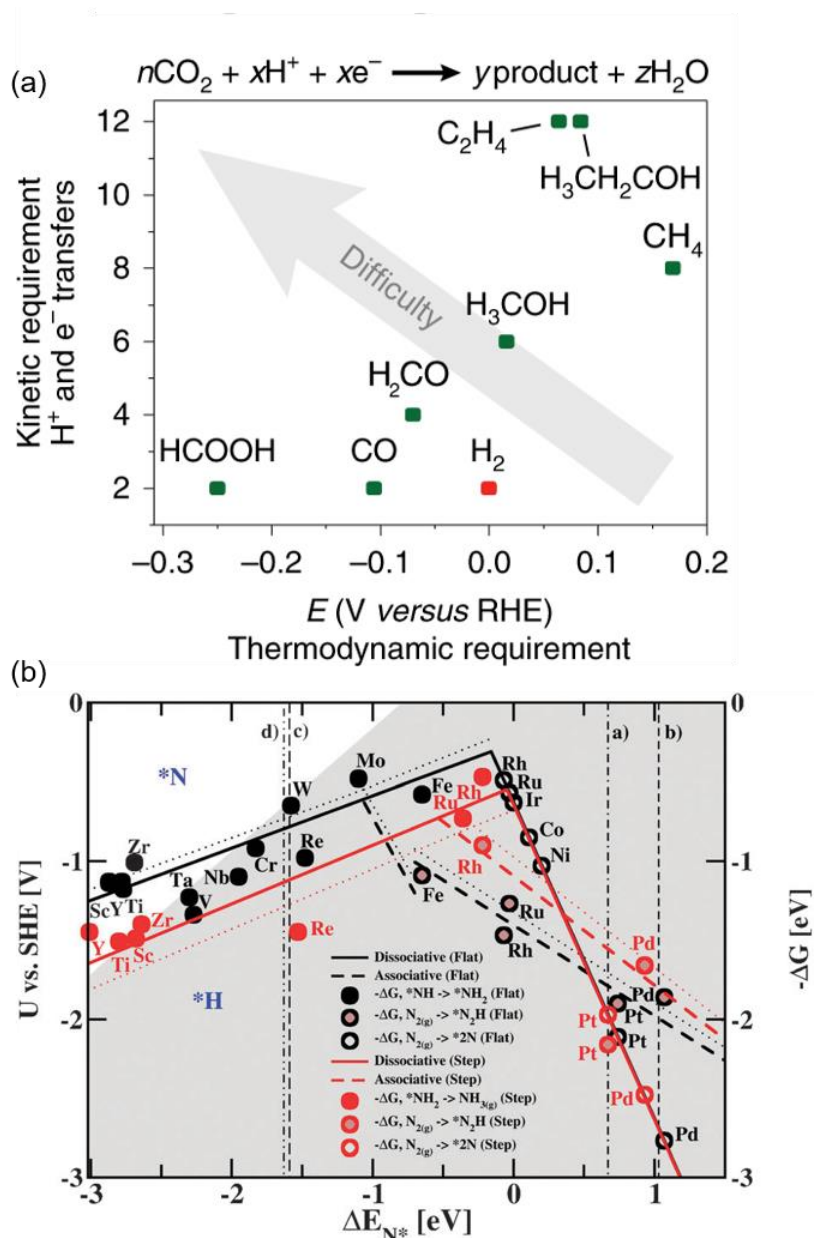


Figure 2. (a) Kinetic versus thermodynamic requirements of various CO_2 reduction reactions. The plotted values are based on the reaction equation given above the graph, made stoichiometric according to the product composition. Reproduced from *Nat. Mater.* **2019**, *18* (11), 1222-1227. Copyright 2019 Springer Nature. (b) Combined volcano diagrams (lines) for the flat (black) and stepped (red) transition metal surfaces for the reduction of nitrogen with a Heyrovsky-type reaction, without (solid lines) and with (dotted lines) H-bond effects. Reproduced from *Phys. Chem. Chem. Phys.* **2012**, *14* (3), 1235-1245. Copyright 2012 Royal Society of Chemistry.

3. Increasing selectivity via catalyst design

3.1 Catalyst nano-structuring for improved mass transport

Advancements in nanotechnology and characterisation techniques have enabled a plethora of morphologies to be explored to improve catalytic activity and product selectivity. Porous materials have attracted particular attention due to their effect on the local chemical environment, including local

pH and the mass transport of the reactant and intermediates.^{24, 25} The ability to increase the number of effective active sites, both by maximising surface area and facilitating the accessibility of such sites, makes porosity useful and interesting across a broad range of fields.²⁶ Such effects are especially crucial when considering the poor solubility of CO₂ and N₂ in aqueous electrolytes, which cause mass transport limitations and barriers to high activity and selectivity.

Hierarchical porous networks are found commonly in biological organisms as a strategy to mitigate mass transport limitations in the utilisation of nutrients.²⁷ The three-dimensional networks were replicated in early work by Huan *et al.* who used gold nanodendrites for electrochemical sensing.²⁸ Their application in catalysis has recently appeared as an efficient strategy to increase current densities and catalyst selectivity in small molecule electroreduction and oxidation.

The dynamic hydrogen bubble templating (DHBT) method has been the most prominent technique to create such hierarchical porosity, which was recently comprehensively reviewed by the Bhargava group²⁹ and specifically for CO₂RR materials by the Broekmann group.³⁰ The process involves the electrodeposition of a metal from aqueous solutions of the respective cations, while co-generated hydrogen bubbles act as a dynamic template to create a metal foam. As the bubbles nucleate, grow and detach, a hierarchical pore structure forms with layers of pores of increasing diameter (**Figure 3a**), including micropores in the submicron range and macropores 10-100 μm.²⁹ The DHBT technique is relatively simple, requiring aqueous solutions and no need for organic or inorganic templates (as in traditional metal foam synthesis),³¹ high temperatures, high pressures or uncommon equipment. Nonetheless, additives such as citrate are common to influence crystal growth.³²⁻³⁴ Bi- and multi-metallic catalysts are also possible by co-electrodeposition, galvanic replacement, stepwise electrodeposition or spontaneous decoration.²⁹ For example, many studies for CO₂RR have coupled copper with one other metal such as Ag, Sn, In or Zn.³⁵⁻³⁹

By fine-tuning parameters such as proton source and concentration, applied overpotential or current density, substrate material, and the metal source and concentration, the nanostructure can be carefully controlled and optimised. Broekmann and co-workers produced a dendritic Cu-based DHBT foam, and demonstrated a strong dependence of the C₂-product selectivity on the surface pore size diameter, with the optimal size being between 50 and 100 μm.⁴⁰ They identified the temporal trapping of gaseous intermediates inside these pores as the key to product selectivity. Intermediates such as CO

and C_2H_4 , which would otherwise be released into the bulk electrolyte, were entrapped in the pores of the foam catalyst, causing them to further react to form C_2H_6 (**Figure 3b**). At -0.8 V vs RHE they achieved a 55 % Faradaic efficiency for C_2 -products.

Such dendritic structures with large surface areas are common in this synthesis due to the deposition taking place at high current densities and therefore in the diffusion limited regime. Copper and oxide derived copper dendrites have had particular interest due to their apparent selectivity for multicarbon products.⁴¹⁻⁴⁴ Huan *et al.* produced a dendritic CuO material from DHBT that could be used both as a CO_2R and OER catalyst.^{45, 46} It consisted of a triple layer structure with a metallic Cu core covered by layers of Cu_2O and CuO (**Figure 3c**). In electrocatalytic conditions, the CuO material is reduced to metallic Cu, generating nano-Kirkendall voids within the dendrite structures. Such voids, that appear at the copper/copper oxides interface upon reduction, are termed nano-Kirkendall voids as they appear as a consequence of the very different diffusivities of Cu and O atoms. The overall external shape of the material is maintained upon reduction but cavities are generated under its external layer due to the lower density of Cu with respect to the original copper oxides.⁴⁷ These gas-accessible voids were proposed to enhance the confinement of secondary CO_2RR products, such as CO, resulting in FE_{C_2+} over 50 %. By using the catalyst in a continuous flow electrolyzer, they were able to reach a stable current of 25 mA/cm² with 2.95 V, equating to 21 % energy efficiency for hydrocarbon production. By coupling the cell to a photovoltaic cell, they achieved a 2.3 % solar-to-hydrocarbon efficiency.

DHBT foams for single-carbon products such as CO and formate have also been reported. A silver-foam with needle-shaped features in the mesopores was produced by using a citrate additive to control growth on the nanometer scale.³² Between -0.3 to -1.2 V vs RHE 90% Faradaic efficiency for CO was observed, however at higher over-potentials they produced C_2 -products, with 51% CH_4 at -1.5V (**Figure 3d**). This unusual activity for Ag was attributed to the catalyst morphology and nanostructure increasing *CO surface concentration and residence time. Recent work by Mayer and co-workers exemplifies the advantages of the simplicity of the DHBT method. In a one-step synthesis they used waste industrial Cu-Sn bronze as a material precursor to deposit a mesoporous $Cu_{10}Sn$ foam.⁴⁸ They achieved over 85 % Faradaic efficiency for CO at -0.8 V vs RHE, over double that of the plain Cu-Sn bronze, with partial current densities three times higher. Du *et al.* prepared a nanoporous tin DHBT

foam on a tin substrate and achieved a Faradaic efficiency for formate of 90 % with current densities of 23 mA/cm^2 .⁴⁹

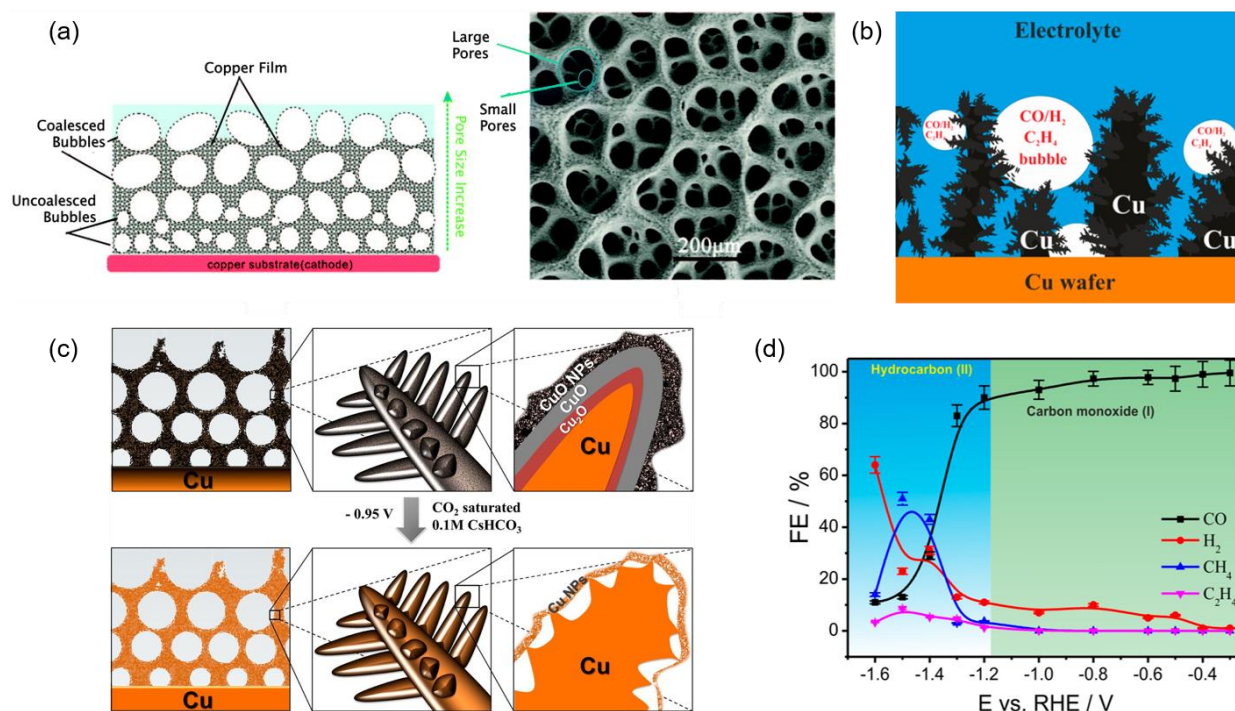


Figure 3. (a) Schematic illustration and SEM image of a copper DHBHT foam, demonstrating the hierarchical pore structure. Reproduced from *Chemical Communications* **2015**, 51 (21), 4331-4346, copyright 2015 Royal Society of Chemistry, and *Journal of The Electrochemical Society* **2013**, 160 (10), D441-D445, copyright 2013 IOP Publishing. (b) Schematic illustration of gaseous CO₂R intermediates (CO and C₂H₄) and by-products (H₂) trapped within the porous Cu foam catalyst. Reproduced from *ACS Catalysis* **2016**, 6 (6), 3804-3814. Copyright 2016 American Chemical Society. (c) Schematic illustration of a dendritic CuO DHBHT-foam before (top) and after (bottom) CO₂ electroreduction in 0.1 M CsHCO₃, showing the material reduction to metallic Cu and the formation of nano-Kirkendall voids. Reproduced from *Proc Natl Acad Sci U S A* **2019**, 116 (20), 9735-9740. Copyright 2019 Proceedings of the National Academy of Sciences. (d) Potential dependent product distribution of the CO₂RR using a Ag-DHBHT-foam catalyst by faradaic efficiency, showing the formation of hydrocarbons at potentials more negative than -1.2 V vs RHE. Reproduced from *ACS Catalysis* **2018**, 8 (9), 8357-8368. Copyright 2018 American Chemical Society.

Other morphology-based strategies have been utilised to modulate mass transport in CO₂ reduction, including the application of nanostructures such as nano-wires, sheets, needles, cones or tubes. Burdyny *et al.* explored the effect of nanomorphology of a silver catalyst on gas-evolution and subsequently bubble-induced mass transport.⁵⁰ By combing mathematical modelling and experimental observations using a dark field microscope, they compared bubble formation on nanoparticles, nanorods and nanoneedles, and found a mean bubble diameter of 97, 31 and 23 μm respectively. They illustrated that the generation of smaller bubbles improved long-range mass transport of CO₂, resulting in a small diffusion thickness and a 4-fold increase in limiting current density of CO production

(**Figure 4a**). Surendranath and co-workers synthesised gold inverse opal thin films and found that changing the mesostructure by increasing porous film thickness could diminish HER 10-fold whilst maintaining activity for CO₂ to CO, enhancing the faradaic efficiency for CO from less than 5 % to over 80 %.⁵¹ They attributed this to the formation of diffusional gradients. Studies into nanocavities and their performance and mechanism of action have emerged in recent years. Yang et al. utilised finite-element method simulations and experimental measurements on a multihollow cuprous oxide catalyst.⁵² Analysis from X-ray absorption studies and operando Raman spectra indicated that the pore cavities confined *CO intermediates, which bound to Cu⁺ sites and locally protected them against reduction during CO₂RR (**Figure 4b**), as well as promoted C-C coupling. They achieved a C₂₊ product Faradaic efficiency of 75 % and partial current density of 267 mA cm⁻².

As N₂ electroreduction is a comparatively less mature field with its own unique challenges, studies into morphological effects on catalytic activity and selectivity are less extensive. Although a range of nanostructures exist amongst the literature,⁵³ specific insight into the role morphology plays in catalysis is limited. Wei et al. loaded ruthenium nanoparticles onto carbon nanotubes, which were also applied as the gas diffusion electrode.⁵⁴ Despite using a typical H-cell set-up, the GDE structure allowed N₂ gas to be flowed through the GDE and porous catalyst, instead of being solely solubilised in the electrolyte as illustrated in **Figure 4c**. They achieved a NH₃ yield rate of 2.1 nmol/cm²s and Faradaic efficiency of 13.5 %.

A great range of nanostructures have been applied to the CO₂RR and NRR to regulate mass transport, and although strong correlations between structure and performance have been made, their mechanisms of action are often highly complex and difficult to define. Most theories focus on the mass transport of reactants and intermediates either through improved diffusion and convection or through their physical confinement in the catalyst pores. Considerable progress has been made by combining computational and experimental research to define and improve catalyst nanomorphology, especially in the CO₂RR field, however their application to new materials and fields such as NRR is still an open area of research.

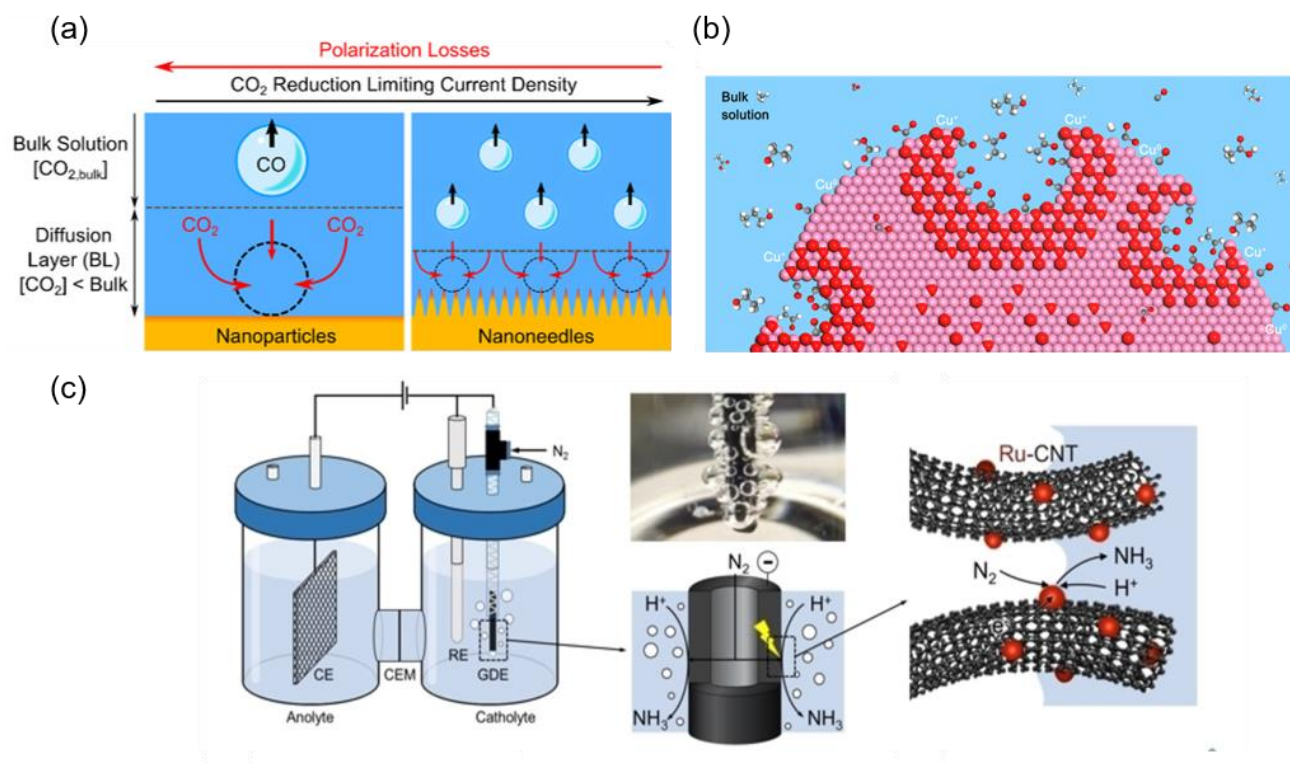


Figure 4. (a) Schematic showing the effect of catalyst nanostructure on bubble departure diameter and its impact on the diffusion boundary layer thickness and CO₂ mass transport. Reproduced from *ACS Sustainable Chemistry & Engineering* **2017**, 5 (5), 4031-4040. Copyright 2017 American Chemical Society. (b) Schematic of a cuprous oxide catalyst with nanocavities that confine carbon intermediates such as CO and C₂H₄. White: hydrogen; grey: carbon; red: oxygen; pink: copper. Reproduced from *Journal of the American Chemical Society* **2020**, 142 (13), 6400-6408. Copyright 2020 American Chemical Society. (c) Schematic illustration (left to right) and picture (middle-top) of the NRR in an H-cell with a microtubular Ru-CNT (carbon nanotube) gas diffusion electrode. Reproduced from *ChemElectroChem* **2020**, 7 (22), 4679-4684. Copyright 2020 European Chemical Societies Publishing.

3.2 Surface functionalization

Functionalization of the electrode or catalyst surface with organic or inorganic ligands has been explored as a strategy to tune the interaction between adsorbed intermediates and catalysts, inhibiting HER and improving product selectivity. In addition to the decoration of the surface of a catalytic material with surface-bound ligands, the covalent grafting of molecular co-catalysts onto the surface of a catalytic material has also been explored as a strategy to further tune the catalyst selectivity.⁵⁵

In this section, we will outline some key examples in the diverse field of catalyst surface functionalization, which has been comprehensively reviewed for the CO₂RR by Reisner and co-workers.²⁵ To date, many organic additives such as amino acids,⁵⁶ amines,^{57, 58} aminothiols,⁵⁹ pyridiniums,^{60, 61} N-heterocyclic carbenes (NHCs),^{62, 63} imidazolium ligands,⁶⁴ porphyrin-based metallic complexes,^{65, 66} polymers,^{67, 68} and inorganic additives,^{69, 70} have been proposed to control the

binding energy of CO₂RR reaction intermediates (**Figure 5a**). For instance, Kim *et al.* demonstrated a 94.2% FE for the production of CO from amine-capped Ag supported on carbon, thanks to the effective suppression of the HER and the intrinsic high selectivity towards the CO₂RR from Ag (**Figure 5b**).⁵⁷ DFT calculations suggested that the amine-capped Ag nanoparticles stabilize the *COOH intermediate while destabilizing *H. Conversely, thiol-capped Ag nanoparticles exhibited superior reaction rates towards both the HER and CO₂ reduction by indiscriminately increasing ΔG^*_{H} and ΔG^*_{COOH} .

As presented in **Figure 5c**, Zhao *et al.* developed a simple modification strategy using amines to depress the hydrogen evolution reaction on ultrasmall Au NPs and enhance CO₂-to-CO conversion.⁵⁸ The amine groups, as well as the molecular configuration, were found to play important roles in tuning the electrocatalytic activity of low-coordinated sites of the nanoparticles. The authors claimed that strong interactions between the Au surface and the amine ligands combined with the peculiar configuration were responsible for the improved CO₂RR performance. Remarkably, linear amines promoted the formation of CO, an effect, which was enhanced by increasing the length of the alkyl chain, whereas the branched polyamine greatly depressed it. Wang *et al.* demonstrated a 55% and 77% selectivity for ethylene and C₂₊ products, respectively, using a tricomponent copolymer to modify the surface of Cu electrodes (**Figure 5d**).⁶⁷ Systematic studies indicated that the three components of the copolymer control electrostatic interactions, gas diffusion, hydrophilicity, which were found to be necessary to improve selectivity. The copolymer was obtained by ring-opening metathesis polymerization, thereby offering a new degree of freedom for tuning the selectivity.

Applying a molecular design approach to tune heterogeneous catalysts has also proved effective in the functionalization of palladium foil with chelating N-heterocyclic carbene (NHC) ligands, demonstrating a 32-fold increase in activity for CO₂ to C₁ products.⁶² N-aryl-pyridinium salts have also proved effective in tuning electronic properties to stabilize intermediates for CO₂RR to ethylene.⁶¹ Porphyrin-based metallic complexes have been used to functionalize copper surfaces to increase the concentration of CO intermediates and promote C-C coupling; a Faradaic efficiency of 41 % for ethanol was achieved at 124 mA/cm² at -0.82 V vs RHE.⁶⁶

Modifying the catalyst surface indirectly has also been implemented by Varela *et al.* through the addition of halides to the electrolyte.⁷⁰ They hypothesized that the adsorption of halides onto copper increased the negative charge of the catalyst surface, altering selectivity. In the case of iodide, the induced negative charge favored the protonation of CO, enhancing CH₄ production.

Applying well-defined molecular approaches to heterogeneous systems can give important insights into catalytic mechanisms and help to fine-tune active sites and product selectivity. Some functionalization strategies operate through molecular coordination and can therefore be carefully controlled by altering functional and side groups so that specific CO₂RR intermediates can be stabilized. Other strategies, such as the addition of halides or ionic liquids, affect the charge on the catalyst surface, increasing CO_{ads} coverage for example.^{70, 71} Both have proven effective in improving product selectivity in the CO₂RR, and similar approaches could be applied to the NRR to help overcome the dominance of HER (Figure 5e). The exact surface binding motifs of ligands and the mechanism for altered selectivity are still unclear. Understanding the precise nature of the interface remains a key challenge for attaining the desired catalytic properties.⁵⁵

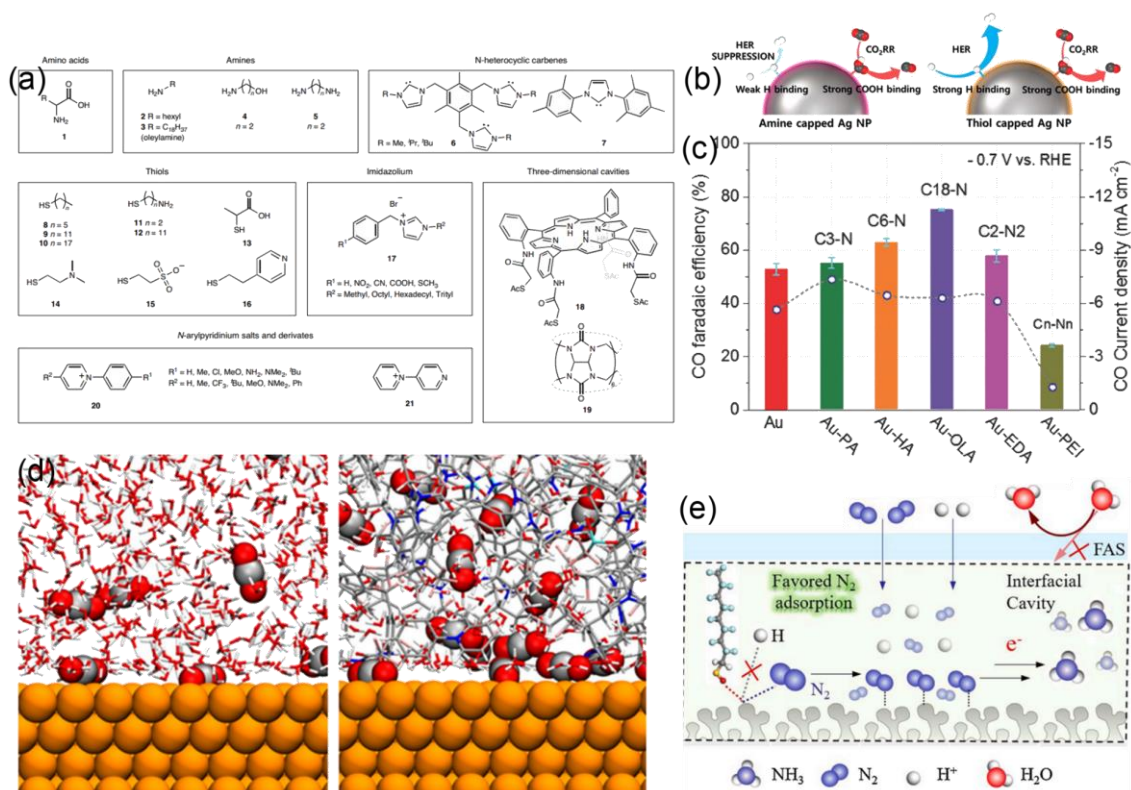


Figure 5. (a) Surface modifiers grouped into different classes used to modulate the local chemical environment around the catalytic site. (amino acids, amines, N-heterocyclic carbenes, thiols, imidazolium, three-dimensional cavities, N-arylpiperidinium salts and derivatives). Reproduced from *Nature Catalysis* **2020**, 3 (10), 775-786. Copyright 2020 Springer Nature. (b) Schematic of the product selectivity, depending on the Ag NPs immobilized with an amine (or thiol)-containing anchoring agent. Reproduced from *ACS Catal.* **2017**, 7 (1), 779-785. Copyright 2017 American Chemical Society. (c) FE_{CO} (column) and j_{CO} (circle) of gold catalysts with different surface amine modifications in CO₂-saturated 0.1 M KHCO₃ at -0.7 V vs. RHE. Reproduced from *Adv. Energy Mater.* **2018**, 8 (25), 1801400. Copyright 2018 Wiley. (d) Interface structure after 12 ns molecular dynamics simulations with a water/Cu interface and random copolymer with a water/Cu interface. Colour code: Cu, orange; C, grey; O, red; N, blue; F, pink; S, cyan; H, white. Reproduced from *J. Am. Chem. Soc.* **2021**, 143 (7), 2857-2865. Copyright 2021 American Chemical Society. (e) Possible NRR mechanism at the surface of the hydrophobic

3.3 Crystal size and facet control

Tremendous advances have recently been made to engineer catalysts in order to lower the HER during the CO₂RR and NRR processes.⁷¹ Compared with their bulk counterparts, nanostructured catalysts show original and often enhanced activity owing to their unique surface electronic and chemical properties. These properties can be finely adjusted to tune the activity and selectivity of electrocatalytic reactions. The surface of a nanomaterial catalyst typically consists of planar areas with single-crystalline orientations separated by steps and kink sites with lower coordination numbers. Complex atomic structures are therefore present at the interface between different grains in polycrystalline and/or nanostructured surfaces. Buonsanti *et al.* investigated the catalytic properties of exposed facets of Cu nanocatalysts at commercially relevant current densities (**Figure 6a**).⁷² The study revealed that facet-dependent selectivity is retained in a gas-fed flow cell, showing greater HER suppression than in a conventional H-cell. The (100) facets of Cu nanocubes have been identified to be selective for the evolution of C₂H₄, whereas the (111) facets of Cu octahedra are selective towards CH₄. Conversely, Cu spheres do not exhibit any specific product selectivity, suggesting that randomly mixed facets cannot depress the HER during the CO₂RR. Chorkendorff *et al.* systematically investigated the structure-selectivity relationship of Au single crystals for electrocatalytic CO₂ reduction (**Figure 6b**).⁷³ Remarkably, they found that the kinetics for the formation of CO strongly depend on the surface structure. Under-coordinated sites, for instance, on the surface of Au(110) or at the step edges of Au(211), show at least 20-fold higher activity than more coordinated configurations – such as Au(100). By selectively poisoning under-coordinated sites with Pb, they identified the selectivity of these active sites towards the reduction of CO₂, effectively suppressing the HER.

Roldan Cuenya, Strasser and coworkers investigated the role of particle size in CO₂ electroreduction using size-controlled Cu nanoparticles (NPs).⁷⁴ A dramatic increase in the catalytic activity and selectivity of CO against H₂ was observed once the particle size was decreased, particularly for NPs smaller than 5 nm, as shown in **Figure 6c**. Changes in the population of low-coordinated surface sites and their stronger chemisorption were linked to H₂ and CO selectivity. As shown in the inset of **Figure 6c**, a drastic increase in undercoordinated atoms is observed below a particle size of 2 nm with a

coordination number lower than 8. These peculiar sites accelerate both hydrogen evolution and CO₂ reduction to CO *via* an increase in binding energy. However, the undercoordinated sites are unfavourable for the subsequent hydrogenation of CO, which lowers the hydrocarbon selectivity of the NPs. A plausible explanation for the observed trend is the reduced mobility of intermediate reaction species (CO and H) on the small NPs due to stronger bonding, which decreases the possibility of further recombination to form hydrocarbons. At intermediate particle sizes, the spherical particle model predicts low and constant populations of (100) and (111) facets, which is consistent with the reduced yet constant hydrocarbon selectivities observed for Cu NPs between 5 and 15 nm compared to Cu bulk surfaces. For these larger NPs, weaker binding of CO and H is expected, favouring hydrocarbon formation.

Another critical parameter for suppressing the HER with metal NP catalysts is the interparticle spacing. Mesoscale phenomena, such as interparticle reactant diffusion and readsorption of intermediates, can play an important role in the product selectivity for multistep reactions.^{75, 76} In this context, Mistry *et al.* showed that for CO₂ electroreduction, decreasing the interparticle spacing for a constant nanoparticle size can suppress the HER, which further increases the selectivity for CH₄ and C₂H₄ owing to the increased possibility of the *CO intermediate re-adsorbing on a neighbouring particle and being further reduced (**Figure 6d** and **6e**).⁷⁷ More importantly, this study uncovers general principles of tailoring NP activity and selectivity by carefully engineering the size and distance. These principles guide the rational design of mesoscopic catalyst architectures to enhance the production of the desired reaction products.⁷⁸

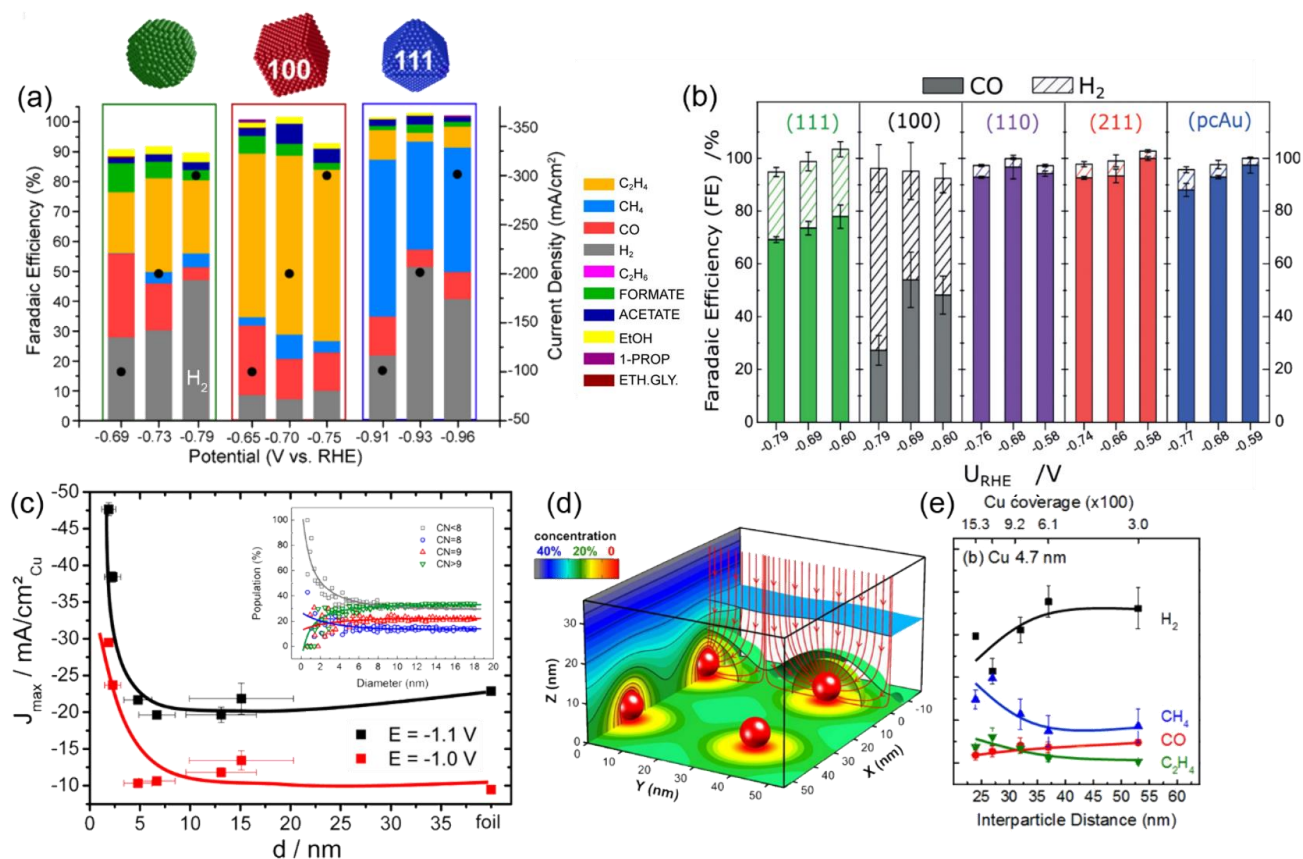


Figure 6. (a) Relation between the Faradaic efficiencies and potentials for different Cu morphologies (sphere, cube and octahedra). Reproduced from *ACS Catal.* **2020**, *10* (9), 4854-4862. Copyright 2020 American Chemical Society. (b) Relation between the Faradaic efficiencies and potentials with the exposure of different Au facets. Reproduced from *Angew. Chem. Int. Ed.* **2019**, *58* (12), 3774-3778. Copyright 2019 Wiley. (c) Particle size effect during catalytic CO₂ electroreduction. The Faradaic current densities at -1.1 and -1.0 V vs. RHE are plotted against the size of the Cu NP catalysts, and the inset shows the population (relative ratio) of surface atoms with a specific coordination number (CN) as a function of particle diameter. Reproduced from *J. Am. Chem. Soc.* **2014**, *136* (19), 6978-6986. Copyright 2014 American Chemical Society. (d) Simulation results of the CO₂ concentration distribution based on diffusion equations. The red arrows show the reactant flux towards the NPs. The colour scale shows the concentration of CO₂ at a given distance from the NPs as a percentage of its value in the bulk of the electrolyte. A diffusion layer thickness of 100 nm was assumed. Reproduced from *ACS Catalysis* **2016**, *6* (2), 1075-1080. Copyright 2016 American Chemical Society. (e) Faradaic selectivity during the electroreduction of CO₂ at -1.1 V vs. RHE with a Cu interparticle distance of 4.7 nm. Reproduced from *ACS Catalysis* **2016**, *6* (2), 1075-1080. Copyright 2016 American Chemical Society.

3.4 Single site engineering

One of the main hurdles to the rational improvement of selectivity using metallic or metal-oxide/sulfide catalysts is the large distribution of accessible sites that may result in different favoured reaction products and decreased selectivities. Single atom catalysts (SACs) hence represent an attractive strategy to increase selectivity *via* a narrower distribution of active sites and an improved control of the first coordination sphere of the active site, bridging the gap between well-defined molecular catalysts and complex heterogeneous materials. The catalytic properties of SACs hence

result from the combination between the molecular tuning of the coordination environment of the active sites and its interaction with the support.⁷⁹ Different types of supports for SACs have been explored to date and include metals, carbon-based materials, and metal (hydr)oxides, nitrides, and carbides. Metal-supported SACs, also called single-atom alloys (SAAs) have also been explored. They generally yield thermodynamically more stable interactions than other atom-support interactions due to strong metal-metal interactions.⁸⁰ Advantageously SAAs can offer different active sites on the host metal (*i.e.* the support) and the individual atoms, providing further opportunities to modulate reaction pathways.⁸¹⁻⁸³ Zhang et al. demonstrated the control of the CO₂RR products between formate and CO by varying the Cu/Sn composition.⁸⁴ They reported that the use of Cu₁Sn₁ comprising a core-shell structure doped with a small amount of Cu using CuSn and SnO alloy as core and shell, respectively, leads to the preferential formation of formate with an FE greater than 95% at -1.2 V. In contrast, single atoms of Sn supported on Cu: Cu₂₀Sn₁ show a high selectivity for CO with a maximum FE_{CO} of 95.3% at -1.0 V.

Carbon substrates have been widely explored in the form of graphite, graphdiyne as well as graphene and its derivatives including heteroatom (N, O, S and P)-doped sp² carbon materials. Carbon supports indeed offer several advantages such as high surface area, high electronic conductivity and strong thermal stability, while they possess numerous coordination environments to stabilize the single atom sites.^{85, 86} The different behaviours of transition metals in the form of nanoclusters or metal-nitrogen-doped carbon catalysts (MNCs) were examined by the Chan and Strasser groups.⁸⁷ The results of their calculations revealed that *CO₂ adsorption is the limiting step on metals, whereas for nitrogen-coordinated SACs, the reaction can be limited either by *CO₂ adsorption or by the formation of *COOH via a proton-electron transfer (**Figure 7a**). Pan and co-authors reported the design of MNC SACs with atomically dispersed Co sites anchored on polymer-derived hollow N-doped porous carbon spheres.⁸⁸ The single-atom Co-N₅ sites were identified as the main active centers for CO₂ activation, and the rapid formation of *COOH as a critical reaction intermediate followed by a rapid desorption of CO. A similar behaviour has also been reported on carbon nanosheet-supported Ni-N₄ sites, which resulted in near-utility selectivity for CO and a single-pass conversion of 2.6% cm⁻² when implemented in a flow cell.⁸⁹ Huan et al. investigated a series of iron-based catalysts synthesized by pyrolysis of Fe-, N-, and C-containing precursors for the electroreduction of CO₂ to CO in aqueous medium and demonstrated that the selectivity of these materials for CO₂ reduction is governed by the proportion of

isolated FeN₄ sites compared to Fe-based nanoparticles.⁹⁰ They demonstrated that the nature of the metal species modulates the selectivity of the reaction pathways and suggested that FeN₄ sites are responsible for CO₂RR whereas the Fe cluster are responsible for HER. In a following work, they demonstrated the strong influence of the electrode support on the catalyst selectivity, highlighting the importance of reducing mass transport limitation to promote a higher selectivity towards CO₂ reduction.⁹¹

In recent years, metal (hydr)oxides, nitrides, carbides and sulfides have become very popular supports of SACs thanks to their high specific surface areas, abundant vacancies, and surface functional groups.⁹² Thanks to their strong corrosion resistance, metal nitrides/carbides with metal centres exposed on their surface are good supports to stabilize isolated metal atoms *via* strong metal-support interactions. In this context, electrically conducting MXenes such as Mo₂C have been explored as supports for the CO₂RR.⁹³ Zhang et al. demonstrated an efficient approach to produce single atom copper immobilized on MXene for the electrosynthesis of methanol from CO₂. The SACs were obtained *via* selective etching of hybrid *A* layers (Al and Cu) in quaternary *MAX* phases (Ti₃(Al_{1-x}Cu_x)C₂).⁹⁴ Combining X-ray absorption spectroscopy analysis and density functional theory calculations, they proposed that the Cu single atoms in the form of Cu^{δ+} with 0 < δ < 2 have a low energy barrier for the rate-determining step corresponding to the conversion of HCOOH* to CHO*, a key reaction intermediate for the reduction of CO₂ to CH₃OH (**Figure 7b**).

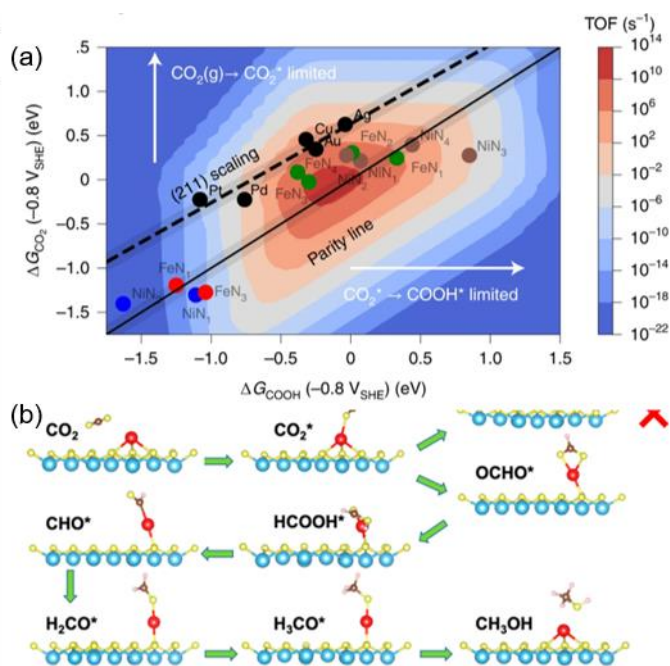


Figure 7. (a) Rate map for CO₂R to CO at -0.8 V_{SHE} and pH=2 obtained from the (211) TM scaling line. The annotated points show MNC SACs either at single or double vacancies. Reproduced from *Nature Catalysis* **2021**, 4 (12), 1024-1031. Copyright 2021 Springer Nature. **(b)** DFT calculations proposed a reaction pathway for the functionalization of CO₂ to methanol on isolated Cu of SA-Cu-MXene. Reproduced from *ACS Nano* **2021**, 15 (3), 4927-4936. Copyright 2021 American Chemical Society.

4. The electrolyte: an active component to drive reactivity and enhance selectivity

4.1 Adjusting the local pH at the electrode/electrolyte interface

The pH value of the electrolyte greatly influences the equilibrium potential of the CO₂RR and NRR, as highlighted in the partial Pourbaix diagrams for the CO₂RR and NRR provided in **Figure 8a** and **Figure 8b**.⁹⁵⁻⁹⁸ A high local pH typically disfavor HER, thus enabling higher faradaic efficiencies for multicarbon products in the context of CO₂RR and for ammonia in the context of NRR.^{99, 100} The groups of Sinton and Sargent have achieved remarkable results for the CO₂RR in highly alkaline media; using 7 M KOH they achieved a 1.3 A cm⁻² partial current density for ethylene in a flow cell.¹⁰¹ Engineering of the triple-phase interface was key to these results and will be discussed further in Section 5. Unfortunately for CO₂ electrolysis, the use of alkaline electrolyte is complicated by the fatal exergonic formation of carbonate ($CO_2 + 2OH^- \rightarrow CO_3^{2-} + H_2O$ / $CO_2 + OH^- \rightarrow HCO_3^-$) which is detrimental to both energy and carbon efficiency.¹⁰² Neutral bicarbonate electrolytes have been applied to reduce electrolyte consumption and to buffer the local pH, although at high currents CO₃²⁻ is still formed from CO₂ and electrogenerated OH⁻. Several studies have explored the dependence of product distribution on local pH at the electrode/electrolyte interface, as well as the concentration and buffer capability of the electrolyte. In that line, a fine tuning of the product selectivity for CO₂RR on Cu electrodes was achieved via the modulation of local pH upon variation of the electrolyte buffer capacity, CO₂ pressure, and current density.¹⁰³ Varela *et al.* proposed that electrolytes with a high buffer capacity could facilitate the transfer of coupled electrons/protons, thus being beneficial for the evolution of hydrogen.¹⁰⁴ By comparison, they found that electrolytes with a low buffer capacity could suppress the formation of H₂ owing to the low concentration of protons near the electrode surface, favouring selectivity towards the formation of C₂H₄ (**Figure 8c**). Conversely, applying a higher current density can also lead to a higher local pH. This is due to a high consumption rate of local protons compared to the rate of mass transport of protons from the bulk electrolyte. Huang *et al.* modelled an electrode surface and found that even in highly acidic electrolytes (pH=1), local neutrality and alkalinity could be created above 200 mA/cm².¹⁰⁵ They required at least 400 mA/cm² to produce multicarbon products.

This improved carbon efficiency considerably, although energy efficiency remains problematic. While a higher CO₂ pressure could result in a lower local pH at a constant electrolyte concentration, they demonstrated that it also favoured ethylene formation by increasing the local *CO concentration and the corresponding *CO surface coverage.¹⁰⁶ Recently, Chen *et al.* reported that adjusting the thickness of a highly porous Au film allows controlling the mass transfer resistance and increasing the local pH at the electrolyte/electrode interface of CO₂ reduction, which results in the promotion of the CO₂RR while inhibiting the HER.¹⁰⁷

For the nitrogen reduction reaction, Xu *et al.* summarized the dependence of the formation of nitrogen-reduction intermediates on pH for aqueous media.¹⁰⁸ Due to the large overpotentials needed to activate N₂ and the low solubility of N₂ in aqueous electrolytes, when the applied overpotential is sufficient to trigger the electrochemical synthesis of NH₃, the reaction at the active sites quickly becomes controlled by the mass transport of N₂ molecules. Consequently, the presence of protons near the electrode surface leads to the undesired production of hydrogen. As illustrated in **Figure 8d**, Wang *et al.* gauged the NRR performance of commercial Pd/C in electrolytes with different pH values. Their observations revealed that the effective suppression of the HER activity in the neutral electrolyte was attributed to a higher barrier for mass and charge transfer.^{109,110}

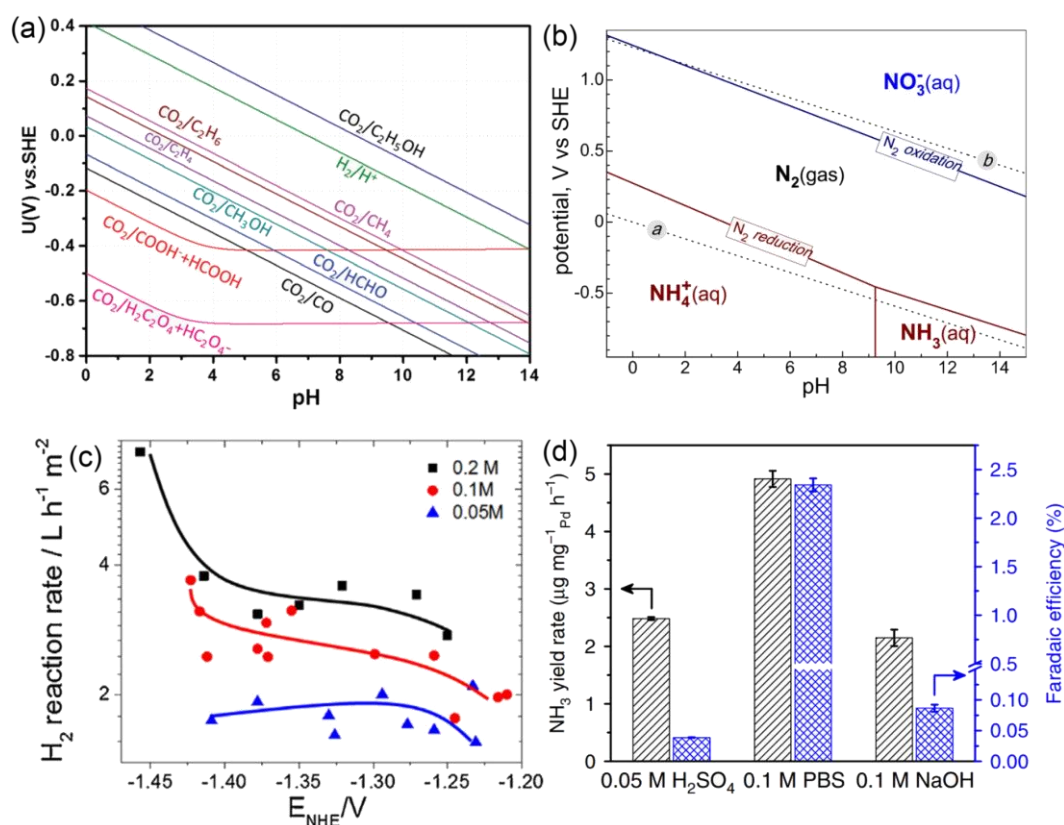


Figure 8. (a) Partial Pourbaix diagram for CO₂ reduction in aqueous solutions that describes the relationship between the equilibrium potential of the associated reaction and pH, which is plotted based on thermodynamic data. Reproduced from *Chem. Soc. Rev.* **2021**, *50*, 4993-5061. Copyright 2021 Royal Society of Chemistry. (b) Partial Pourbaix diagram for the N₂-H₂O system. Solid lines correspond to N₂ reduction to NH₄⁺ or NH₃ (red) and N₂ oxidation to NO₃⁻ (blue). Dotted lines *a* and *b* straddle the region of water reduction to H₂ and oxidation to O₂, respectively. Reproduced from *Science* **2018**, *360* (6391), eaar6611. Copyright 2018 AAAS. (c) Formation rates of gas products as a function of applied electrode potentials in CO₂ saturated electrolytes with different buffer capacities. Reproduced from *Catal. Today* **2016**, *260*, 8-13. Copyright 2016 Elsevier. (d) NH₃ yield rate and Faradaic efficiency of Pd/C processed in N₂-saturated electrolytes with different pH values. Reproduced from *Nat. Commun.* **2018**, *9* (1), 1-7. Copyright 2018 AAAS.

4.2 Optimizing the components of the electrolyte: alkali metal cation effects

Bicarbonate or carbonate are the most investigated electrolyte salts employed for the CO₂RR as they provide a near-neutral pH but most importantly allow to maintain a stable and high dissolved CO₂ concentration upon operation.^{111, 112} Hence, while the nature of the anions are rarely explored in electrochemical studies, a wide range of studies has investigated the variation of the alkali cations. In CO₂RR, while the influence of alkali cations on product selectivity and catalyst efficiency are commonly accepted^{72, 113 114} the origin of this effect is still largely debated in the literature. The influence of the used alkali metal cations on the CO₂RR activity and selectivity is generally attributed to the relatively high concentration of alkali cations in the outer Helmholtz plane (OHP). Early work from Monteiro *et al.* proposed that large cations are specifically adsorbed more easily on the catalyst surface because of the fewer coordinated water molecules.¹¹⁵ Adsorbed cations can also elevate the potential at the OHP and decrease the local proton concentration, suppressing HER.¹¹⁶ Alternatively, it was suggested that the cation size can significantly affect the rate of water hydrolysis by tuning the hydration energy.¹¹⁷ For instance, the pK_a value of Li⁺ was calculated to be three times higher than that of Cs⁺. The hydrated Cs⁺ acts as a buffer, maintaining a locally low pH near the electrode and increasing the local CO₂ concentration compared to Li⁺ by 28 times (**Figure 9a**). To gain more insight into the role of cations in electrocatalysis, Ringe *et al.* developed a combined ab initio/continuum model of cation and electric double layer field effects based on a continuum modified Poisson-Boltzmann approach (**Figure 9b**).¹¹⁸ By applying a single set of cation sizes derived from experimental data, the model showed quantitative agreement with the experiments for the catalyst system on both Ag and Cu. Their theoretical model and experimental results indicate that the repulsive interactions derived from the hydrated cations in the Helmholtz layer should be responsible for the change of surface charge and their electric field. The use of high-valent cations with a small hydration radius also

increases the potential of zero charges or capacitance, which maximizes the surface charge density and the corresponding interfacial electric fields.¹¹⁹ Bell's group provided insights regarding the beneficial effect of cations, particularly at relatively low overpotentials, for which the reaction rate does not perturb the local pH.^{120, 121} Notably, the hydrogen and CH₄ partial currents remained steady, while formate, C₂H₄, and C₂H₅OH formation rates increased when using large alkali cations. The cation size-independent production of H₂ and CH₄ was attributed to the zero dipole moment of *H and *CHO, which are the corresponding reaction intermediates of the reactions (**Figure 9c**).

Alkali metal cations have also been used to promote the CO₂RR in strongly acidic medium. A key advantage to operating at a low pH is the improved carbon utilisation efficiency, which is limited in neutral and alkaline media due to the formation of carbonate. Sargent and co-workers utilised a cation-augmenting layer to sustain a high K⁺ concentration at the copper catalyst surface.¹⁰⁵ They achieved 61 % Faradaic efficiency for CO₂RR products and 40 % for C₂₊ products at 1.2 A/cm², and by lowering the CO₂ flow they reached a single pass conversion efficiency of 77%. Gu et al. explored the effect of alkali cations on the CO₂RR in acid with tin oxide, gold and copper catalysts, achieving 90% faradaic efficiencies for formic acid and CO.¹²² Using a simulation based on the Poisson-Nernst-Planck (PNP) model, they predicted that the origin of such striking effects was the modulation of electric fields, which inhibited the migration of hydronium ions.

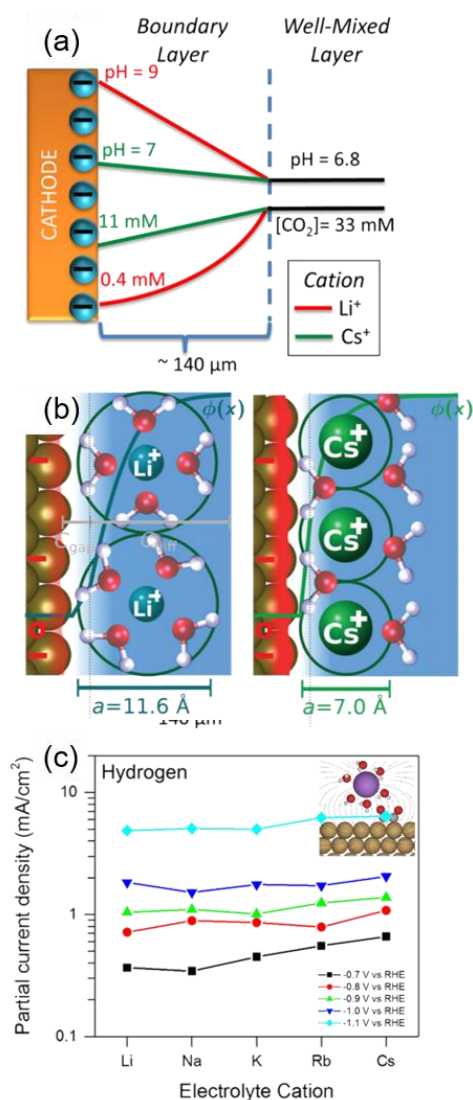


Figure 9. (a) Effect of cation hydrolysis on the electrochemical reduction of CO_2 over Ag. Distribution of pH and CO_2 concentration in the boundary layer. Hydrated Cs^+ buffers the cathode to maintain the pH close to 7 and to increase the CO_2 concentration. Reproduced from *J. Am. Chem. Soc.* **2016**, *138* (39), 13006-13012. Copyright 2016 American Chemical Society. (b) Illustration of the origin of cation effects in field-driven electrocatalysis. Repulsive interactions between hydrated cations at the outer Helmholtz plane reduce the local concentration of cations, the surface charge density (depicted by the red-coloured region) and the electric double layer field. The diffuse layer that is explicitly modelled by the size-modified Poisson-Boltzmann (MPB) model is depicted, as well as the Helmholtz gap capacitance region and the interfacial ion diameter. Reproduced from *Energy Environ. Sci.* **2019**, *12* (10), 3001-3014. Copyright 2019 Royal Society of Chemistry. (c) Average current densities obtained during bulk electrolysis as a function of metal cations at different potentials. Reproduced from *J. Am. Chem. Soc.* **2017**, *139* (32), 11277-11287. Copyright 2017 American Chemical Society.

4.3 The search for novel electrolytes: ionic liquids and non-aqueous electrolytes

Ionic liquids (ILs), which are defined as salts that remain liquid below $100\text{ }^\circ\text{C}$, have been proven to be a promising new class of environmentally benign solvents.¹²³ By tuning the molecular structure and polarity of the IL, the CO_2 and N_2 absorption capacity and the ability to stabilize charged CO_2 and N_2 species can be tuned and optimized. ILs also possess several advantages, such as a wide

electrochemical windows, thermal and chemical stability, negligible volatility and electron transfer mediation for redox catalysis, which make them an interesting alternative to promote the CO₂RR and NRR.¹²⁴ As they are nonaqueous by nature, ILs allow control of the aqueous content to an optimum level to provide protons for hydrocarbon formation while suppressing the HER.¹²⁵⁻¹²⁹

ILs have been extensively investigated for the CO₂RR because the cations of ILs can form a complex with CO₂ and further activates it. Rosen *et al.* reported the use of 1-ethyl-3-methylimidazolium tetrafluoroborate (EMIM-BF₄) as an IL electrolyte for the electrochemical conversion of CO₂ to CO on silver (**Figure 10a**).¹³⁰ The IL system lowers the energy of the *CO₂ intermediate via the formation of a complex intermediate, which lowers the energy associated with the initial step of the reduction reaction.¹³¹ The formation of CO occurred at a very low onset overpotential, and the IL system demonstrated sustained production of CO for 7 hours with a FE_{CO} of more than 96%. ILs have also been applied with transition metal dichalcogenides, which are known to be more prone to promote the HER over other reduction reactions. Remarkably, Asadi *et al.* exfoliated WSe₂ nanoflakes to perform the electroreduction of CO₂ to CO using a 50 vol.% [Emim]BF₄/H₂O solution.¹³² The current density, FE, and TOF in producing CO were all superior at lower overpotentials, suggesting a high selectivity for the CO₂RR (**Figure 10b**). Copper selenide nanocatalysts have been identified to convert CO₂ to CH₃OH at low overpotentials in a [Bmim]PF₆/acetonitrile-H₂O mixed electrolyte.¹³³ In addition, in a [Bmim]BF₄-H₂O electrolyte, MoTe₂ could also be used as a catalyst for CO₂ reduction to CH₄ with a high FE of 83% at a relatively low overpotential.¹³⁴ Atifi *et al.* demonstrated that protic ionic liquids (PILs) derived from 1,8-diazabicyclo[5.4.0]undec-7-ene (DBU) effectively promote the electrochemical reduction of CO₂ to formate (HCOO⁻) with high selectivity (**Figure 10c**).¹³⁵ The use of PILs composed of the conjugate acid of DBU, [DBU-H]⁺, efficiently catalysed the reduction of CO₂ to HCOO⁻ (FE_{HCOOH} ≈ 80%) with significant suppression of CO and H₂ production (FE_{CO} + FE_{H₂} ≈ 20%) in either acetonitrile or an acetonitrile/H₂O mixed electrolyte.

Ionic liquids and nonaqueous electrolytes with high N₂ solubility under ambient conditions can also increase the local concentration of N₂ near the catalyst surface by as much as 20 times compared to water on a volumetric basis.¹³⁶ MacFarlane and co-workers reported the use of ionic liquids with high N₂ solubility for the electroreduction of N₂ to ammonia at room temperature and atmospheric pressure.¹³⁷ As presented in **Figure 10d**, FE_{NH₃} as high as 60% was achieved in [P6,6,6,14][eFAP].

Ortuño *et al.* used DFT calculations to explore the nature of N₂ adsorption on different ions, and found that a stronger interaction accompanied by charge - delocalization will result in stronger adsorption of N₂.¹³⁸ As shown in **Figure 10e**, they found that on a Ru surface the presence of ILs reduces the relative electronic energy of the N₂RR intermediate N₂H* more significantly than that of the HER intermediate, H₂*, lowering the energy by 0.34 eV and 0.11 eV, respectively. Suryanto *et al.* identified the importance of the IL molar fraction (X_{IL}) on the physicochemical properties of the electrolyte mixture and the NRR performance.¹³⁹ An FE as high as 23.8 ± 0.8% with an NH₃ yield rate of 1.58 ± 0.05 × 10⁻¹¹ mol s⁻¹ m⁻² was achieved for X_{IL} = 0.23 at an optimal potential of -0.65 V vs. NHE (**Figure 10f**). Note that in this study, which predates the publication of standard NRR protocols mentioned in the above text, no ¹⁵N labelling studies were provided, but an extensive purification of the N₂ reactant was carried out. The significant drop in the NRR performance when further increasing X_{IL} highlights the role of 1H,1H,5H-octafluoropentyl 1,1,2,2-tetrafluoroethylene ether (FPEE) in facilitating the mass transport of N₂ in the electrolyte. The authors suggested that other factors correlating FE and X_{IL} could play a role, such as the presence of complex molecular interactions and the different diffusion behaviours of neutral N₂ molecules and polar H₂O within the mixed electrolyte system, a known phenomenon with ionic liquids.¹⁴⁰

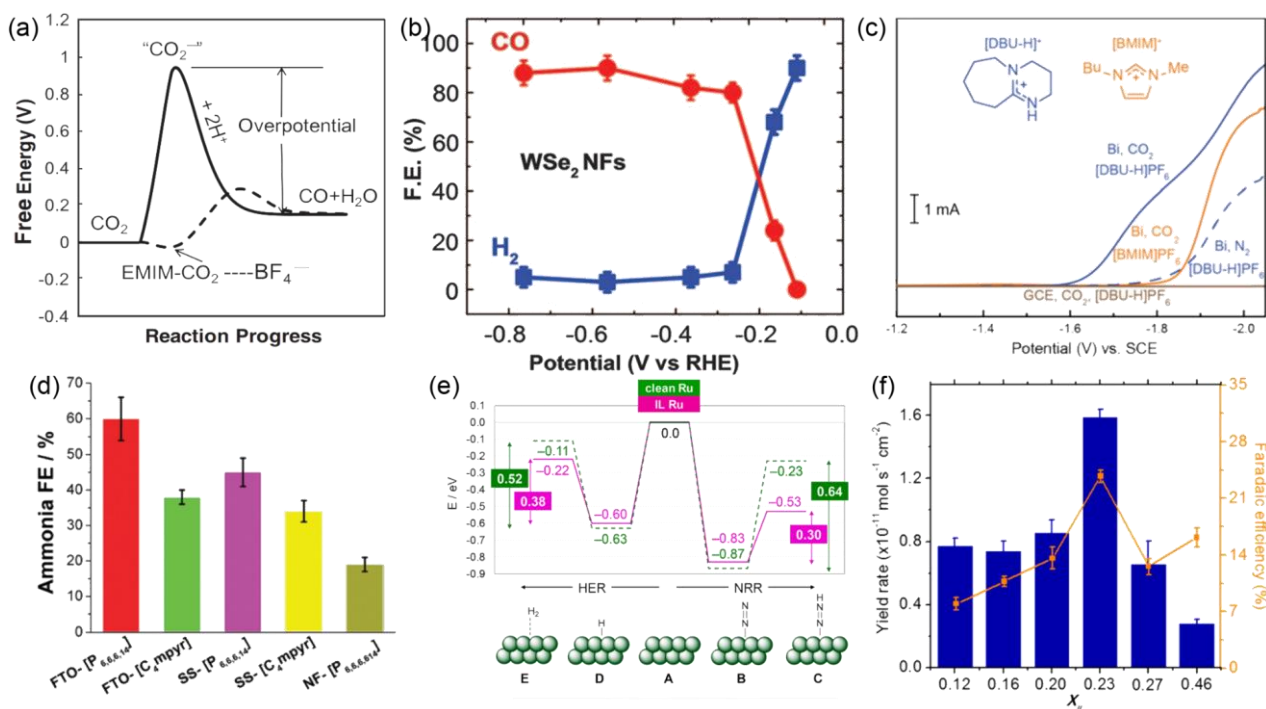


Figure 10. (a) Schematic of how the free energy of the system changes during the CO₂ + 2H⁺ + 2e⁻ ⇌ CO + H₂O reaction in water, acetonitrile (solid line) or EMIM-BF₄ (dashed line). Reproduced from *Science* **2011**, 334 (6056), 643-644. Copyright 2011 AAAS. (b)

Overall FE_{CO} and FE_{H_2} at different applied potentials for WSe_2 NFs. The error bars represent the standard deviation of four measurements. Reproduced from *Science* **2016**, 353 (6298), 467-470. Copyright 2016 AAAS. (c) Linear sweep voltammograms were recorded for Bi-based and bare GCEs in MeCN containing 250 mM IL and 0.1 M TBAPF₆ under the saturation of Ar, N₂, or CO₂. Reproduced from *ACS Catal.* **2018**, 8 (4), 2857-2863. Copyright 2018 American Chemical Society. (d) Faradaic efficiency for electroreduction of N₂-saturated ILs on various electrodes at a constant potential of 0.8 V vs. NHE. Reproduced from *Energy Environ. Sci.* **2017**, 10 (12), 2516-2520. Copyright 2017 Royal Society of Chemistry. (e) Corresponding reaction energy profiles of such intermediates during the NRR (right) and HER (left) for clean (dashed green line) and IL-decorated (solid purple line) Ru surfaces. Reproduced from *J. Phys. Chem. Lett.* **2019**, 10 (3), 513-517. Copyright 2019 American Chemical Society. (f) Solvent-IL ratio (X_{IL}) dependence of the NH₃ yield and FE at -0.65 V vs. NHE. Reproduced from *ACS Energy Lett.* **2018**, 3 (6), 1219-1224. Copyright 2018 American Chemical Society.

4.4 Solid-state electrolyte designs

Conventional liquid electrolytes used in the CO₂RR and NRR, such as KHCO₃, Na₂SO₄, or KOH, mainly have three purposes: *i*) to transport ions between the cathode and anode for efficient current flow, *ii*) to provide protons for successive PCET and *iii*) to solvate liquid products. The mixture of liquid products and ion impurities requires energy- and cost-intensive downstream separation steps to obtain pure products, which complicates the infrastructure for delocalized production.¹⁴¹ To tackle this problem, the concept of solid-state electrolytes was proposed, inspired by progress in solid-state electrolytes for batteries.¹⁴² A solid-state electrolyte is typically placed between ion-exchange membranes with close contact to efficiently transport the generated ions and minimize the ohmic loss of the device.¹⁴³ Remarkably, solid-state electrolytes were found to be very effective in suppressing the HER by limiting the flow of protons to the catalyst active sites during the electrochemical CO₂RR.¹⁴⁴ The Wang group have reported the continuous electrocatalytic conversion of CO₂ to pure liquid fuels using two electrode systems with solid electrolytes.^{145, 146} They applied a porous solid electrolyte (PSE) layer composed of styrene-divinylbenzene copolymer microspheres with sulfonic acid functional groups for proton conduction. Using a formic-acid-selective bismuth catalyst ($FE_{HCOOH} \sim 97\%$), the electrochemically generated protons and formate anions could combine at the PSL to produce formic acid (**Figure 11a**). By directly flowing a carrier gas instead of deionized water through the PSL, they were able to collect product vapours that could be condensed to form the pure product (almost 100 wt.% formic acid), alongside impressive current density and stability (**Figure 11b**).

Sheets *et al.* proposed a novel polymer gel approach to convert N₂ to NH₃ at mild temperatures (30-60 °C) and pressures (20 psig).¹⁴⁷ As illustrated in **Figure 11c**, the polymer gel electrolyte helped to control the rate of the HER by limiting water transport and boosting N₂ transport, thus improving the selectivity towards the NRR.

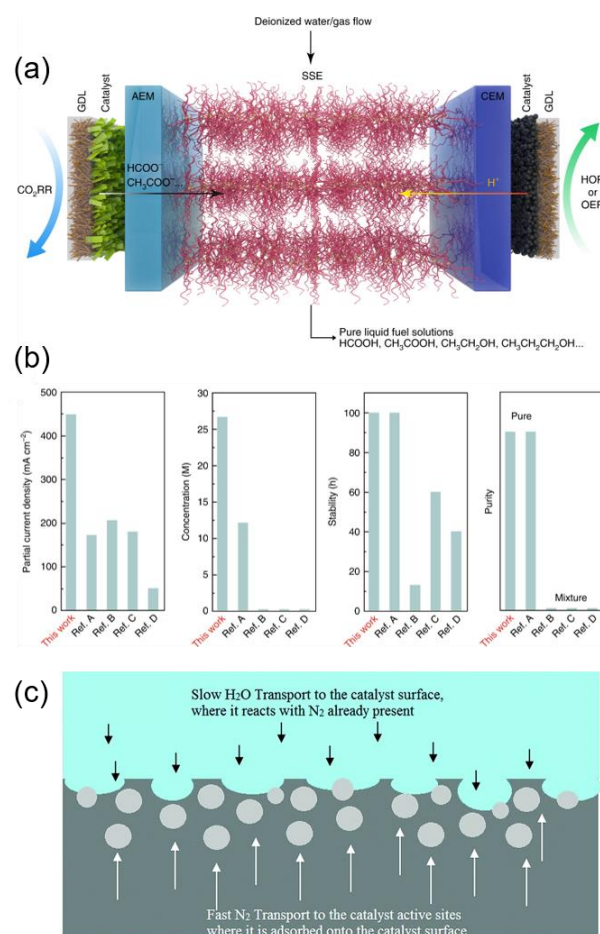


Figure 11. (a) Schematic illustration of the CO₂ reduction cell with a solid electrolyte. Reproduced from *Nat. Energy* **2019**, 4 (9), 776-785. Copyright 2019 Nature Publishing Group. (b) Electrochemical performance of an all-solid-state CO₂RR reactor compared with previous literature. Reproduced from *Nat. Commun.* **2020**, 11 (1), 1-9. Copyright 2020 Springer Nature. (c) Cathode species transport diagram illustrating the advantage of the polymer gel electrolyte to limit water transport. Reproduced from *Chem. Comm.* **2018**, 54 (34), 4250-4253. Copyright 2018 Royal Society of Chemistry.

5. Three-phase interface engineering

The abundance of protons near the catalyst active sites constitutes a significant challenge for the catalyst selectivity vs. the competing HER in aqueous electrolytes, resulting in low selectivity and activity of the CO₂RR and NRR. A mitigation strategy resides in facilitating the accessibility of the catalyst to high concentrations of CO₂ or N₂ molecules. While protons (H⁺) are readily available in aqueous solutions *via* water ionization, the supply of CO₂ and N₂ molecules to the catalyst surface is limited by their low concentration and slow diffusibility. In saturated aqueous electrolytes, the solubility of CO₂ in H₂O is 33 mmol L⁻¹ at 298 K and 1 atm pressure, whereas the value for N₂ in H₂O remains as low as 0.7 mmol L⁻¹.¹⁸ By comparison, the concentration of protons in a neutral aqueous

electrolyte is typically 2.7-fold and 132-fold higher than the concentrations of CO₂ and N₂, respectively.

In the context of CO₂RR, Raciti *et al.* demonstrated that the local concentration of CO₂ at the catalyst surface can reach zero under strong reaction driving force conditions, hence lowering selectivity by limiting the supply of the reactant.¹⁴⁸ Significant advances to minimize this reactant supply issue at the electrode have been made thanks to the implementation of efficient three-phase interfaces between gaseous CO₂, the liquid electrolyte and the solid catalyst. The most typical realisation of such a three-phase interface involve porous gas diffusion layer (GDL) electrodes, which allow the delivery of gas-phase CO₂ directly to the catalyst active sites. Such a strategy, resulting in higher CO₂ and lower H⁺ surface concentrations, has the potential to improve CO₂RR performances while significantly lowering competitive HER. The properties of the GDL can affect CO₂ and water transport heavily, and main advances in this field have been recently reviewed.^{149, 150} and will not be extensively reviewed here in the context of CO₂RR. Thinner GDL/catalyst layers shorten the CO₂ diffusion distance, raising the relative CO₂ concentration; however, excessively high concentrations can decrease multicarbon product formation by competing with intermediates such as CO for binding sites. Tan *et al.* found that by adjusting the catalyst layer structure and the CO₂ feed concentration and flow rate, they could establish a moderate local CO₂ concentration that was optimal for multi-carbon product selectivity.¹⁵¹

Alternatively to requiring GDL-based electrodes, the catalyst support itself can be modulated to modulate the three-phase interface, via a fine-tuning of the local microenvironment near the catalyst surface through nanostructuring and surface functionalisation. Inspired by biological strategies to entrap a gas layer at the surface of a solid, and in particular by plastron effect enabling the diving bell spider to breathe underwater, Wakerley *et al.* functionalized porous dendritic Cu electrodes generated via the DHBT strategy mentioned above in section 3 with long-chain alkanethiols. The resulting superhydrophobic Cu electrodes demonstrated a 6 fold decrease of HER upon treatment with the alkanethiol and a subsequent drastic increase in CO₂ reduction selectivity.²¹ They proposed that the hydrophobicity establishes triple phase interfaces at the electrode where CO₂ mass transport is omnidirectional and H⁺ mass transport is unilateral (**Figure 12a**). This increases the local CO₂ concentration and thereby the surface concentration of Cu-COOH* and Cu-CO*, enhancing C-C coupling. This study led to the identification of the role of hydrophobicity and the formation of gaseous voids as effective levers to orient the reaction pathway towards the formation of multicarbon products. Khan and coworkers explored the idea of gas-trapping further by using a gasphilic silicon substrate in

proximity to the catalyst layer. Creating a CO₂ plastron adjacent to the catalyst improved mass transfer, enriching and maintaining the local CO₂ concentration. Using a smooth copper catalyst they recorded improved activity and a decrease in FE_{H₂} (13 % compared to 29 % with bulk CO₂ bubbling). These trends were replicable using nanostructured copper, demonstrating the transferability of such an approach to different catalysts.¹⁵² Moreover, Xing *et al.* showed that a hydrophobic microenvironment can significantly enhance CO₂ electrolysis by facilitating reactant diffusion (**Figure 12b**).¹⁵³ Using commercial copper nanoparticles dispersed with hydrophobic polytetrafluoroethylene (PTFE) nanoparticles, they reported improved activity and Faradaic efficiency for CO₂ reduction with a partial current density >250 mA cm⁻² and a single-pass conversion of 14% at moderate potentials. Importantly, this performance was approximately twice as large as that of regular electrodes without added PTFE. Similar findings were also observed from a Bi-based catalyst modified with PTFE nanoparticles in the catalyst layer to demonstrate a partial current density of 677 mA cm⁻² for formate and 35% single-pass CO₂ conversion at -0.7 V vs. RHE (**Figure 12c**).¹⁵⁴ Pham *et al.* compared various ionomeric binders on a Cu catalyst, and achieved a 77 % Faradaic efficiency and 600 mA cm⁻² partial current density for C₂₊ products at -0.76 V vs RHE using a fluorinated ethylene propylene (FEP) binder.¹⁵⁵ They attributed these results to the hydrophobic properties of FEP. The Sinton and Sargent groups have also done notable work on modulating the three-phase interface in continuous flow and membrane electrode assembly (MEA) electrolyzers, enabling high current densities (e.g. > 1 A cm⁻²) to be achieved.^{101, 144} For example, they presented a catalyst:ionomer bulk heterojunction (CIBH) architecture, which had both hydrophilic and hydrophobic functionalities. By having different domains that favoured gas and ion transport routes, they were able to decouple gas, ion and electron transport, extending the reaction interface from the submicrometer to the several micrometer range.¹⁰¹ These examples illustrate that moderate hydrophobicity of the catalyst layer can establish a microenvironment with a balance between gaseous CO₂ and liquid electrolytes inside the catalyst layer. Such microenvironments – equivalent to microreactors – reduce the thickness of the diffusion layer, accelerate CO₂ mass transport and link highly active reaction zones at the interfaces between the three phases involved in the reaction.¹⁵⁶ The triple-phase interface can also be further tuned by applying ionomers to control pH and CO₂/H₂O concentrations. Bell and co-workers postulated that anion-exchange ionomers (e.g. sustainion) increase CO₂ solubility, cation-exchange ionomers (e.g. nafion) increase local pH by trapping OH⁻ ions, and both types increase water concentration.¹⁵⁷ By optimising a bilayer ionomer

coating and coupling to pulsed electrolysis, they achieved 90 % Faradaic efficiency for C₂₊ products and just 4 % for H₂.

In the case of NRR, when applying large potentials at the electrodes, the kinetically facile HER becomes preferable to the reduction of N₂ due to the relatively low energy barrier associated with the reaction. It was suggested that the HER should always dominate at normal proton concentrations near the metal electrode surface. However, when few protons or electrons are provided, the NRR may preferentially occur, as recently observed experimentally. Designing a triple-phase interface for NRR can increase the local N₂ concentration and improve *N₂ adsorption, whilst limiting the availability of protons by reducing contact with the electrolyte.¹⁵⁸ Using this strategy, Zhang *et al.* realized triple-phase electrolysis *via in situ* fabrication of Au nanoparticles located on hydrophobic carbon fibre paper (Au/CFP) (**Figure 12d**).¹⁵⁹ The hydrophobic carbon fibres facilitated the formation of three-phase contact points (TPCPs) for N₂, the liquid electrolyte and the Au NPs. Xiao *et al.* successfully modified the *d*-band structure of a self-supporting nanoporous Mo₄P₃ catalyst by capping with a fluorosilane hydrophobic layer.¹⁶⁰ This approach aims at weakening the ability of the material surface to adsorb protons while simultaneously preventing lowering the amount of water available at the active sites, thus further lowering competitive HER. This hydrophobic Mo₄P₃ material exhibits decent NRR performances, with a FE of 10.1% and an NH₃ yield rate of 17.3 μg h⁻¹ cm⁻². According to Wang and co-workers, excessive suppression of the HER is not, however, beneficial to NRR activity, although it can lead to higher Faradic efficiency (**Figure 12e**). A sharp decrease in the local concentration of protons does not benefit the NRR process, as protons are necessary for the successive PCET steps associated with the formation of ammonia. These investigations point out that although the release of hydrogen is a competitive reaction, protons are paradoxically essential to increase the ammonia yield rates.¹⁶¹

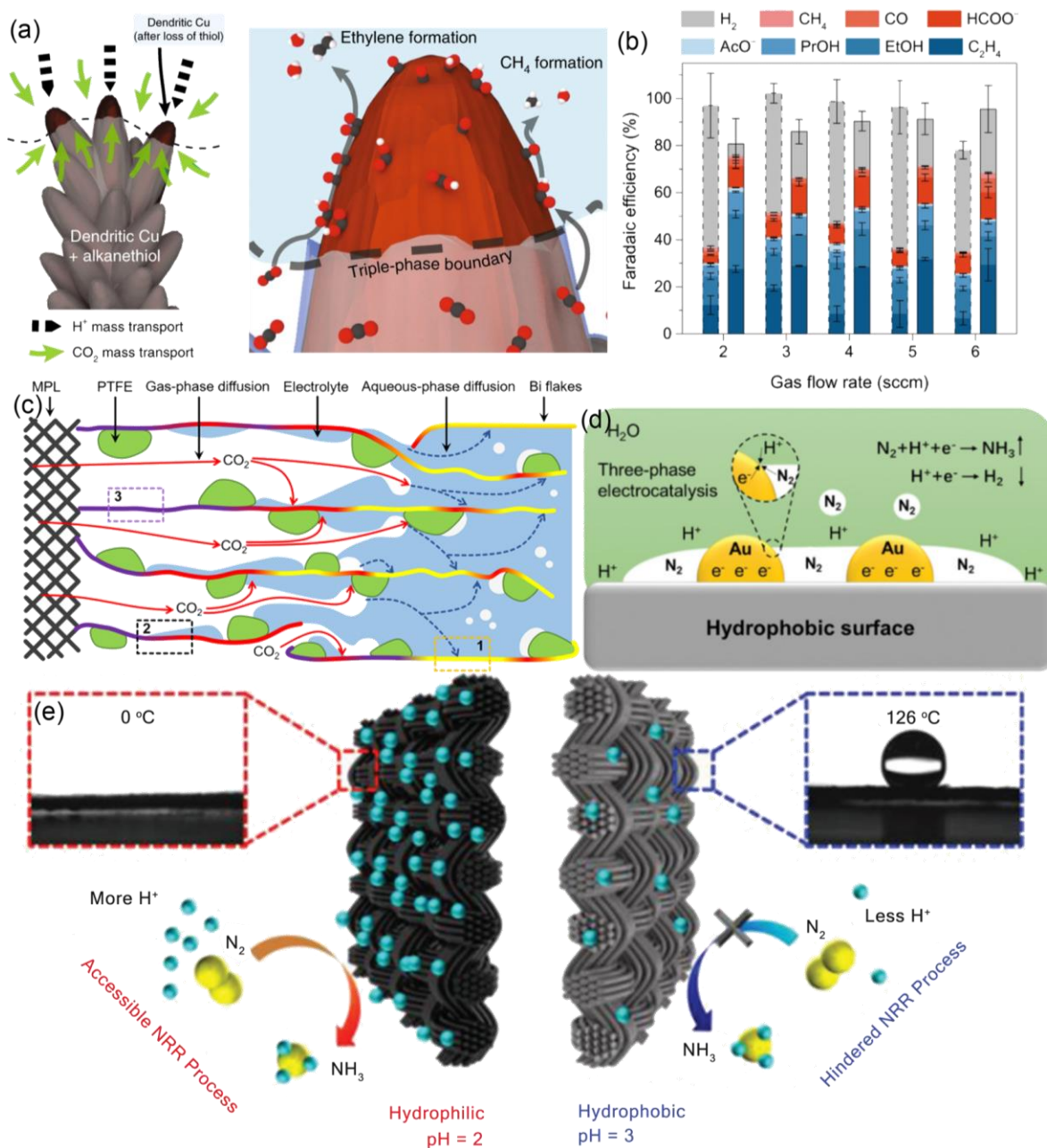


Figure 12. (a) Operation of the hydrophobic dendrite, illustrating the enhanced CO_2 mass transport from the triple-phase boundary between the electrolyte, the electrode and gaseous CO_2 and the resultant formation of key products on the surface. Reproduced from *Nat. Mater.* **2019**, *18* (11), 1222-1227. Copyright 2019 Springer Nature. (b) Faradaic efficiencies for the CO_2 RR on the two electrodes (dash: AvCarb MGL370 + Cu/C; solid: AvCarb GDS2230 + Cu/C) at -1.0 V vs. RHE with various CO_2 flow rates. Reproduced from *Nat. Commun.* **2021**, *12* (1), 1-11. Copyright 2021 Springer Nature. (c) Schematic illustration of CO_2 mass transport inside the catalyst layer with added PTFE, including gas-phase diffusion (solid red arrows) and aqueous-phase diffusion (dashed blue arrows). The dashed rectangles indicate catalyst areas that are only exposed to the electrolyte, exposed to both electrolyte and gaseous CO_2 , and only exposed to gaseous CO_2 . Reproduced from *ACS Energy Lett.* **2021**, *6*, 1694-1702. Copyright 2021 American Chemical Society. (d) Schematic illumination of three-phase contact for N_2 (gas), the electrolyte (liquid), and the catalyst (solid) at the hydrophobic interface. Reproduced from *Adv. Sci.* **2020**, *7* (22), 2002630. Copyright 2020 Wiley. (e) NRR catalytic mechanism of Mo_2C/C under proton-suppressed and proton-enriched conditions. Reproduced from *Adv. Mater.* **2018**, *30* (46), 1803694. Copyright 2018 Wiley.

6. Conclusions and Perspectives

The industrial development of the CO₂RR and NRR is currently plagued by low Faradaic and energy efficiencies. The successive PCET steps associated with the corresponding reaction intermediates increase the complexity and complicate the search for an ideal catalyst. In contrast, the simplicity of the HER mechanism and the abundant presence of protons in traditional electrolytes make the production of hydrogen a competitive and parasitic reaction that consumes a significant amount of electrons to the detriment of the fixation of CO₂ and N₂. Additionally, the selectivity towards a single product, particularly important in the context of CO₂RR is a central point to be considered. Multiple strategies have shown promise but still require the elaboration of a robust and rational framework; they have demonstrated that optimal activity and selectivity can be obtained upon modulating thermodynamics and kinetics of the reaction. By engineering the catalyst, electrolyte and reaction interface, three main strategies have been applied towards that goal (**Table 2**): i) targeting a narrow distribution of molecularly defined active sites, ii) increasing the reactant/proton ratio at the three-phase interface where the reaction takes place to lower the undesired formation of H₂ and iii), the stabilisation and confinement of reaction intermediates in the electrode vicinity to favour the formation of multi-electron reduction products.

The complexity of the parameters involved to address these challenges simultaneously further highlights the interest in combining experimental and theoretical approaches to guide the design of both catalysts and electrolyzers for the CO₂RR and NRR. From this perspective, machine learning will help rapid screening of catalysts with high selectivity based on massive data in the silico database by focusing on near-optimal bond energy with adsorbates, such as *CO and *N₂H. In addition, enabling a better understanding and control of the PCET steps, notably via the elaboration of a robust framework to link the relative contribution of charge transfer and protonation steps on overpotential and on the distribution of surface species will be key to further rationally improve electrocatalysts.

This review illustrated several examples displaying industry-relevant performances in terms of selectivity and current densities, highlighting the potential of electrochemical approaches for the preparation of carbon and nitrogen containing molecules. However, for the CO₂RR, many studies have been performed in alkaline or neutral media, resulting in carbonate formation in the electrolyte. This is detrimental to carbon utilization and energy efficiency, especially considering the energy that would be required to regenerate spent electrolyte. This problem has been considerably underestimated and

overlooked for some time, however, an increasing number of studies over recent years have attempted to tackle this issue. Using acidic electrolyte prevents carbonate crossover to the anode and regenerates CO₂ close to the cathode surface, improving carbon utilization efficiency. Naturally this media poses challenges regarding hydrogen evolution, and the application of strategies covered in this review will be pivotal in overcoming this.

Moreover, most of the presented strategies introduced in the present review enable improving catalyst selectivity for a relatively short period of time, but have not been investigated over industrially relevant time scales. Maintaining high selectivity for the CO₂RR and NRR over long operation time remains the largest challenge to date, as rapid loss in activity and selectivity is observed for most of the systems reported. This notably results from the fact that in operation undesirable intermediates or poisonous byproducts preferably deposit on the catalyst surface and affect the catalysis process. This phenomenon may decrease the effective area of the electrocatalyst, accelerate cathodic degradation and increase selectivity towards competitive HER. The demonstration of catalysts with ultralong stability of > 5000 hours constitutes in our view the last milestone to be reached in order to validate the industrial potential of the CO₂RR and NRR.

Table 2. Reference numbers of key examples of the three strategies for enhanced product selectivity in carbon dioxide and nitrogen reduction reactions and how they are implemented. References related to CO₂RR are in normal text while references related to NRR are underlined.

	Strategy	i) Molecularly Defined Active Sites	ii) High Local Reactant Concentration	iii) Stabilizing and Confining Intermediates
Catalyst Design	Porous Networks		30	32, 40, 45
	Nanostructures (e.g. wires, films, needles etc.)		50, 51, <u>54</u>	52
	Surface functionalization	61, 62, 66		57, 58, 61, 66, 70
	Control of crystal size, facet & spacing	72-74, 77		
	Single-site engineering	81, 86-91		81
Electrolyte Engineering	Adjusting local pH		101, 103-105, 107	106, <u>108</u>
	Alkali metal cation effects		105, 122	120
	Ionic liquids		<u>137, 139</u>	130, 131, <u>138</u>
	Solid-state electrolyte		143-145, <u>147</u>	
Three-Phase Interface Engineering	Gas diffusion layer electrodes		150-152	
	Gas trapping		21, 152-154, <u>159, 160</u>	
	Utilizing ionomers		101, 155, 157	

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