

REVIEW

[View Article Online](#)
[View Journal](#) | [View Issue](#)Cite this: *J. Mater. Chem. A*, 2024, **12**, 10597Advances in the direct electro-conversion of captured CO₂ into valuable productsKezia Langie,^{†ab} Gwangsu Bak,^{†c} Ung Lee,^{ade} Dong Ki Lee,^{aef} Chan Woo Lee,^{ib} Yun Jeong Hwang^{ib*cg} and Da Hye Won^{ib*adh}

The direct electrochemical conversion of captured CO₂ (capt-eCO₂R) into valuable chemicals has recently emerged as a promising carbon capture and utilisation technology that will contribute to achieving net-zero carbon emissions. Conventional electrochemical CO₂ (eCO₂R) typically uses pure CO₂ gas as a reactant; thus, this system requires substantial energy and capital allocation across the entire process, from the initial CO₂ capture to the post-CO₂ conditioning for product separation. The capt-eCO₂R addresses these limitations and presents a compelling economic advantage by integrating the CO₂ capture and direct electro-conversion of captured CO₂ in the form of carbamate and (bi)carbonate without a CO₂ conditioning process. The capt-eCO₂R is still in the early stages of development and is not as mature as the conventional eCO₂R; thus, several challenges remain to be addressed to improve system performance. This review provides a comprehensive overview of the capt-eCO₂R system, including various system configurations, suitable catalysts, and strategies to enhance performance within captured media. The reaction mechanisms depend on the form of captured CO₂; therefore, we categorised them according to the type of CO₂ absorbent. The outlook, ongoing challenges, and strategies for future development are also presented.

Received 21st February 2024
Accepted 2nd April 2024

DOI: 10.1039/d4ta01178c

rsc.li/materials-a

1. Introduction

The concentration of CO₂ in the atmosphere has increased over hundreds of years owing to anthropological activities, reaching 413.2 ppm in 2020, which represents an increase higher than the average growth rate observed over the past decade (2.5 ppm per year).¹ Carbon capture, utilisation, and storage (CCUS) technology has attracted considerable interest as a means to overcome the climate crisis caused by CO₂.^{2–4} Electrochemical CO₂ reduction (eCO₂R) to produce value-added chemicals is one of the most significant CCUS technologies owing to its high potential to achieve net-zero carbon emission through

integration with renewable energy sources. Although eCO₂R has a relatively low level of technological readiness, recent developments have markedly improved its practical feasibility.

Among the various components of the eCO₂R system, a state-of-the-art electrolyser or reactor that directly supplies gaseous CO₂ to the catalyst layer through a gas diffusion electrode (GDE) considerably improves the scale and selectivity of the reaction (Fig. 1a), leading to record-breaking current densities ranging from hundreds of mA cm^{−2} to several A cm^{−2}.^{5,6} In addition, the conventional eCO₂R achieves a high Faradaic efficiency (FE) of more than 90% and 80% in the case of C₁ (*i.e.* CO and formate)^{7,8} and C₂₊ (*i.e.* ethylene, ethanol acetate, *n*-propanol, *etc.*)^{9,10} compounds, respectively. Nevertheless, techno-economics analysis (TEA) indicates that the economic viability of the eCO₂R system is limited considering the current levels of technology and the market price of the target chemicals.^{11,12} The energy-intensive upstream utilisation of CO₂ as a reactant gas is one of the factors undermining the technological competitiveness of eCO₂R.¹³ Expensive, energy-intensive processes such as CO₂ absorption, desorption, and CO₂ compression are required to supply high-purity CO₂ from flue gas or even direct air capture (DAC). For instance, CO₂ capture using a commercially available CO₂ absorbent, such as monoethanolamine, requires up to 4.3 GJ t_{CO₂}^{−1} for amine regeneration.^{14,15} This drawback compromises the environmental sustainability of the technology. To implement a profitable and environmentally

^aClean Energy Research Center, Korea Institute of Science and Technology, Seoul 02792, Republic of Korea. E-mail: dahye0803@kist.re.kr^bDepartment of Chemistry, Kookmin University, Seoul 02707, Republic of Korea^cDepartment of Chemistry, Seoul National University, Seoul 08826, Republic of Korea. E-mail: yjhwang1@snu.ac.kr^dDivision of Energy and Environmental Technology, KIST School, Korea University of Science and Technology (UST), Seoul 02792, Republic of Korea^eGraduate School of Energy and Environment (Green School), Korea University, Seoul 02841, Republic of Korea^fDepartment of Chemical and Biomolecular Engineering, Yonsei-KIST Convergence Research Institute, Yonsei University, Seoul 03722, Republic of Korea^gCenter for Nanoparticle Research, Institute for Basic Science (IBS), Seoul 08826, Republic of Korea^hKHU-KIST Department of Converging Science and Technology, Kyung Hee University, Seoul 02477, Republic of Korea[†] These authors contributed equally to this work.

sustainable eCO₂R system, an advanced system linking low-concentration CO₂ capture and conversion is critical.

The direct electrochemical conversion of captured CO₂ (capt-eCO₂R) has recently been recognised as a state-of-the-art eCO₂R system that simplifies the system configuration (Fig. 1b). Rather than relying on gaseous CO₂ as a reactant, this system directly utilises captured CO₂ in CO₂ absorbent and thus integrates the CO₂ capture and electrochemical conversion processes. Unreacted CO₂ gas is absorbed in the reaction media (*i.e.*, CO₂ absorbent), allowing concentrated gas products to be obtained from the capt-eCO₂R. In contrast, the conventional gas-fed eCO₂R system (gas-eCO₂R) necessitates an additional process to separate the gas products from the unreacted CO₂ gas. Thus, in principle, systems that directly convert captured CO₂ conversion do not require energy-intensive processes such as CO₂ conditioning (CO₂ desorption and compression) and product separation, which are essential in gas-eCO₂R (Fig. 1c).¹⁶ Based on these intrinsic advantages, a few studies have compared the life cycle analysis (LCA) and TEA of capt-eCO₂R systems with those of gas-eCO₂R systems. Langie *et al.* compared the LCA and TEA of the direct electrochemical conversion of triethylamine (TREA) captured CO₂ with those of several syngas production CCU systems such as the reverse water gas shift reaction and conventional gas-eCO₂R.¹⁷ The capt-eCO₂R system using TREA recorded the most favourable operating expenditure (OPEX) and lowest break-even syngas price. Moreover, evaluation of the global warming potential (GWP) through global sensitivity analysis showed that the capt-eCO₂R system using TREA can achieve near net-zero CO₂ emissions, making the capt-eCO₂R system industrially feasible, environmentally sustainable, and economically viable. Despite these advantages, the performance of capt-eCO₂R is rather low;

however, LCA/TEA analyses suggest that improving the current density at lower overpotentials may make the system more compelling. A similar direction of development was proposed by Gao *et al.*, who found that lower capital expenses could be achieved with capt-eCO₂R, but a carbon-negative system still requires high current densities of over 0.15 A cm⁻².¹⁸ Therefore, to implement a profitable and environmentally sustainable system, improving technological maturity through technical development and a greater theoretical understanding of the capt-eCO₂R is urgently required.

In this study, we investigated various capt-eCO₂R systems that employ different types of electrolyzers and absorbents. The solvent in such systems acts as both a CO₂ absorbent and an electrolyte, thus determining the form of the captured CO₂ (*e.g.*, carbamate, bicarbonate/carbonate, *etc.*). Furthermore, because the CO₂ reactant is captured in a liquid solvent, system components such as electrodes, membranes, electrolytes, electrolyzers, and other parameters should differ from those of the conventional gas-eCO₂R system. Therefore, this review has a particular focus on the role of the system elements, the reaction mechanism of the capt-eCO₂R, and strategies to improve performance based on the form of the captured CO₂ and the type of solvent. A perspective on an advanced future capt-eCO₂R system with industrial applications is also presented.

2. Direct conversion of captured CO₂

The capt-eCO₂R system achieves the electrochemical conversion of captured CO₂ in a liquid electrolyte without additional feeding of gaseous CO₂ (Fig. 1b). In this system, carbamates or bicarbonate/carbonates, the dominant forms of captured CO₂

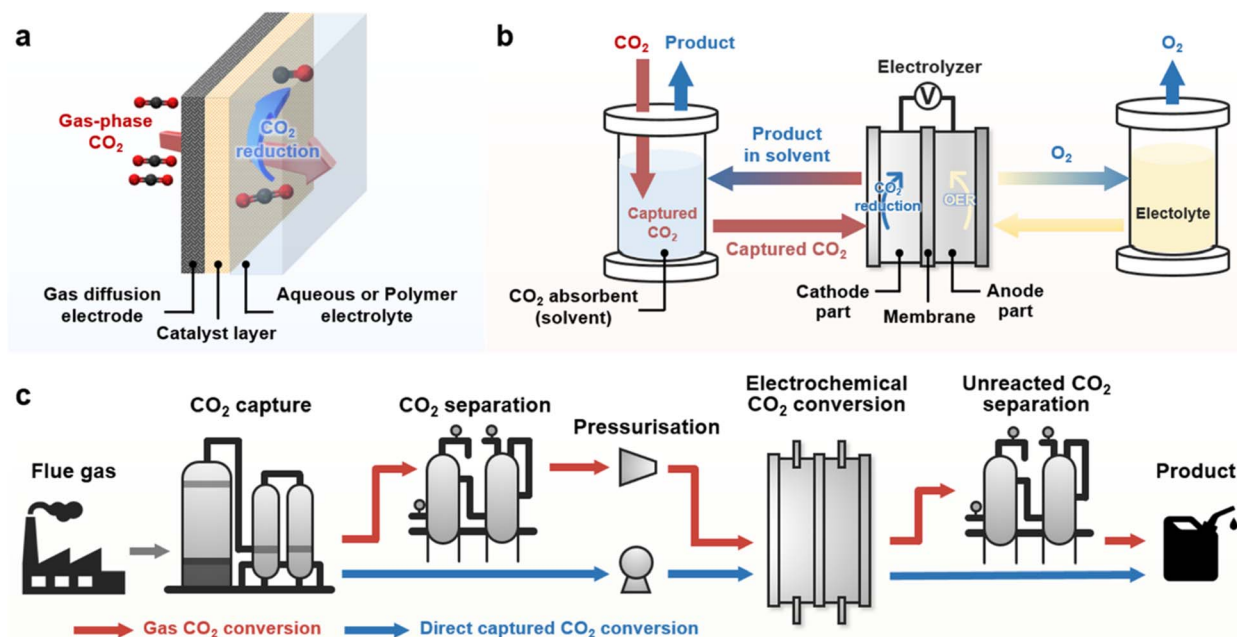


Fig. 1 (a) Schematic of the GDE electrolyser for eCO₂R. (b) System for capt-eCO₂R. (c) Comparison of the systematic pathway of conventional gas-eCO₂ (red line) and capt-eCO₂R (blue line).



Table 1 CO₂ capture agents and its capture ability and capture form of CO₂

CO ₂ capture agent		Capture ability	Concentration	Capture condition	Capture form	Reference
Metal hydroxide	KOH	~0.745 mol _{CO₂} mol _{KOH} ⁻¹	0.412 mol L ⁻¹	35 °C, 4 bar ^a	Bicarbonate	19
Primary amine	Monoethanolamine	0.58 mol _{CO₂} mol _{amine} ⁻¹	30 wt%	40 °C, 1 bar ^b	Carbamate	20
Secondary amine	Diethanolamine	0.53 mol _{CO₂} mol _{amine} ⁻¹	30 wt%	40 °C, 1 bar ^b	Carbamate	20
Tertiary amine	Triethanolamine	0.39 mol _{CO₂} mol _{amine} ⁻¹	30 wt%	40 °C, 1 bar ^b	Bicarbonate	20
	Triethylamine	~0.193 mol _{CO₂} mol _{amine} ⁻¹	3 M	R. T. ambient ^c		17

^a The capture ability is evaluated by feeding 99% CO₂. ^b The capture ability is evaluated by feeding 15% CO₂ and 85% N₂ concentration. ^c The capture ability is evaluated by feeding 5% CO₂ concentration in flue gas.

in the aqueous absorbent, are utilised directly as reactants. The formation of carbamate or bicarbonate is determined by the type of absorbent used. The capture abilities and forms of major CO₂ absorbents are outlined in Table 1. Primary and secondary amines capture one CO₂ molecule per two amine molecules, primarily as a carbamate. Theoretically, tertiary amines can capture one CO₂ molecule per one amine molecule, but the reported low CO₂ capture ability appears to be due to a slow CO₂ absorption rate. Potassium hydroxide (KOH) captures one CO₂ molecule per one molecule, but salt formation hinders the recording of the theoretical CO₂ capture ability. Considering that the optimal CO₂ absorbent concentrations for primary/secondary amines and tertiary/KOH are 30 wt% and 3 M, respectively, the concentrations of carbamate and bicarbonate/carbonate would be similar regardless of the types of CO₂ absorbents used. Therefore, in this review, we categorised the previous studies based on the forms of the captured CO₂ (Fig. 2). As carbamate and (bi)carbonate electrolytes have different reaction mechanisms, their utilisation requires different strategies and approaches.

2.1 CO₂ captured in carbamate form

Many primary and secondary amines, such as monoethanolamine, diethanolamine (DEA), 2-amino-2-methyl-1-propanol (AMP), and their mixtures, are commercially available CO₂ absorbent, primarily capturing CO₂ as carbamates (RNHCOO⁻). Therefore, CO₂ captured by these types of amines can be easily integrated into established industrial processes; however, the

utilisation of carbamate can prove challenging owing to the very strong C–N bond, which necessitates the use of an energy-intensive amine regeneration process.^{21,22} Furthermore, the negatively charged moiety of the carbamate results in electrostatic repulsion between the cathode and carbamate. Several approaches have been developed to overcome the limitations imposed by the intrinsic properties of carbamate.^{2,23,24}

2.1.1 Monoethanolamine media. Monoethanolamine is the most common adsorbent used in the direct conversion of captured CO₂. Chen *et al.* first achieved capt-eCO₂R in a monoethanolamine medium by experimentally screening several metal candidates, including In, Sn, Bi, Pb, Pd, Ag, Cu, and Zn to identify high-performing catalysts in 30% (w/w) monoethanolamine solution.²⁵ In this experiment, the monoethanolamine solution was saturated with CO₂ for 30 min before the experiment, and capt-eCO₂R was conducted in a gas-tight H-type cell under an Ar atmosphere (Fig. 3a). The main products of these catalysts primarily yielded the same products as conventional gas-eCO₂R, *i.e.*, CO and formic acid; however, the hydrogen evolution reaction (HER), which competes with the eCO₂R, occurred significantly under these operating conditions, thereby reducing the efficiency of the system. Several classes of organic surfactants were added to the monoethanolamine solution to modulate the water access near the electrode and reduce the HER activity, including the cationic cetyltrimethylammonium bromide (CTAB), anionic sodium dodecyl sulfate (SDS), and the non-ionic Triton X-100 (Fig. 3b and c). CTAB outperformed the other surfactants in increasing the

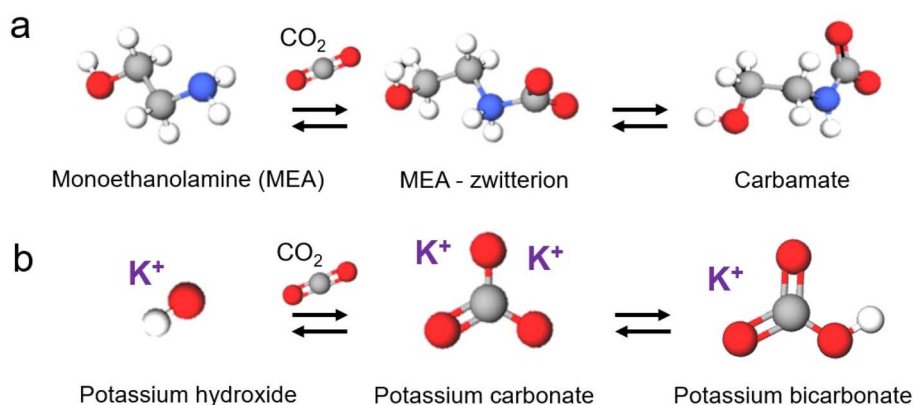


Fig. 2 Reaction mechanism of the direct conversion of CO₂ into (a) carbamate and (b) bicarbonate.



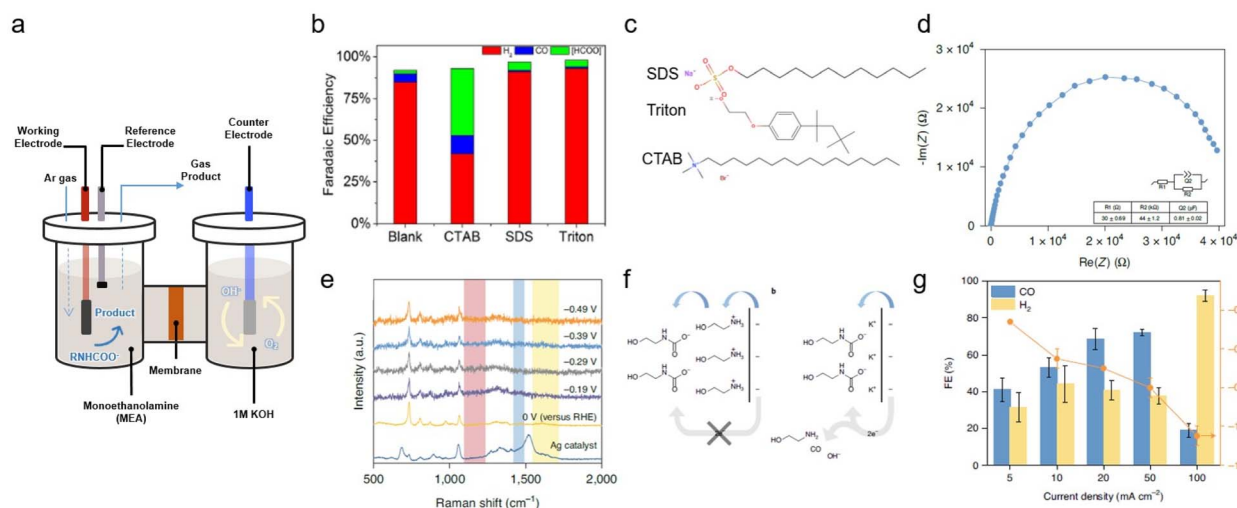


Fig. 3 (a) Schematic of an H-type cell. (b) FE of products obtained with a smooth indium plate at a potential of -0.8 V vs. RHE in monoethanolamine containing 0.1% (w/w) of various surfactants. (c) Structures of the examined surfactants. Copyright 2017,²⁵ Wiley. (d) EIS of the monoethanolamine/KCl electrolyte collected at open circuit potential, approximately -0.145 V vs. Ag/AgCl. Inset: the equivalent circuit and fit values of each component. R_1 and R_2 are the series and charge transfer resistances, respectively; Q_2 is the constant phase element. The error of each component denotes the standard deviation of three independent measurements. (e) *In situ* surface-enhanced Raman spectra of the monoethanolamine/KCl electrolyte. (f) Proposed interfacial structure near the electrode surface. (g) Product distribution of monoethanolamine- CO_2 conversion to H_2 and CO at different applied current densities, ranging from 5 mA cm^{-2} to 100 mA cm^{-2} in a flow cell system. Copyright 2020,²⁴ Springer Nature.

capt-e CO_2R selectivity over the HER (Table 2). For instance, the In electrode achieved enhanced the FE of CO from 4.8 to 17% and that of formate from 2.4 to 45.4%.

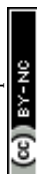
A similar monoethanolamine utilisation system was developed by Lee *et al.*; however, this system also exhibited low capt-e CO_2R performance.²⁴ In CO_2 -saturated monoethanolamine solution, a Ag catalyst showed a CO FE of just 5% in the potential range from -0.46 to -0.78 V. Electrochemical impedance spectroscopy (EIS) and *in situ* Raman analysis of capt-e CO_2R in monoethanolamine solution showed the formation of an electrochemical double layer (EDL) composed of ethanol-ammonium ions and carbamate. Due to the cationic ethanol-ammonium ion forming an inner Helmholtz layer owing to the negatively biased electrode surface, electron transfer must first pass through the ethanol-ammonium ion before reaching carbamate, which consequently inhibits the facile conversion of carbamate (Fig. 3f). Various alkali cations, such as Li^+ , Na^+ , K^+ , Rb^+ , and Cs^+ , were added to the electrolyte to systematically tune the EDL. Among the various alkali cations, K^+ significantly improved the capt-e CO_2R performance,

by facilitating the adjustment of EDL, as evidenced by data from EIS and *in situ* Raman analysis (Fig. 3d and e). The addition of Li^+ and Na^+ resulted in the production of less CO than the pure monoethanolamine solution. This trend correlates with the ionic radius of the cations owing to differences in water solvation. Increasing the reaction temperature to facilitate cleavage of the C–N bond can also improve the performance of the system. The current density at 60°C was 15 times higher than that at room temperature. By adjusting the EDL with a high concentration of alkali salt, 2 M KCl, and circulating the catholyte at 60°C in the flow cell, CO production remarkably increased to 72% at 50 mA cm^{-2} and -0.8 V *versus* reversible hydrogen electrode (RHE) by directly utilising monoethanolamine captured CO_2 (Fig. 3g).

To understand the capt-e CO_2R system, its catalytic activity was investigated in various absorbent media. Atomically dispersed nickel-doped carbon (Ni–N/C), a promising capt-e CO_2R catalyst, was applied in CO_2 -captured 5 M monoethanolamine solutions (Fig. 4a and b).²⁶ The Ni–N/C catalyst has demonstrated high CO production selectivity comparable to

Table 2 FE of various products during capt-e CO_2R on various metal electrodes in CO_2 -saturated 30% (w/w) monoethanolamine solution with 0.1% (w/w) CTAB²⁵

Catalyst		In			Sn			Bi			Pb			Pd			Ag			Cu			Zn		
F.E. (%)	$-V_{\text{RHE}}$	0.8	1.1	1.3	0.8	1.1	1.3	0.8	1.1	1.3	0.8	1.1	1.3	0.1	0.5	0.8	0.8	1.1	1.3	0.8	1.1	1.3	0.8	1.1	1.3
H_2		41.9	42.0	44.3	68.6	78.5	93.4	69.5	87.1	93.4	79.6	79.2	85.1	91.6	87.3	96.0	62.8	84.7	89.5	79.7	98.1	91.0	103.0	91.4	102.0
CO		17.0	10.7	11.2	9.0	3.6	2.6	7.0	4.9	2.6	1.9	3.0	3.5	—	—	—	33.4	15.9	9.2	1.7	—	—	3.7	2.9	0.5
HCOOH		45.4	39.4	36.5	19.0	16.4	2.0	24.3	7.1	3.9	8.5	8.7	6.1	4.0	4.1	0.1	2.0	2.8	1.7	19.1	0.5	0.1	5.4	2.0	2.0
Total		104.3	92.1	92.0	96.6	98.5	98.0	100.8	99.1	99.9	90.0	90.9	94.7	95.6	91.4	96.1	100.8	103.4	100.4	100.5	98.6	91.1	112.1	96.3	104.5



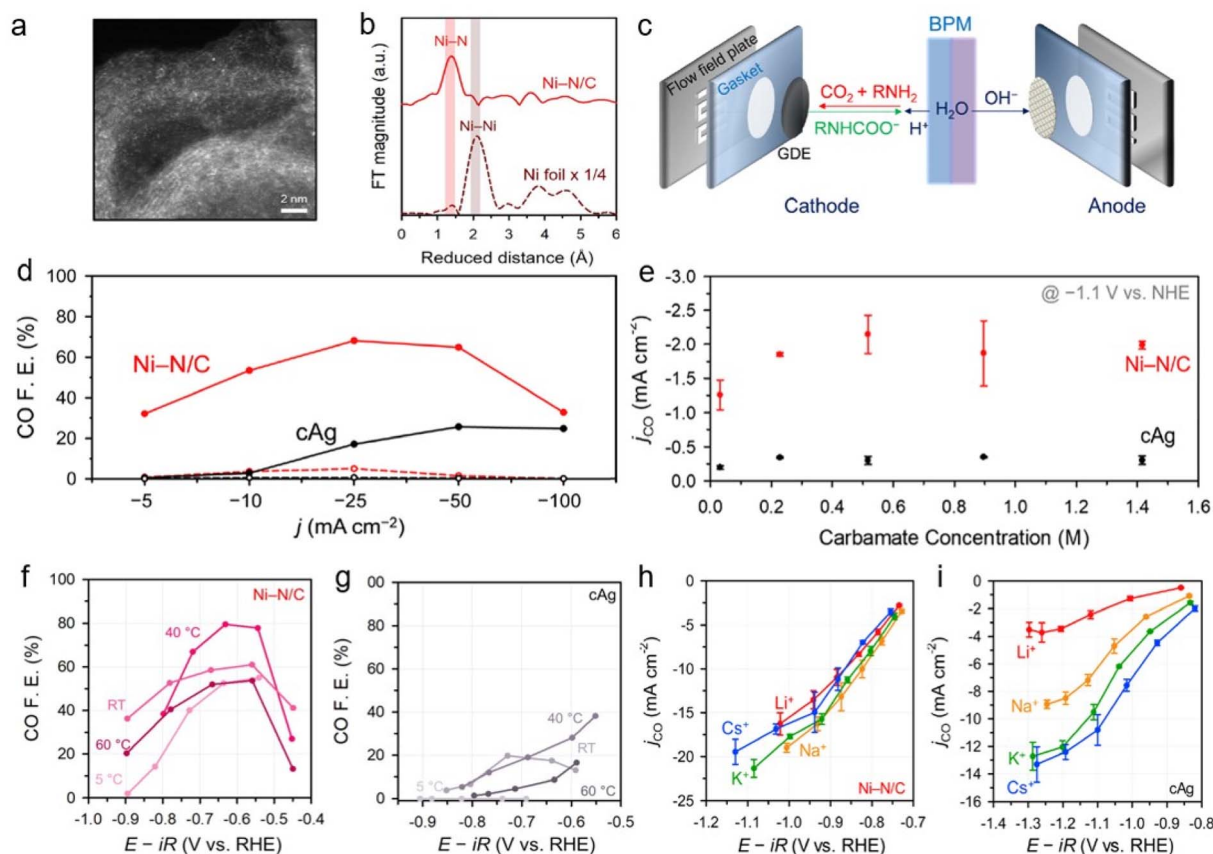
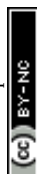


Fig. 4 Physical characterisation of Ni-N/C (a) high-angle annular dark-field scanning transmission electron microscopy image. (b) Extended X-ray absorption fine structure spectra of Ni-N/C and Ni-foil. (c) Schematic of capt-eCO₂R in MEA cell with BPM, anode and cathode. (d) Ni-N/C and cAg performance in membrane electrode assembly using 5 M monoethanolamine. (e) j_{CO} of Ni-N/C and cAg with respect to carbamate concentration. CO FE of (f) Ni-N/C and cAg (g) at various temperatures. Cation effect and sensitivity of Ni-N/C (h) and cAg (i) by occupying conventional gas-eCO₂R. Copyright 2020,²⁶ Royal Society of Chemistry.

that of a Ag catalyst, while the atomically dispersed active sites efficiently suppress the HER. Using Ni-N/C in an H-type electrolyser using CO₂-captured 5 M monoethanolamine solutions, they achieved a CO partial current density three times higher than that obtained with a commercial Ag electrode (cAg). Furthermore, the introduction of a zero-gap membrane electrode assembly (MEA) equipped with a bipolar membrane (BPM) (Fig. 4c) improved the system efficiency, with Ni-N/C resulting in a 64.9% CO FE at -50 mA cm^{-2} , 2.5 times higher than that obtained using the cAg (Fig. 4d). To understand the catalytic performance of the Ni-N/C catalyst, the reaction mechanisms were investigated by adjusting the carbamate concentration (from 1 M to 5 M monoethanolamine) (Fig. 4e). Interestingly, the reaction rate was unaffected by the carbamate concentration, indicating a zeroth-order dependence on the carbamate concentration in this range. This implied that the CO₂ released from the carbamate, rather than the carbamate itself, is the main reactant in the capt-eCO₂R. Temperatures higher than 60 °C typically favoured H₂ generation and thus reduced CO selectivity (Fig. 4f and g). The similarity between the capt-eCO₂R pathway and the conventional gas-eCO₂R pathway leads to the high performance of the Ni-N/C catalyst in the CO₂-

capturing amine-based solvent, which is attributed to its outstanding intrinsic activity. The performances of Ni-N/C and cAg in the capt-eCO₂R were further investigated in different CO₂ capturing media, such as 1 M 3-amino-1-propanol, 2-(methyloamino)ethanol, AMP, DEA, and KHCO₃, in order to investigate the effect of bulky cations. Considering that alkali metal cations with a smaller effective size have a higher surface density and thus stabilise the CO₂-intermediate, the Ni-N/C exhibited remarkable CO FE performance irrespective of the type of electrolyte. Additionally, the CO FE of cAg decreases significantly as the cation size increases. The weak cation sensitivity of Ni-N/C was further demonstrated in M₂CO₃ electrolytes containing various alkali metal cations (M = Li⁺, Na⁺, K⁺, and Cs⁺) (Fig. 4h and i). This unique feature of Ni-N/C may be derived from its potential for zero charge (PZC), which is higher than that of cAg. The high PZC of Ni-N/C was thought to increase the surface charge density, which in turn reduces the impact of the steric bulk of the cation on surface intermediate stabilisation, thereby showing universal performance. This further highlights the importance of charge density at the EDL in improving the capt-eCO₂R, in addition to its inherent catalytic activity.



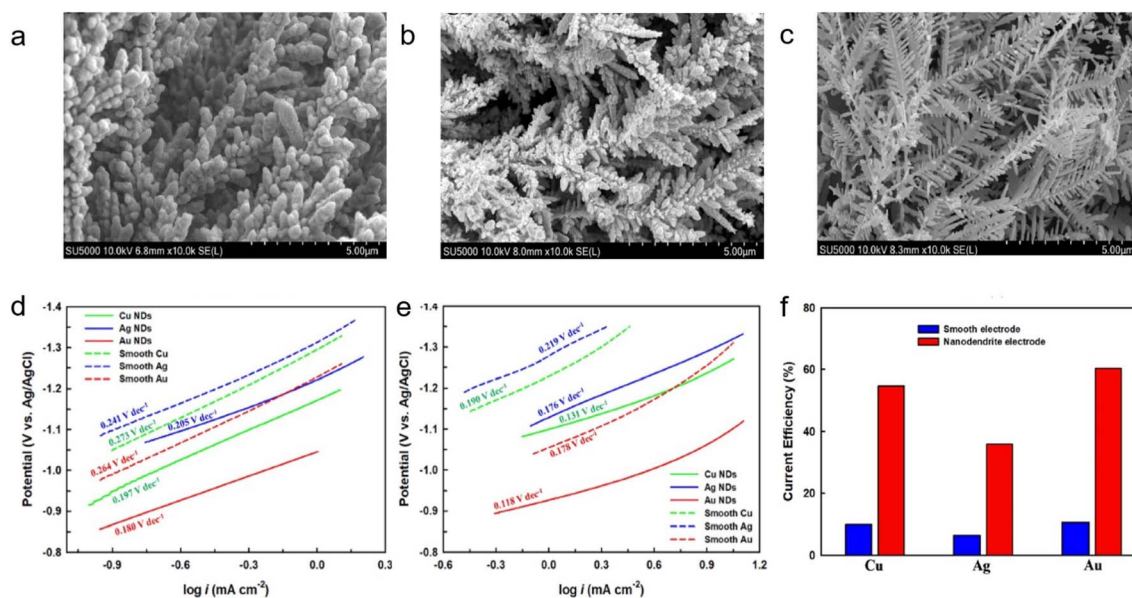


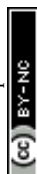
Fig. 5 SEM images of as prepared (a) Cu, (b) Ag, and (c) Au nanodendrites. Tafel slope of various electrodes in CO₂-saturated solutions of (d) 0.1 M NaHCO₃ and (e) 0.05 mol fraction ethanolamine solution. (f) Current efficiency of smooth and nanodendritic electrodes in CO₂-saturated 0.05 mol fraction ethanolamine solution at −1.0 V (vs. Ag/AgCl). Copyright 2021,²⁷ Wiley.

Modification of the catalyst morphology is an effective strategy for improving performance and has been applied to the capt-eCO₂R system. Hossain *et al.* prepared Cu, Ag, and Au nanodendrites and conducted the capt-eCO₂R in CO₂-saturated 0.05 M ethanolamine solutions (Fig. 5a–c).²⁷ The nanodendrites exhibited a high electrochemically active surface area with a large double-layer capacitance, and consequently showed a higher eCO₂R current density than a smooth surface. The Au nanodendrites outperformed the other electrodes, showing the highest current efficiency and lowest onset potential. These properties may arise from the synergistic effect between the dendritic structure and the intrinsic activity of Au, which is a well-known catalyst for CO production. The as-synthesised Au nanodendrites showed the most efficient eCO₂R with a high current density in the linear sweeping voltammetry studies and recorded the lowest Tafel slope (Fig. 5d and e). A comparison of the eCO₂R performance of nanodendrites in CO₂-saturated NaHCO₃ and CO₂-saturated ethanolamine demonstrated that ethanolamine is a promising absorbent for the capt-eCO₂R system. Ethanolamine plays an important role in achieving high current density and catalytic activity because the protonated NH₃⁺ group of ethanolamine increases CO₂ solubility.^{28,29} Furthermore, as-synthesised Au nanodendrites exhibited a significant performance in capt-eCO₂R in ethanolamine, demonstrating its potential as a catalyst in this system (Fig. 5f).

Recent studies have identified the active species in the reduction of CO₂ in monoethanolamine-containing aqueous electrolytes.³⁰ The carbamate moiety is not the only active site in aqueous monoethanolamine, and bicarbonate formation occurred as the partial pressure of CO₂ increased, as detected by ¹³C NMR. This bicarbonate was formed *via* carbamate hydrolysis and was confirmed by the reduction in pH as the

bicarbonate concentration and CO₂ pressure increased. Though the reduction of CO₂ to CO in monoethanolamine is limited by the adsorption of dissolved CO₂, the addition of proton-donating ammonium cations influences the HER. Further study is required to achieve a better understanding of the active state of monoethanolamine during CO₂ reduction.

2.1.2 Other types of solvents media. Although monoethanolamine is the most common industrial CO₂ absorbent, capt-eCO₂R systems can employ other types of solvents. Bhat-tacharya *et al.* studied the ability of a series of primary, secondary, and tertiary amines to utilise carbamates/carbamic acid as a CO₂ source.³¹ The primary amines included tetramethylguanidine (TMG) and aniline, the secondary amines were morpholine, DEA, and dimethylamine (DMA), and the tertiary amines were TREA and triethanolamine (TEOA) (Fig. 6a). The tertiary amine may absorb CO₂ as bicarbonate, as discussed in detail in Section 2.2.2. Employing glassy carbon as the working electrode, the only product produced was formate; CO and H₂ were not detected in this system. The contribution of amines was examined by cyclic voltammetry (CV) in the presence and absence of CO₂. TMG traps more CO₂ than the other amines and therefore drives the equilibrium towards carbamate. TMG reacts as a Brønsted base towards CO₂ owing to its steric bulk, while the other amines are Lewis bases. Upon the addition of CO₂, the Infrared (IR) spectra showed the presence of carbamate species. No carbonate or bicarbonate vibrations were detected in the IR spectra when water was added, indicating that no other species were formed. In this study, TMG was added to stabilise the carbamates of the other selected amines. In the absence of TMG, no significant peaks related to any amines were observed. In the presence of TMG, all the CV data shifted owing to the formation of carbamate. Among the tertiary



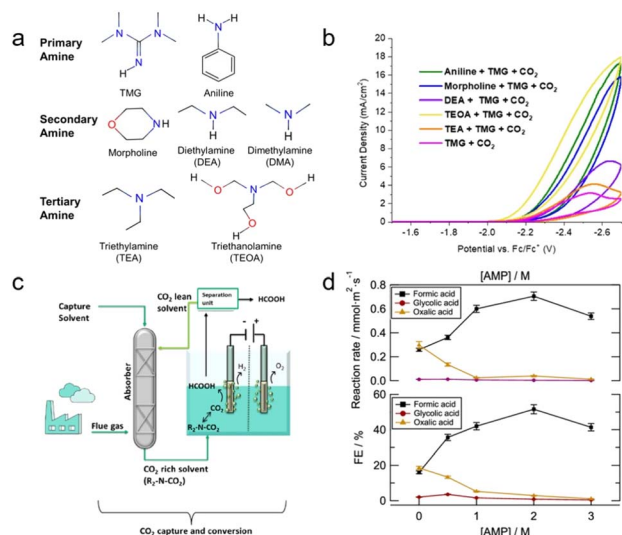


Fig. 6 (a) Primary, secondary, and tertiary amines tested in this experiment. (b) CV responses of various amines in the presence of CO₂ and 0.1 M TMG at a scan rate of 0.1 V s⁻¹, scanning negative. Copyright 2020,³¹ The Electrochemical Society. (c) The proposed CO₂ capture and conversion system. (d) AMP concentration effect of reaction rate and FE of formate, glycolate, and oxalate during electrolysis at -2.5 V vs. Ag/AgCl at 75 °C. Copyright 2021,³² American Chemical Society.

amines, only TEOA shifted the onset potential to a less negative value in the presence of TMG. Aniline, TEOA, and morpholine underwent complexation with CO₂, resulting in higher current densities in CV tests (Fig. 6b). This suggests that these amines have a greater ability to bind redox-active CO₂ in solution. Interestingly, the bicarbonate content increased in the presence of secondary amines and aniline through an anodic shift of the onset potentials when water was added, while in the carbamate content decreased. These amines resulted in a reduction in potential upon the addition of water to the saturated CO₂ solution, indicating that proton or bicarbonate reduction occurs due to the addition of water. The CV data of the secondary amines and aniline exhibited an anodic shift upon the addition of water, except for TMG, indicating the conversion of carbamate species to bicarbonate. The addition of water had a negligible effect on the CV of TREA because no carbamate species were present.

Pérez-Gallent *et al.* developed an integrated system for CO₂ capture and conversion by combining chemical and physical absorption solvents as electrolytes and increasing the temperature of the system in a mixed solution of AMP and propylene carbonate (PC).³² The system is functioned by combining these two solvents, where AMP is CO₂ capturing media and PC is the physical solvent. This method involves heating an electrochemical reactor to liberate the captured CO₂ (Fig. 6c). The concentration of AMP affected the reduction of CO₂ at an optimal concentration of 2 M. Further increasing the concentration of AMP to 3 M results in a highly viscous solution that limits the conductivity and mass transfer properties of the solvent (Fig. 6d). Increasing the temperature also affects the performance of the system,^{24,26} which will be discussed in detail

in Section 3.3. A Pb catalyst produced formic acid with a FE of up to 45% and a reaction rate of 0.56 mmol m⁻² s⁻¹. Recent research has indicated that AMP predominantly captures CO₂ in the carbamate form, which undergoes hydrolysis in an aqueous solution to form bicarbonate.^{33,34} This phenomenon enhances the efficacy of AMP relative to other primary amines. Bicarbonate conversion is further discussed in Section 2.2.

2.2 (Bi)carbonate conversion

Metal hydroxides such as NaOH and KOH chemically absorb CO₂ to form bicarbonates or carbonates depending on the pH of the solution. Such absorbents offer several advantages over amine absorbents, including lower volatility and absence of odour; however, their industrial application is limited by the drawbacks of salt formation during CO₂ conditioning, high regeneration energy, and corrosivity arising from their high alkalinity.^{35–43} Despite these drawbacks, metal hydroxides are currently suitable CO₂ absorbents for DAC systems, owing to their high selective absorption capacity in low-concentration CO₂ environments. Moreover, the application of metal hydroxides to the capt-eCO₂R system can address the high regeneration energy issue because the high-temperature (>700 °C)⁴⁴ calcination process to release captured CO₂ is replaced by electrochemical conversion of bicarbonate/carbonate. Therefore, capt-eCO₂R systems using metal hydroxides are promising systems for achieving net-zero CCUS and carbon-negative direct air capture and utilisation (DACU).

Tertiary amines absorbents chemically absorb CO₂ in the form of bicarbonate. Unlike primary and secondary amines, tertiary amines do not directly bond with CO₂ but act as base catalysts in the CO₂ capture process (Fig. 7). Compared to primary and secondary amines, the C–H bonds of tertiary amines give an energetical advantage over the C–N bonds of primary and secondary amines in the desorption process. Additionally, tertiary amines generally have a higher capacity because one tertiary amine molecule captures one CO₂ molecule, whereas two molecules of primary and secondary amines are required to capture one CO₂ molecule as mentioned in Section 2. The performance of tertiary amine-based systems is expected to depend on the amine type, given that metal hydroxides provide the alkali cations required for electrochemical conversion reactions.

2.2.1 Alkaline media. The direct electrochemical conversion of KHCO₃ is known to exhibit low selectivity and current density in a conventional H-type cell system.^{45,46} Li *et al.* introduced a new reactor configuration to present a new reaction pathway for the direct utilisation of bicarbonate (Fig. 8a).⁴⁷ This method employs a MEA electrolyser with a BPM to separate water into H⁺ and OH⁻ using an induced electric field. The cation- and anion-exchange layers of the BPM contacted the cathode and anode, respectively, to deliver H⁺ to the cathode under a reverse bias (Fig. 8b). This creates an acidic environment at the cathode surface, and consequently, the bicarbonate in the catholyte circulating towards the cathode is instantly converted to CO₂ because of the pH equilibrium reaction. The CO₂ generated in this manner is called *in situ* CO₂. The *in situ*



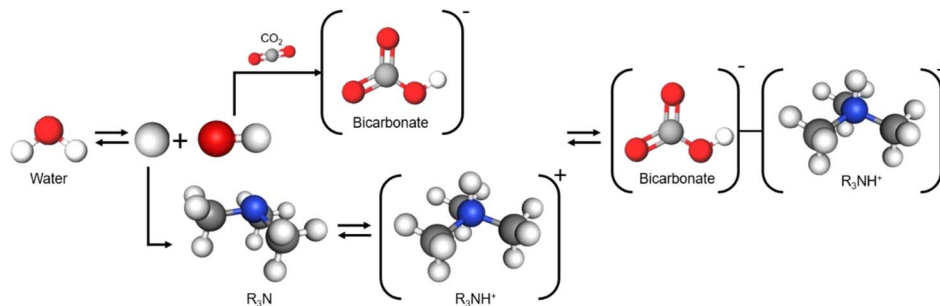


Fig. 7 Representation of the process of CO₂ capture by tertiary amines.

CO₂ receives electrons from the electrode and is reduced to produce CO. The suggested reaction pathway was verified experimentally by replacing the BPM with an anion exchange membrane (AEM) that does not transfer protons in the same cell system. The efficiency of the reactor containing the AEM was significantly lower than that of the reactor containing the BPM, indicating that the reaction occurred *via* the proposed reaction pathway. In this system, the bicarbonate conversion efficiency increased to 81% CO FE at -25 mA cm^{-2} and 37% CO FE at -100 mA cm^{-2} in 3 M KHCO₃ (Fig. 8c), thereby overcoming the low selectivity and activity observed in existing systems. This advancement paves the way for further exploration into the direct conversion of bicarbonate. In a follow-up

study, Lees *et al.* analysed the same flow cell configuration using a continuum model.⁴⁸ The modeling data presented the concentration distribution of bicarbonate ions, CO₂, pH, and CO₃²⁻ from the cation-exchange layer of BPM to the catalyst layer as a function of the current density and water dissociation performance of BPM (Fig. 8d–g). The model supported a previously proposed mechanism⁴⁷ based on simulations and experimental results and also suggests that performance results can be improved by adjusting factors such as the thickness of the catalyst layer and the efficiency of the BPM.

The bicarbonate conversion products were diversified by using different catalyst materials. Similar to conventional gas-eCO₂R, the major products are determined by the type of

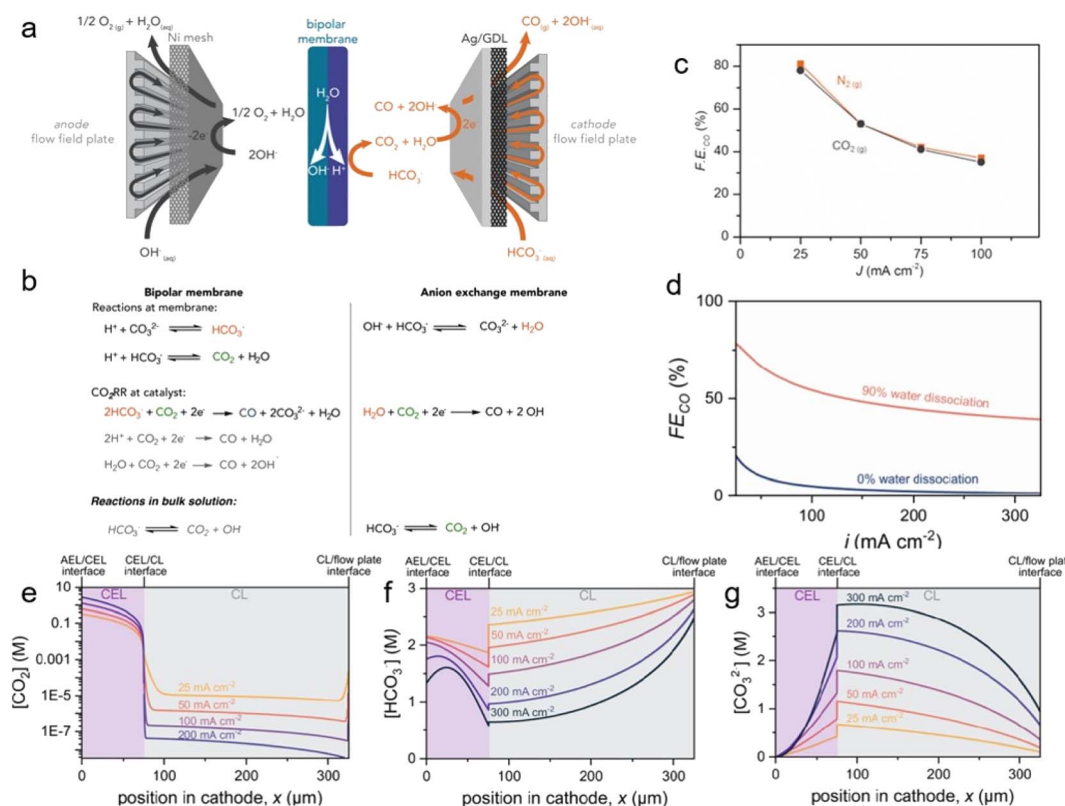
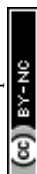


Fig. 8 (a) Schematic of the MEA electrolyser with BPM. (b) The proposed dominant reactions that occur using each membrane. (c) CO FE between 25 and 100 mA cm⁻² in a 3.0 M KHCO₃ solution. Copyright 2019,⁴⁷ Elsevier. (d) CO FE as a function of the water dissociation performance of BPM in the bicarbonate electrolyser cathode. Modeling results of (e) CO₂, (f) HCO₃⁻, (g) CO₃²⁻ concentrations at various current densities. Copyright 2022,⁴⁸ American Chemical Society.



catalyst. CO is primarily produced using Ag catalysts. Zhang *et al.* used a cathode consisting of porous Ag foam that showed high activity of 59% CO FE at -100 mA cm^{-2} in 3 M KHCO_3 (Fig. 9b).⁴⁹ Kim *et al.* reported that Ni–N/C showed significantly higher selectivity (91.0%) than cAg catalyst (28.8%) at $-0.65 \text{ V}_{\text{RHE}}$ in 1 M KHCO_3 (Fig. 9c).²⁶ In addition, Li *et al.* noted that the H_2 :CO ratio of syngas, the feedstock for the Fischer–Tropsch reaction, varied from 2.5 : 1 to 7 : 1, and they produced a 3 : 1 ratio of syngas from CO_2 captured 2 M KOH using a Ag catalyst.⁵⁰ In particular, they also showed operational stability of 145 hours in the system. As a study on the production of formate, Bonet Navarro *et al.* used a bulk Sn catalyst to obtain 18% and 47% formate FE in 1.5 M KHCO_3 in the case of CO_2 pre-saturated and CO_2 pre-saturated 1.5 M KHCO_3 , respectively, at $-1.6 \text{ V vs. Ag/AgCl}$.⁵¹ In addition, Li *et al.* reported that electrodeposited Bi on carbon paper produced formate with 64% FE at -100 mA cm^{-2} and 27% FE at -400 mA cm^{-2} in 3 M KHCO_3 (Fig. 9a).⁵² Lees *et al.* also demonstrated that Cu foam produced methane with a partial current density of $-120 \pm 10 \text{ mA cm}^{-2}$ in $34 \pm 7\%$ yield from capt- eCO_2R in 3 M KHCO_3 with 3 mM CTAB.⁵³ 1D continuum modeling suggests that multi-carbon products are formed if the H^+ supply from the membrane is smoother and the catalyst surface becomes more acidic. To study the multi-carbon products, Lee *et al.* composed a bilayer with one layer consisting of a mixture of Ag catalyst, Nafion, and poly(tetrafluoroethylene) (PTFE) and the other comprising a mixture of Cu catalyst and Sustainion.⁵⁴ A flow reactor equipped with this bilayer recorded $41.6 \pm 0.39\%$ C_{2+} FE at -100 mA cm^{-2} in 3 M KHCO_3 (Fig. 9d).

Another study targeting multi-carbon products found that controlling the microenvironment was crucial, in addition to catalyst design. Lee *et al.* investigated instantaneous CO_2 detachment by a pH equilibrium reaction of the absorbed CO_2 species according to distances between the membrane and the cathode.⁵⁵ The use of a 0.5 M H_2SO_4 anolyte and Nafion as a cation exchange membrane (CEM) with various thicknesses of interposers enabled the construction of well-defined spacing between 135 and 540 μm (Fig. 10a–c). In this configuration, high C_{2+} FE of 38% at a distance of 135 μm and a current density of -300 mA cm^{-2} were obtained in the presence of Cu nanoparticle catalysts (Fig. 10d). Also noteworthy, the CEM lowered the overpotential of the system by facilitating proton transfer to the cathodic side owing to the concentration gradient (Fig. 10e). To enhance C_{2+} production, cobalt phthalocyanines on carbon nanotubes (CoPc–CNTs) was applied to the Cu to produce CO from CO_2 with high turnover frequencies, resulting in a higher C_{2+} FE of 47% at -300 mA cm^{-2} (Fig. 10f). These efficient systems highlight the importance of adjusting the mass transport to improve current density.

2.2.2 Tertiary amine media. Process engineering has been applied to combine the capture and conversion of CO_2 . Diaz *et al.* applied the concept of switchable polarity solvents (SPS) using a tertiary amine, 1-cyclohexylpiperide (CHP), in which the polarity changes upon CO_2 absorption, which results in a change in its water solubility.⁵⁶ Based on this property, a reversible amine cycle can be established when CHP is applied as a CO_2 absorbent in the capt- eCO_2R system (Fig. 11a). A CEM (Nafion) applied MEA with a buffer layer between CEM and cathode was prepared, along with a 1.25 M CHP- H_2CO_3 solution as a catholyte by

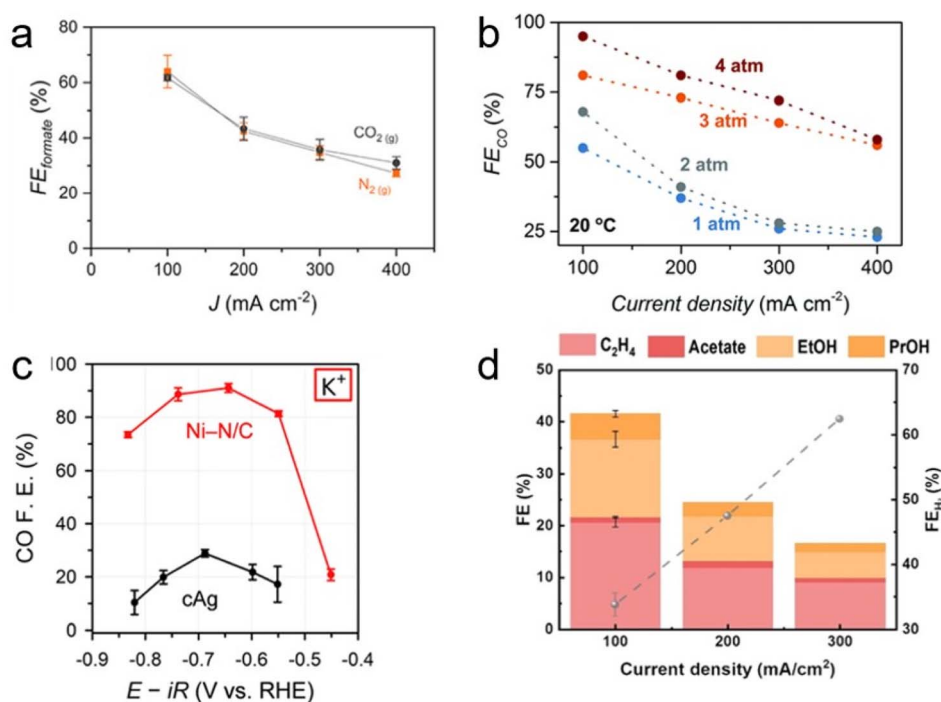


Fig. 9 (a) Formate FE as a function of current density in 3.0 M KHCO_3 solution purged with $\text{CO}_2(\text{g})$ or $\text{N}_2(\text{g})$. Copyright 2020,⁵² American Chemical Society. (b) CO FE as a function of pressure at different current densities. Copyright 2022,⁴⁹ Royal Society of Chemistry. (c) CO FE of Ni–N/C and cAg in 1 M KHCO_3 . Copyright 2022,²⁶ Royal Society of Chemistry. (d) FE of products as a function of current density. Copyright 2022,⁵⁴ Wiley.

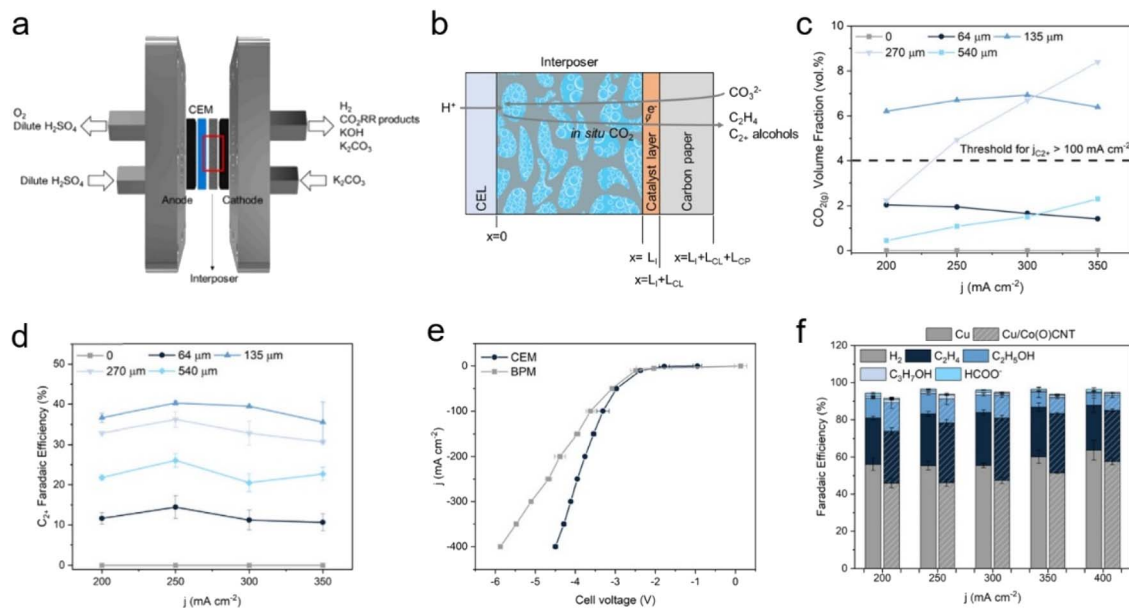


Fig. 10 (a) Schematic of CO_3^{2-} -fed electrolyser. (b) Representation of the cation-exchange layer, interposer, catalyst layer, and carbon paper. (c) $\text{CO}_2(\text{g})$ volume fraction under different spacing conditions and current densities. (d) C_2+FE of CO_3^{2-} -fed electrolyser with Cu electrocatalyst. (e) Full cell j - V curve with Cu electrocatalyst with a CEM (Nafion) and a BPM in 1.5 M K_2CO_3 solution with 135 μm interposer. (f) Product distribution of Cu electrocatalyst and Cu/CoPc-CNTs catalyst at applied current densities ranging from 200 to 400 mA cm^{-2} . Copyright 2023,⁵⁵ Elsevier.

capturing CO_2 in CHP. This concentration (1.25 M) $\text{CHP-H}_2\text{CO}_3$ affords the maximum concentration of CO_2 and ionic conductivity. In this configuration, a Ag catalyst showed approximately 30% CO FE at -104 mA cm^{-2} (Fig. 11b). These studies of bicarbonate reduction activity in amine solutions elucidate the behaviour and activity of tertiary amines in the capt- eCO_2R .

Langie *et al.* developed a unique bicarbonate conversion system known as a reaction swing absorption (RSA) using TREA as a CO_2 absorbent.¹⁷ The tertiary amine TREA was used in place of a metal hydroxide owing to its high selectivity towards bicarbonate and industrially applicable CO_2 absorption rate. TREA demonstrated a CO_2 absorption rate of 84.5% at a flow rate of $0.8 \text{ m}^3 \text{ h}^{-1}$ with 3% CO_2 , and 95.1% at $0.5 \text{ m}^3 \text{ h}^{-1}$ with 3% CO_2 , measured in an absorption tower with a height of 3 m (Fig. 12a and b). Various membranes, substrates, and cathode catalysts were evaluated to identify the viable configuration for the utilisation of TREA-captured CO_2 . The optimal catalyst for syngas consisted of hydrophilic nano-coral structured Ag with carbon (coral Ag/C), which achieved a CO FE of 68.7% at -20 mA cm^{-2} and 29.2% at

-200 mA cm^{-2} in a BPM equipped MEA electrolyser (Fig. 12c). Remarkably, the CO_2 concentration in the outlet gas is nearly zero, highlighting the effective absorption of TREA and its suitability as a CO_2 absorbent for the capt- eCO_2R system. Also, they have reported up to 70 hours of stable performance in TREA capt- eCO_2R through the use of optimized catalyst. The feasibility of the RSA system compares favourably with traditional CO_2 -to-syngas technologies, such as the reverse water-gas shift reaction and gas- eCO_2R , according to TEA and LCA. RSA is in the early stages of development and thus incurs higher capital expenditure (CAPEX) (Fig. 12d); however, it offers the lowest OPEX and break-even prices among the evaluated CCU processes (Fig. 12e). In the optimistic scenario, with potential improvements in CO FE and reduced voltage, the break-even price of RSA could drop to as low as \$0.65 per kg of syngas, significantly lowering both the CAPEX and OPEX. The RSA also showed the highest net present value (NPV), indicating a substantial positive increase in profitability (Fig. 12f). LCA showed that RSA had the lowest GWP in all tested scenarios, potentially being as low as $0.27 \text{ kg CO}_2 \text{ eq. kg}^{-1}$ of syngas, moving

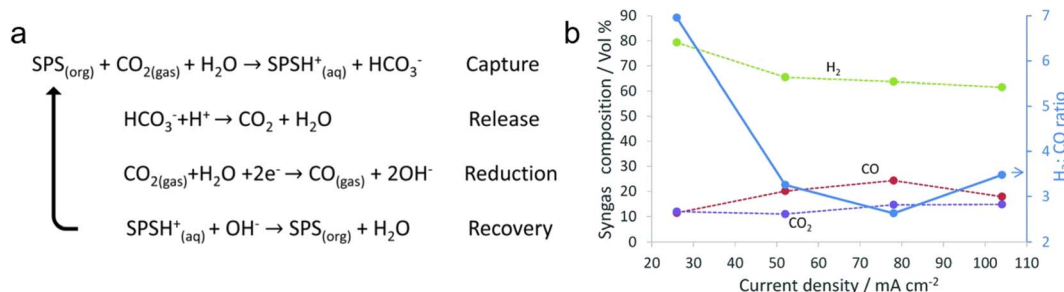


Fig. 11 (a) A proposed amine cycle. (b) $\text{H}_2 : \text{CO}$ ratio of the syngas obtained at different current densities. Copyright 2018,⁵⁶ Royal Society of Chemistry.



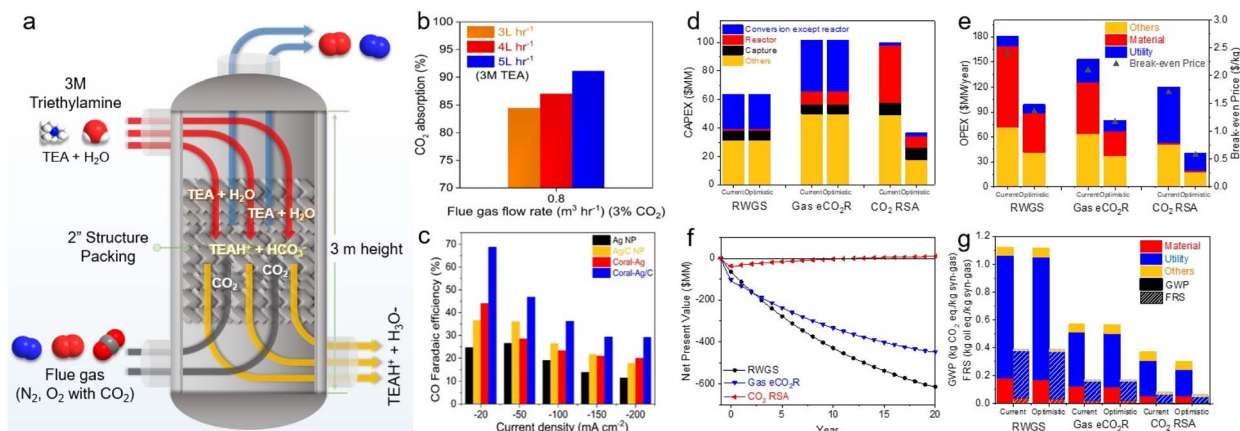


Fig. 12 (a) Schematic of the CO₂ absorption column used in the capture experiments. (b) CO₂ absorption rates at various TREA flow rates at fixed gas flow rate and CO₂ concentration. (c) CO FE of Ag nanoparticle, Ag/C, coral-Ag, and coral Ag/C in 3 M TREA. (d) CAPEX of three processes. (e) OPEX and break-even price of three processes. (f) LCA of the current and optimistic scenarios of three processes. (g) NPV of an optimistic scenario with a selling price of \$0.8 per kg syngas. (h) LCA of the current and optimistic scenarios of three processes. Copyright 2022,¹⁷ Springer Nature.

towards net-zero CO₂ emissions (Fig. 12g). These findings highlighted the economic and environmental advantages of bicarbonate systems. Recent achievements in bicarbonate conversion are summarised in Table 3.

To investigate the dependence of the electrochemical activity on the type of tertiary amine, Hosseini *et al.* analysed solutions of monoethanolamine, diethylenetriamine (DETA), diisopropylamine (DIPA), and aminoethylpiperazine (AEP) with and without the addition of bicarbonate using CV.⁵⁷ Of these; only DIPA showed a significant catalytic effect on bicarbonate reduction, then the mechanism of carbonate reduction in DIPA was explored electrochemically by scan rate control experiments and EIS analysis.

3 Strategy for improve performance of capt-eCO₂R

Recent studies that aimed to identify the active characteristics of primary amine-based systems have proposed that the release or dissolution of CO₂ may constitute an active component rather than the carbamate itself. Increasing the ratio of free CO₂ to the captured absorbent is important to promote more efficient

utilisation regardless of the absorbent type. Similar strategies have therefore been employed to improve the performance of capt-eCO₂R system, regardless of the type of CO₂ absorbent.

3.1 Control the wettability of electrodes

In the case of an MEA cell or flow cell for the gas-eCO₂R, where reactions occur at a triple-phase boundary, various strategies have been applied to reduce the involvement of water on the solid electrode surface to inhibit the HER and flooding.^{60–62} For example, porous carbon paper treated with hydrophobic materials, such as PTFE and microporous layers (MPL), is preferred in the gas-eCO₂R system because it alleviates flooding and enables gaseous CO₂ to access the catalyst layer. In contrast, the capt-eCO₂R requires different mass transport approaches to achieve feasibility because the reactant is delivered in the liquid phase rather than the gas phase. Hydrophobic treatment of the carbon substrate inhibits intimate contact between the liquid reactants and the catalyst layer and may therefore have the opposite effect.

Lees *et al.* compared the performance of Ag catalysts on carbon papers under different PTFE and MPL conditions.⁵⁹ Placing the Ag catalyst on carbon paper without PTFE and MPL

Table 3 The capt-eCO₂R performance of reported bicarbonate electrolyzers

Catalyst	Catholyte	Product	Faradaic efficiency (%)	Current density (mA cm ⁻²)	Temperature	Pressure	Reference
Ag	1.25 M CHP-H ₂ CO ₃	CO	30	−104	25 °C	40 psig	56
Coral Ag/C	3 M TREA	CO	30	−200	R.T.	Ambient	17
Ag foam	1.5 M CsHCO ₃	CO	80	−100	R.T.	Ambient	58
Ag	3 M KHCO ₃	CO	82 ± 2	−100	R.T.	Ambient	59
Ag foam	3 M KHCO ₃	CO	95	−100	R.T.	4 atm	49
Bi	3 M KHCO ₃	HCOO [−]	62 ± 1	−100	R.T.	Ambient	52
Cu foam	3 M KHCO ₃	CH ₄	30	−400	R.T.	Ambient	53
Cu/Ag bilayer	3 M KHCO ₃	C ₂ +	41.6 ± 0.39	−100	R.T.	Ambient	54
Cu/CoPc-CNTs	1.5 M K ₂ CO ₃	C ₂ +	47	−300	R.T.	Ambient	55



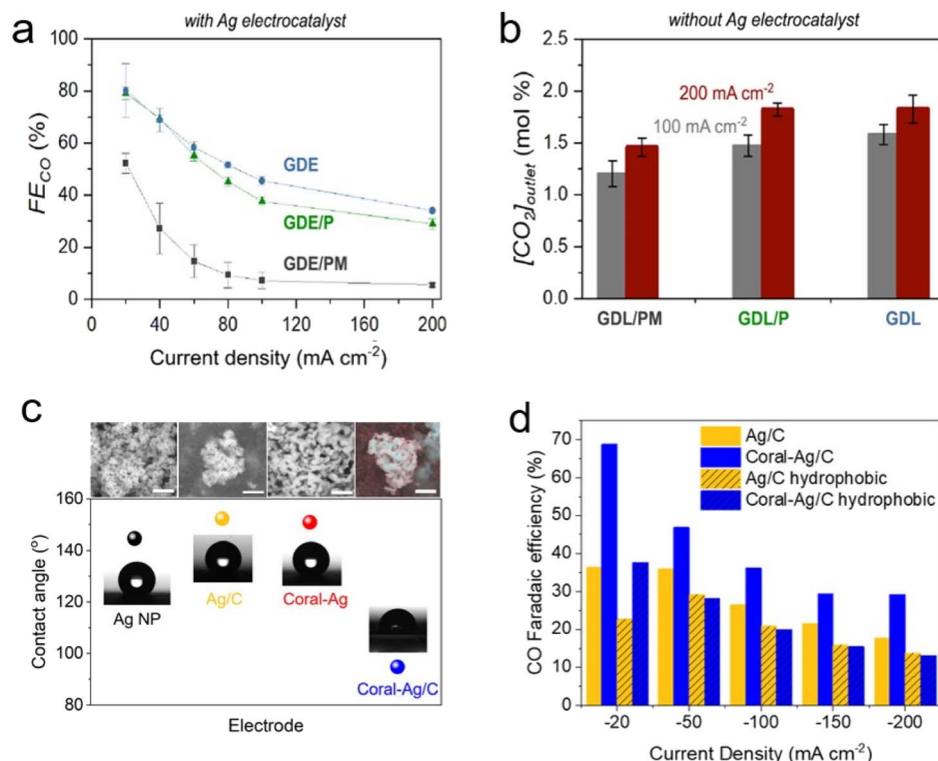


Fig. 13 (a) CO FE of three different electrode conditions; GDE with neither PTFE nor an MPL, GDE/P containing PTFE but with an MPL, and GDE/PM containing both PTFE and an MPL. (b) The concentration of CO₂ in the outlet gas under the three electrode conditions. Copyright 2020,⁵⁹ American Chemical Society. (c) SEM images (scale: 500 nm) and contact angles of all Ag electrodes. (d) CO FE of prepared hydrophobic electrodes in the RSA system displayed with those of Ag/C and coral Ag/C. Copyright 2022,¹⁷ Springer Nature.

achieved the highest performance (Fig. 13a and b), indicating that a well-wetted environment facilitates the approach of the reactants to the electrode in the capt-eCO₂R. Based on this observation, Zhang *et al.* used porous Ag foam as an electrode, which showed significantly higher performance than the control carbon cloth as mentioned in Section 2.2.1.⁴⁹ Langie *et al.* also constructed an MEA cell using carbon papers that were not treated with PTFE or MPL.¹⁷ To enhance the hydrophilicity of the catalyst and improve CO production, the coral Ag/C was prepared through electrochemical oxidation and reduction processes. The performance of coral Ag/C was compared with Ag nanoparticles, coral Ag, and Ag/C to determine the origin of the high performance associated with the coral Ag/C. They found that the hydrophilicity of the catalyst material, through the transition from Ag nanoparticles to coral Ag and Ag/C structures, along with the measurement of the contact angle with water, contributes significantly to the performance improvement (Fig. 12c and 13c). Additionally, the CO production performance of various carbon papers, both treated and untreated with PTFE, showed a consistent trend where hydrophobic carbon paper resulted in reduced CO production (Fig. 13d).

3.2 Addition of various additives

CO₂ absorbents such as amino-alcohol solutions can interfere with the catalyst surface in the capt-eCO₂R system. As explained in Section 2.1.1, Lee *et al.* observed an increase in performance

by adding alkali cation, especially K⁺, to the monoethanolamine solution, and observed the effect of alkali cation on the composition of electrode surface EDL through Raman spectroscopy and EIS analysis.²⁴ These results demonstrate that while bulky ammonium cations inhibit reaction rates by preventing the direct transfer of electrons from the cathode to the reactants, this can be overcome by adding alkali cations.

Fink *et al.* confirmed a cation effect similar to that observed in conventional gas-eCO₂R by adjusting the type and concentration of alkali cations in MHCO₃ (M = Li, Na, K, Cs) solutions.⁵⁸ Furthermore, the influence of the cation on the chemical desorption and electrochemical conversion of CO₂ was investigated. The amount of *in situ* CO₂ was similar regardless of the identity of the cation; however, larger cations increased the CO selectivity, consistent with the known trend of the cation effect. They concluded that solvated alkali cations affect the stabilisation of the reaction intermediate, similar to the gas-eCO₂R.

The organic chains of cationic surfactants are known to inhibit the HER, which competes with the eCO₂R; thus, cationic surfactants have been used as alternative additives to metal cations. Banerjee *et al.* reported that a cationic surfactant suppressed the HER in the gas-eCO₂R by adsorbing onto the surface of the cathode and thereby preventing the approach of water or hydrated cations in the electrolyte.⁶³ Several similar studies using surfactants have been conducted in the capt-eCO₂R. Lees

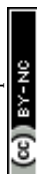


Table 4 Structure and properties of the cationic surfactants

Surfactant	Abbreviation	Charge	Counter ion	Structure
Hexadecyl trimethylammonium bromide	CTAB	+	Br [−]	
Benzyltrimethyl hexadecylammonium chloride	CKC	+	Cl [−]	
Sodium dodecyl sulphate	SDS	−	Na ⁺	
Hexadecyltrimethylammonium chloride	CTAC	+	Cl [−]	

et al. added 3 mM CTAB to 3 M KHCO₃ to convert captured CO₂ to methane and compared this method to that using a solution without CTAB using a porous copper electrode.⁵³ The selectivity of the conversion reaction increased from nearly 0% to approximately 30%. In another study, Gutiérrez-Sánchez *et al.* compared the influence of four different surfactants, CTAB, CKC, SDS, and CTAC, on the formate production performance (Table 4) using a Sn wire as a working electrode.⁶⁴ In 2 M KHCO₃, the 1000 μM of CKC addition resulted in the highest activity, increasing the FE from 12% to 66% at −0.9 V_{RHE}. The aromatic ring of CKC was hypothesised to increase resistance to H⁺, thereby improving the FE by more than 20% relative to CTAB, which exhibited the second-best performance.

3.3 Temperature and pressure

During capt-eCO₂R, desorption of CO₂ from the absorption medium is expected to be necessary for electrochemical

conversion into a product. Optimising the physical conditions, such as temperature and pressure, should improve the capt-eCO₂R performance by facilitating the breaking of chemical bonds between the absorbent and CO₂; however, the additional energy required for this process must be considered to ensure overall energy efficiency.

The optimal temperatures vary according to the components and target products of the capt-eCO₂R because temperature affects multiple properties, including the desorption–absorption equilibrium, catalytic reaction rates, and mass transfer. Lee *et al.* adjusted the temperatures of a flow cell system with a Ag catalyst as a cathode.²⁴ The current density at 60 °C was 15 times higher than at room temperature (RT) and CO FE also increases with temperature (Fig. 14a and b). The additional thermal energy facilitated cleavage of the C–N bond between the amine and CO₂, and accelerated the transfer of reactants. Kim *et al.* studied the effect of temperature in capt-eCO₂R at RT, 40 °C,

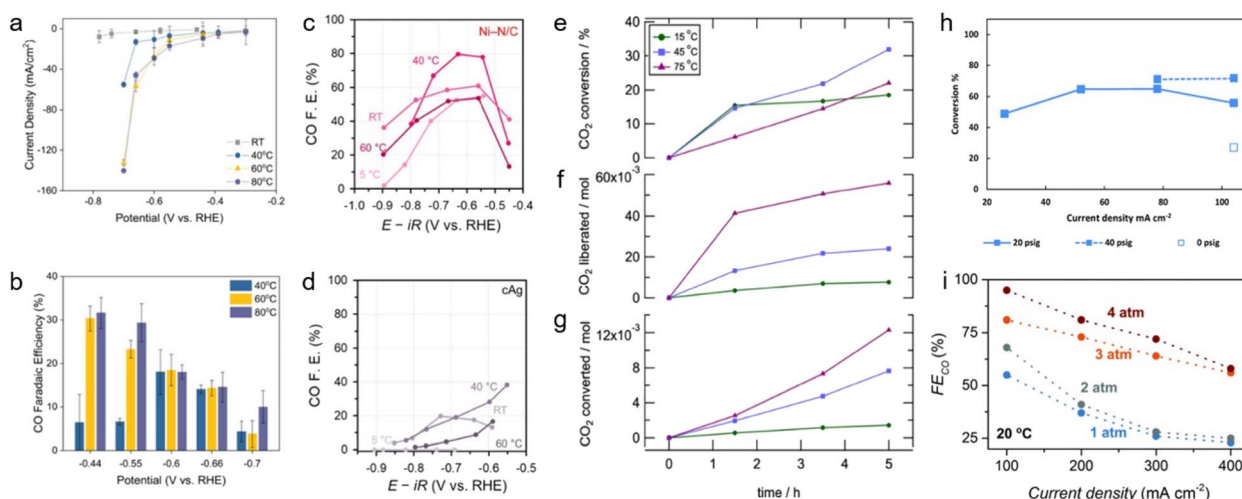


Fig. 14 (a) *j*–*V* curve, (b) CO FE of 2 M monoethanolamine with 2 M KCl electrolyte at different temperatures. Copyright 2020,²⁴ Springer Nature. CO FE of (c) Ni–N/C, and (d) cAg at different temperatures. Copyright 2022,²⁶ Royal Society of Chemistry. (e) CO₂ conversion, (f) liberated CO₂, and (g) converted CO₂ on a Pb electrode in a 1 M AMP in PC solution as functions of time and temperature. Copyright 2021,³² American Chemical Society. (h) Conversion profiles of CO₂ released from the CP–H₂CO₃ solution at different current densities and pressures. Copyright 2018,⁵⁶ Royal Society of Chemistry. (i) FE_{CO} as a function of pressure at different current densities of Ag foam in 3 M KHCO₃. Copyright 2022,⁴⁹ Royal Society of Chemistry.



and 60 °C and found that the capt-eCO₂R demonstrated peak efficiency at 40 °C (Fig. 14c and d).²⁶ A higher temperature of 60 °C had positive effects on the desorption of CO₂ but also simultaneously accelerated HER; thus, 40 °C was determined to be the optimal temperature for the capt-eCO₂R. In addition, Pérez-Gallent *et al.* estimated the CO₂ conversion rate, along with the amounts of liberated and converted CO₂ at 15 °C, 45 °C, and 75 °C with a Pb electrode in 1 M AMP and PC solution.³² The highest CO₂ liberation and conversion amounts were achieved at 75 °C, demonstrating 8 times higher performance than at 15 °C (Fig. 14f and g). Additionally, the highest CO₂ conversion rate was observed at 45 °C (Fig. 14e). While additional thermal energy assists in the desorption and reactions of CO₂, the rate of CO₂ conversion has an optimal temperature of 45 °C.

Pressure also significantly affects the solubility and reaction rates of gaseous molecules. To understand the effect of pressure effect on capt-eCO₂R, Diaz *et al.* measured the CO₂ conversion ratio at various pressures of 0, 20, and 40 psig at a current density of -104 mA cm^{-2} in a CHP-H₂CO₃ mixed solution.⁵⁶ The highest CO₂ conversion of 71.4% was obtained at 40 psig. High pressure facilitated the redissolution of unreacted CO₂ and its participation in the reaction (Fig. 14h). Zhang *et al.* improved the performance on an MEA cell with a porous Ag foam cathode at a constant current density of -100 mA cm^{-2} , by tuning the pressure and temperature.⁴⁹ Increasing the temperature from 20 °C to 70 °C increased the CO FE from 59% to 78%, while increasing the pressure from 1 atm to 4 atm increased the CO FE from 55% to 95% (Fig. 14i). Temperature may affect CO₂ desorption and mass transfer, whereas increasing the pressure increases the kinetic supply and reaction rate of CO₂.

3.4 Other modification

The electrode surface was modified to use the flue gas directly *via* physical separation rather than chemical separation. Al-Attas *et al.* fabricated a permselective GDE composed of a sputtered Ag catalyst on one side and a metal-organic framework (MOF) mixed with Nafion on the other.⁶⁵ Owing to its strong ability to physisorb CO₂, the MOF called Calgary framework-20 (CALF-20) was used to increase CO₂ permselectivity among the multiple gases that make up the flue gas. With the introduction of 10% CO₂ into the N₂-based gas stream, the permselective GDE showed 95% CO FE while the Ag/PTFE electrode showed only 58% CO FE at $-1.32 \text{ V}_{\text{RHE}}$. CALF-20 showed superior performance in suppressing the oxygen reduction reaction with an oxygen-containing gas stream and improved the wet stability with a wet gas stream.

Kim *et al.* demonstrated that the activity of a given electrode is strongly dependent on the electrolyte composition because it can overcome the interference of the bulky cations typically present in CO₂ absorbents.²⁶ Developing catalyst materials affords control over the EDL and enables the manipulation of PZC of the electrode, which determines the electric field required for ion accumulation under electrocatalytic reaction conditions and tailored the catalyst to be insensitive to the cation species, as explained in Section 2.1.1. Developing catalyst materials is a promising strategy to enhance performance because it can be

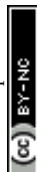
implemented into existing systems and circumvents the undesirable side effects of additives such as salt formation.

4. Perspective

The capt-eCO₂R system is a promising system that complements the weaknesses of both CO₂ capture and CO₂ conversion systems. In the field of CO₂ capture, capt-eCO₂R systems eliminate the requirement for energy-intensive solvent regeneration, gas conditioning, and compression. Concurrently, because the majority of unreacted CO₂ remains within the capture medium, eCO₂R systems significantly improve the economic feasibility of eCO₂R by simplifying or eliminating product separation and recycling process. Although the capt-eCO₂R seems a seamless integration of CO₂ capture and CO₂ conversion, coupled studies considering both reactions thoroughly are essential for developing an advanced, high-performance system.

In the CO₂ capture process, the captured CO₂ is regenerated by environmental factors such as temperature and pressure. The capt-eCO₂R exploits the proton conductivity of the capture medium such that CO₂ can be regenerated and activated by pH changes or applied potentials. To date, various CO₂ absorbents, such as primary, secondary, and tertiary amine, as well as metal hydroxides, have been preferentially screened as well-known CO₂ capture materials, while mixed absorbents or additives have been applied to improve the performance of capt-eCO₂R systems. However, in the capt-eCO₂R system, the CO₂ absorbent acts as both a CO₂ absorbent and an electrolyte. Given the active participation of the electrolyte-like cation effect observed in the capt-eCO₂R system, further research is required to elucidate the role of the absorbent and identify or develop an ideal absorbent for the capt-eCO₂R. In the screening of a promising CO₂ absorbent for the capt-eCO₂R, we propose to consider multiple properties that specifically address the binding energy and electroconductivity, such as pK_a , molecular structure and size, solvation effect, and charge distribution. An understanding of the effects of these parameters is expected to afford control over the strength of CO₂ binding, adsorption, and desorption of the reactants and products, as well as interactions with the catalyst surface. Beyond empirical approaches for selecting a CO₂ absorbent, further studies are required to understand the key parameters and their respective impacts.

The electrolyser configuration must also be redesigned to enhance the circulatory efficiency of CO₂ absorption, desorption, and conversion. Recent studies commonly employ MEA-type electrolyzers for capt-eCO₂R systems; however, because MEA electrolyzers are designed for use with gas-phase CO₂ rather than liquid reactants, the system configuration must be adjusted to ensure their optimal applicability in capt-eCO₂R systems. Strongly hydrophobic substrates appear to limit the interaction between the liquid reactant and the catalyst, indicating that superior performance may be achieved using hydrophilic substrates (Section 3.1). This discrepancy may arise from the current designs of the flow path and substrate location, which are optimised for gas delivery. Consequently, a new generation of electrolyzers more suitable for liquid reactants is required. In addition, most studies employed carbon



substrates, while other types of substrates remain relatively unexplored.

The selection and development of membranes should be a major focus of future work to improve the performance of the MEA configuration. Current capt-eCO₂R systems typically use BPM as a membrane owing to its high H⁺ and OH[−] reflux to each cathode and anode, respectively, which provides sufficient acidity at the cathode side to enable the release and conversion of CO₂. The BPM would have been a good choice to ensure good separation, because most previous studies conducted the cathodic and anodic reactions under different electrolyte conditions. However, the structure of the BPM overlaps with those of the CEM and AEM, resulting in a high resistance due to its thickness, where consequently limits the current density that the BPM can provide to approximately 600 mA cm^{−2}.⁶⁶ Recent progress in enhancing the performance of the BPM can ensure the longevity of the capt-eCO₂R; however, a newly designed or configured electrolyser that does not rely on a BPM should also be considered. Recently, a novel CEM-based system with an interposer was introduced to increase the concentration of *in situ* CO₂ at the desired local pH at the catalyst; however, this configuration still requires a high overpotential owing to the presence of an interposer.⁵⁵ A new membrane optimised for the capt-eCO₂R system should therefore be developed to generate a higher current density and low overpotential.

A thorough understanding of the overall reaction mechanisms could also provide a breakthrough in the development of this system. Opinions regarding the identity of the active species in the capt-eCO₂R differ. Some studies have suggested that captured CO₂ is reduced directly; however, (bi)carbonate/carbamate are generally considered to be preferentially regenerated to *in situ* CO₂ that is then reduced on the catalyst surface. Further reductions to afford the desired products are expected to be similar to those in the gas-eCO₂R system; however, this remains to be conclusively demonstrated. Hence, a comprehensive investigation of these mechanisms requires direct observations by *operando/in situ* spectroscopy, alongside thermodynamic calculations and multiphysics modeling. Fundamental research in this area can provide valuable insights that inform the development of effective strategies and identify promising approaches.

In the current state of research, significant emphasis has been placed on advancing system development and catalyst engineering to enhance the performance of capt-eCO₂R. Given the relatively early stage of development in this field, only a few publications reported on the long-term stability of their system comprehensively.^{17,50} To achieve a long-term stable capt-eCO₂R system, we may encounter challenges similar to those in conventional gas-eCO₂R systems, such as catalyst degradation, membrane stability, and electrolyte changes during prolonged operation. Furthermore, depending on the CO₂ absorbent used, different degradation phenomena and their degrees may be expected, necessitating a thorough investigation of the effects of the CO₂ absorbent. For instance, with a KOH solution, salt formation on the catalyst surface and flow path is probably unavoidable during extended operation.^{38,40,42} Amine-type absorbents may alleviate this issue, but some amines can

potentially damage the catalyst and carbon substrate due to their corrosiveness. Moreover, it is essential to consider that catalysts can degrade due to fluctuations in reactant concentration and physical delamination under a liquid reactant circulating system. Therefore, observing the phenomenon of prolonged reactions should be a primary focus, followed by comprehensive studies to address it. All components of the electrochemical system, including the membrane, cathode, anode, and electrolyte, are crucial areas that require further study for long-term stability.

This technology is still in the early stages of research and development; thus, its performance is currently inferior to that of gas-eCO₂R technologies. Nevertheless, significant performance improvements are expected over time as the development of the aforementioned component technologies continues to mature, driven by empirical insights gained through gas-CO₂R research. In addition, the simplicity of this system will afford economic and environmental benefits that far outstrip those of existing CCUS technologies.

Author contributions

K. L. and G. B. contributed equally to this work, collected and analysed data, and wrote the manuscript. U. L., D. K. L., and C. W. L. contributed to the investigation of recent research trends and wrote the perspective part. Y. J. H. and D. H. W. supervised this study.

Conflicts of interest

The authors declare no competing financial interest.

Acknowledgements

This work was supported by “DACU project” (No. RS-2023-00259920), “Carbon to X Project” (Project No. 2020M3H7A1098229), and NRF-2022R1A2B5B02001380 from the National Research Foundation of Korea (NRF) grant, funded by Ministry of Science and ICT, Republic of Korea, and a KIST institutional project.

References

- 1 WMO, *WMO Greenhouse Gas Bulletin (GHG Bulletin) – No. 17: the State of Greenhouse Gases in the Atmosphere Based on Global Observations through 2020*, WMO, 2021.
- 2 O. Gutiérrez-Sánchez, B. Bohlen, N. Daems, M. Bulut, D. Pant and T. Breugelmans, *ChemElectroChem*, 2022, **9**, e202101540.
- 3 A. Dubey and A. Arora, *J. Cleaner Prod.*, 2022, **373**, 133932.
- 4 A. S. Reis Machado and M. Nunes da Ponte, *Curr. Opin. Green Sustainable Chem.*, 2018, **11**, 86–90.
- 5 G. Y. Duan, X. Q. Li, G. R. Ding, L. J. Han, B. H. Xu and S. J. Zhang, *Angew Chem. Int. Ed. Engl.*, 2022, **61**, e202110657.
- 6 M. G. Kibria, J. P. Edwards, C. M. Gabardo, C. T. Dinh, A. Seifitokaldani, D. Sinton and E. H. Sargent, *Adv. Mater.*, 2019, **31**, e1807166.



- 7 C.-T. Dinh, F. P. García de Arquer, D. Sinton and E. H. Sargent, *ACS Energy Lett.*, 2018, **3**, 2835–2840.
- 8 L. Fan, C. Xia, P. Zhu, Y. Lu and H. Wang, *Nat. Commun.*, 2020, **11**, 3633.
- 9 B. Zhang, J. Zhang, M. Hua, Q. Wan, Z. Su, X. Tan, L. Liu, F. Zhang, G. Chen, D. Tan, X. Cheng, B. Han, L. Zheng and G. Mo, *J. Am. Chem. Soc.*, 2020, **142**, 13606–13613.
- 10 T. Jaster, A. Gawel, D. Siegmund, J. Holzmänn, H. Lohmann, E. Klemm and U. P. Apfel, *iScience*, 2022, **25**, 104010.
- 11 P. Yue, Q. Fu, J. Li, X. Zhu and Q. Liao, *Green Chem.*, 2022, **24**, 2927–2936.
- 12 J. Fernández-González, M. Rumayor, A. Domínguez-Ramos and Á. Irabien, *Int. J. Greenhouse Gas Control*, 2022, **114**, 103549.
- 13 Y. Hori, H. Konishi, T. Futamura, A. Murata, O. Koga, H. Sakurai and K. Oguma, *Electrochim. Acta*, 2005, **50**, 5354–5369.
- 14 R. Veneman, *Adsorptive System for Post-combustion CO₂ Capture*, University of Twente, Enschede, 2015.
- 15 F. Yulia, R. Sofianita, K. Prayogo and N. Nasruddin, *Case Stud. Therm. Eng.*, 2021, **26**, 101093.
- 16 A. J. Welch, E. Dunn, J. S. DuChene and H. A. Atwater, *ACS Energy Lett.*, 2020, **5**, 940–945.
- 17 K. M. G. Langie, K. Tak, C. Kim, H. W. Lee, K. Park, D. Kim, W. Jung, C. W. Lee, H.-S. Oh, D. K. Lee, J. H. Koh, B. K. Min, D. H. Won and U. Lee, *Nat. Commun.*, 2022, **13**, 7482.
- 18 N. S. J. Gao, C. Quiroz-Arita, L. A. Diaz and T. E. Lister, *J. CO₂ Util.*, 2021, **43**, 101365.
- 19 Z. Rastegar and A. Ghaemi, *J. Heat Mass Transf. Res.*, 2021, **58**, 365–381.
- 20 N. El Hadri, D. V. Quang, E. L. V. Goetheer and M. R. M. Abu Zahra, *Appl. Energy*, 2017, **185**, 1433–1449.
- 21 D. Fernandes, W. Conway, R. Burns, G. Lawrance, M. Maeder and G. Puxty, *J. Chem. Thermodyn.*, 2012, **54**, 183–191.
- 22 L. B. Hamdy, C. Goel, J. A. Rudd, A. R. Barron and E. Andreoli, *Mater. Adv.*, 2021, **2**, 5843–5880.
- 23 B. Yoon, D. C. Calabro, L. S. Baugh, S. Raman and G. S. Hwang, *J. Environ. Chem. Eng.*, 2022, **10**, 108987.
- 24 G. Lee, Y. C. Li, J.-Y. Kim, T. Peng, D.-H. Nam, A. Sedighian Rasouli, F. Li, M. Luo, A. H. Ip, Y.-C. Joo and E. H. Sargent, *Nat. Energy*, 2020, **6**, 46–53.
- 25 L. Chen, F. Li, Y. Zhang, C. L. Bentley, M. Horne, A. M. Bond and J. Zhang, *ChemSusChem*, 2017, **10**, 4109–4118.
- 26 J. H. Kim, H. Jang, G. Bak, W. Choi, H. Yun, E. Lee, D. Kim, J. Kim, S. Y. Lee and Y. J. Hwang, *Energy Environ. Sci.*, 2022, **15**, 4301–4312.
- 27 M. N. Hossain, S. Ahmad, I. S. da Silva and H. B. Kraatz, *Chemistry*, 2021, **27**, 1346–1355.
- 28 Y. Qiu, H. Zhong, W. Xu, T. Zhang, X. Li and H. Zhang, *J. Mater. Chem. A*, 2019, **7**, 5453–5462.
- 29 M. Abdinejad, Z. Mirza, X.-a. Zhang and H.-B. Kraatz, *ACS Sustain. Chem. Eng.*, 2020, **8**, 1715–1720.
- 30 G. Leverick, E. M. Bernhardt, A. I. Ismail, J. H. Law, A. Arifuzzaman, M. K. Aroua and B. M. Gallant, *ACS Catal.*, 2023, **13**, 12322–12337.
- 31 M. Bhattacharya, S. Sebghati, Y. M. Vercella and C. T. Saouma, *J. Electrochem. Soc.*, 2020, **167**, 086507.
- 32 E. Pérez-Gallent, C. Vankani, C. Sánchez-Martínez, A. Anastasopol and E. Goetheer, *Ind. Eng. Chem. Res.*, 2021, **60**, 4269–4278.
- 33 Y. Matsuzaki, H. Yamada, F. A. Chowdhury, S. Yamamoto and K. Goto, *Ind. Eng. Chem. Res.*, 2019, **58**, 3549–3554.
- 34 H. M. Stowe and G. S. Hwang, *Phys. Chem. Chem. Phys.*, 2017, **19**, 32116–32124.
- 35 J. J. Lv, M. Jouny, W. Luc, W. Zhu, J. J. Zhu and F. Jiao, *Adv. Mater.*, 2018, **30**, e1803111.
- 36 J. Zhang, W. Luo and A. Züttel, *J. Mater. Chem. A*, 2019, **7**, 26285–26292.
- 37 E. J. Dufek, T. E. Lister and M. E. McIlwain, *Electrochem. Solid-State Lett.*, 2012, **15**, B48.
- 38 Y. Xu, J. P. Edwards, S. Liu, R. K. Miao, J. E. Huang, C. M. Gabardo, C. P. O'Brien, J. Li, E. H. Sargent and D. Sinton, *ACS Energy Lett.*, 2021, **6**, 809–815.
- 39 P. Jeanty, C. Scherer, E. Magori, K. Wiesner-Fleischer, O. Hinrichsen and M. Fleischer, *J. CO₂ Util.*, 2018, **24**, 454–462.
- 40 C. Chen, Y. Li and P. Yang, *Joule*, 2021, **5**, 737–742.
- 41 C. M. Gabardo, C. P. O'Brien, J. P. Edwards, C. McCallum, Y. Xu, C.-T. Dinh, J. Li, E. H. Sargent and D. Sinton, *Joule*, 2019, **3**, 2777–2791.
- 42 S. Popovic, M. Smiljanic, P. Jovanovic, J. Vavra, R. Buonsanti and N. Hodnik, *Angew Chem. Int. Ed. Engl.*, 2020, **59**, 14736–14746.
- 43 F. Li, Y. C. Li, Z. Wang, J. Li, D.-H. Nam, Y. Lum, M. Luo, X. Wang, A. Ozden, S.-F. Hung, B. Chen, Y. Wang, J. Wicks, Y. Xu, Y. Li, C. M. Gabardo, C.-T. Dinh, Y. Wang, T.-T. Zhuang, D. Sinton and E. H. Sargent, *Nat. Catal.*, 2019, **3**, 75–82.
- 44 E. S. Sanz-Perez, C. R. Murdock, S. A. Didas and C. W. Jones, *Chem. Rev.*, 2016, **116**, 11840–11876.
- 45 X. Min and M. W. Kanan, *J. Am. Chem. Soc.*, 2015, **137**, 4701–4708.
- 46 Y. Hori and S. Suzuki, *J. Electrochem. Soc.*, 1983, **130**, 2387.
- 47 T. Li, E. W. Lees, M. Goldman, D. A. Salvatore, D. M. Weekes and C. P. Berlinguette, *Joule*, 2019, **3**, 1487–1497.
- 48 E. W. Lees, J. C. Bui, D. Song, A. Z. Weber and C. P. Berlinguette, *ACS Energy Lett.*, 2022, **7**, 834–842.
- 49 Z. Zhang, E. W. Lees, F. Habibzadeh, D. A. Salvatore, S. Ren, G. L. Simpson, D. G. Wheeler, A. Liu and C. P. Berlinguette, *Energy Environ. Sci.*, 2022, **15**, 705–713.
- 50 Y. G. C. Li, G. Lee, T. G. Yuan, Y. Wang, D. H. Nam, Z. Y. Wang, F. P. G. de Arquer, Y. Lum, C. T. Dinh, O. Voznyy and E. H. Sargent, *ACS Energy Lett.*, 2019, **4**, 1427–1431.
- 51 A. Bonet Navarro, A. Nogalska and R. Garcia-Valls, *Electrochem*, 2021, **2**, 64–70.
- 52 T. Li, E. W. Lees, Z. Zhang and C. P. Berlinguette, *ACS Energy Lett.*, 2020, **5**, 2624–2630.
- 53 E. W. Lees, A. Liu, J. C. Bui, S. Ren, A. Z. Weber and C. P. Berlinguette, *ACS Energy Lett.*, 2022, **7**, 1712–1718.
- 54 J. Lee, H. Liu and W. Li, *ChemSusChem*, 2022, **15**, e202201329.
- 55 G. Lee, A. S. Rasouli, B.-H. Lee, J. Zhang, D. H. Won, Y. C. Xiao, J. P. Edwards, M. G. Lee, E. D. Jung,



- F. Arabyarmohammadi, H. Liu, I. Grigioni, J. Abed, T. Alkayyali, S. Liu, K. Xie, R. K. Miao, S. Park, R. Dorakhan, Y. Zhao, C. P. O'Brien, Z. Chen, D. Sinton and E. Sargent, *Joule*, 2023, **7**, 1277–1288.
- 56 L. A. Diaz, N. Gao, B. Adhikari, T. E. Lister, E. J. Dufek and A. D. Wilson, *Green Chem.*, 2018, **20**, 620–626.
- 57 S. Hosseini, H. Moghaddas, S. Masoudi Soltani, M. K. Aroua, S. Kheawhom and R. Yusoff, *J. Environ. Chem. Eng.*, 2018, **6**, 6335–6343.
- 58 A. G. Fink, E. W. Lees, Z. Zhang, S. Ren, R. S. Delima and C. P. Berlinguette, *ChemElectroChem*, 2021, **8**, 2094–2100.
- 59 E. W. Lees, M. Goldman, A. G. Fink, D. J. Dvorak, D. A. Salvatore, Z. Zhang, N. W. X. Loo and C. P. Berlinguette, *ACS Energy Lett.*, 2020, **5**, 2165–2173.
- 60 M. D. Gálvez-Vázquez, P. Moreno-García, H. Xu, Y. H. Hou, H. F. Hu, I. Z. Montiel, A. V. Rudnev, S. Alinejad, V. Grozovski, B. J. Wiley, M. Arenz and P. Broekmann, *ACS Catal.*, 2020, **10**, 13096–13108.
- 61 S. Verma, Y. Hamasaki, C. Kim, W. Huang, S. Lu, H.-R. M. Jhong, A. A. Gewirth, T. Fujigaya, N. Nakashima and P. J. A. Kenis, *ACS Energy Lett.*, 2017, **3**, 193–198.
- 62 M. E. Leonard, L. E. Clarke, A. Forner-Cuenca, S. M. Brown and F. R. Brushett, *ChemSusChem*, 2020, **13**, 400–411.
- 63 S. Banerjee, X. Han and V. S. Thoi, *ACS Catal.*, 2019, **9**, 5631–5637.
- 64 O. Gutiérrez-Sánchez, N. Daems, W. Offermans, Y. Y. Birdja, M. Bulut, D. Pant and T. Breugelmans, *J. CO₂ Util.*, 2021, **48**, 101521.
- 65 T. Al-Attas, S. NabilK, A. S. Zeraati, H. S. Shiran, T. Alkayyali, M. Zargartalebi, T. Tran, N. N. Marei, M. A. Al Bari, H. Lin, S. Roy, P. M. Ajayan, D. Sinton, G. Shimizu and M. G. Kibria, *ACS Energy Lett.*, 2022, **8**, 107–115.
- 66 C. Shen, R. Wycisk and P. N. Pintauro, *Energy Environ. Sci.*, 2017, **10**, 1435–1442.

