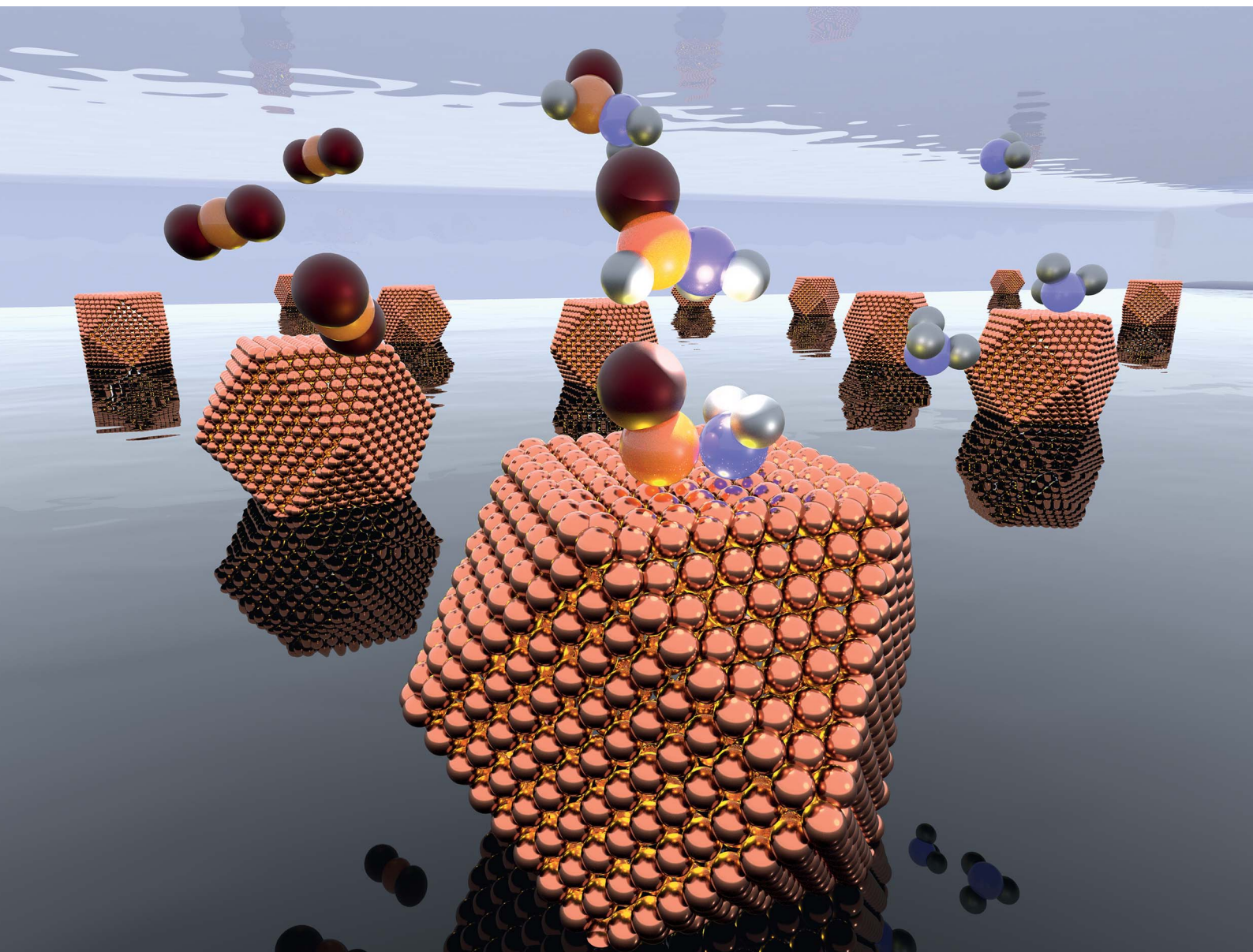


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# Electrochemically driven C–N bond formation from CO<sub>2</sub> and ammonia at the triple-phase boundary†

Junnan Li and Nikolay Kornienko \*

Electrosynthetic techniques are gaining prominence across the fields of chemistry, engineering and energy science. However, most works within the direction of synthetic heterogeneous electrocatalysis focus on water electrolysis and CO<sub>2</sub> reduction. In this work, we moved to expand the scope of small molecule electrosynthesis by developing a synthetic scheme which couples CO<sub>2</sub> and NH<sub>3</sub> at a gas–liquid–solid boundary to produce species with C–N bonds. Specifically, by bringing in CO<sub>2</sub> from the gas phase and NH<sub>3</sub> from the liquid phase together over solid copper catalysts, we have succeeded in forming formamide and acetamide products for the first time from these reactants. In a subsequent complementary step, we have combined electrochemical analysis and a newly developed *operando* spectroelectrochemical method, capable of probing the aforementioned gas–liquid–solid boundary, to extract an initial level of mechanistic analysis regarding the reaction pathways of these reactions and the current system's limitations. We believe that the development and understanding of this set of reaction pathways will play significant role in expanding the community's understanding of on-surface electrosynthetic reactions as well as push this set of inherently sustainable technologies towards widespread applicability.

## Introduction

With the increased focus on attaining global sustainability as a means to mitigate climate change and environmental degradation, the development of green technologies to enable the transition is increasingly important. Within this context, renewable electricity-powered electrosynthetic routes towards generating the fuels and chemicals that drive modern society stand to play a significant role if they manage to displace currently used fossil-fuel dependent methods.<sup>1–4</sup> While the recent decade of academic research has largely focused on water electrolysis<sup>5</sup> and CO<sub>2</sub> reduction<sup>6</sup> to generate H<sub>2</sub> and carbon-based fuels, respectively, there is no reason that the scope of heterogeneous electrosynthesis needs to be limited to these reactions. In principle, almost any commodity chemical can be synthesized from abundant small molecule building blocks (N<sub>2</sub>, H<sub>2</sub>O, CH<sub>4</sub>, biomass...) if the proper catalytic system would be developed.<sup>7</sup> The difficulty in realizing this ambitious aim is that at this point, only relatively simple electrosynthetic reactions over heterogeneous catalysts are well-understood and can be carried out at high rates and selectivity.

To this end, we moved to develop electrosynthetic routes to C–N bond formation using CO<sub>2</sub> and NH<sub>3</sub> as model building blocks. In general, despite the biological, societal and

technological importance of many chemicals containing C–N bonds,<sup>8–11</sup> the area of electrochemical C–N bond formation is very nascent. While biological<sup>12–14</sup> and chemical<sup>8,15</sup> routes are established, only few examples exist in carrying out C–N coupling on heterogeneous electrocatalysts. Thus, new reaction schemes and mechanistic insights in this context stand to provide a significant boost to the community.<sup>16–22</sup> In the context of heterogeneous catalysis, urea has previously been synthesized from co-electrolysis of N<sub>2</sub> or nitrate together with CO<sub>2</sub>.<sup>17,19,20,22</sup>  $\alpha$ -Keto acids have been converted into amino acids with hydroxylamine as a N-source.<sup>23</sup> Further, CO was co-electrolyzed with a series of different amines to generate amide products.<sup>18</sup> Finally, a host biomass-derived furans were reductively aminated to produce to amine derivatives.<sup>21</sup> To

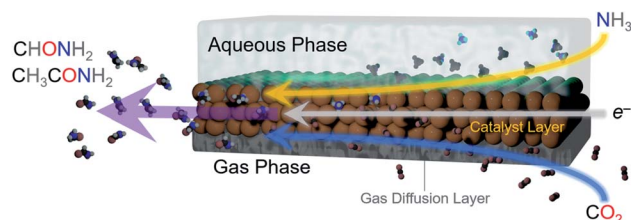


Fig. 1 Illustration of electrosynthetic strategy for on-surface C–N bond formation. A gas diffusion electrode was employed in which the reactants were simultaneously brought in from the gas phase (CO<sub>2</sub>) and from the liquid phase (NH<sub>3</sub>) and reacted over a solid Cu catalyst onto which an electrochemical potential was applied. This configuration enabled the generation of formamide and acetamide C–N bond containing products.

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expand the scope of possibilities of heterogeneously catalyzed C–N bond formation, we have developed a novel electro-synthetic scheme. Here,  $\text{NH}_3$  from the liquid phase would react with  $\text{CO}_2$  from the gas phase over a heterogeneous Cu catalyst at a gas–liquid–solid boundary (Fig. 1). As such, we generated formamide and acetamide from  $\text{CO}_2$  and  $\text{NH}_3$  for the first time, opening up a new avenue to the research community. Through quantitative reaction analysis and newly-developed infrared spectroelectrochemical investigations, we have built up a set of mechanistic insights in terms of elucidating reaction pathways and performance limitations, thus enabling the rational design of next-generation electrosynthetic systems.

## Catalyst construction

As a starting point, we selected two types of commercially available copper catalysts, Cu and CuO nanoparticles, for use in our study. Copper was selected as the element of choice because it has an intermediate binding energy to many carbon-based species.<sup>24</sup> This is a favorable metric in  $\text{CO}_2$  reduction because it enables the retention of surface intermediates en route to the formation of highly reduced products like ethylene. At the same

time, the binding strength to the intermediates is not too high to poison the surface. Thus, we reasoned that the same argument would apply in retaining  $\text{CO}_2$  reduction intermediates long enough for their coupling with  $\text{NH}_3$  would hold. While there is a plethora of studies of Cu-based catalysts and how defects, surface crystallographic facets, ligands, oxygen species and more dictate reaction pathways, we chose to leave such catalyst modifications for future follow-up works given the novelty of this reaction path.<sup>24</sup> The one variable that we did choose to investigate was the use of CuO as a starting material, which, when reduced to Cu under cathodic potentials, would likely contain additional binding sites in the form of defects. As such, Cu (Fig. 2a) and CuO (Fig. 2b) with no deliberate surface or structural modifications and size around 100 nm were used. The catalysts were mixed with a Nafion binder to generate an ink which was then drop cast onto a gas diffusion electrode. This type of electrode featured a gas-permeable gas diffusion layer and microporous layer through which  $\text{CO}_2$  could reach the catalyst layer on top (Fig. 2c). The goal here was to drive the C–N coupling reaction at this interfacial gas–liquid–solid boundary. This type of electrode geometry is particularly beneficial in overcoming the limited solubility of  $\text{CO}_2$  in aqueous electrolyte through the use of alkaline electrolytes

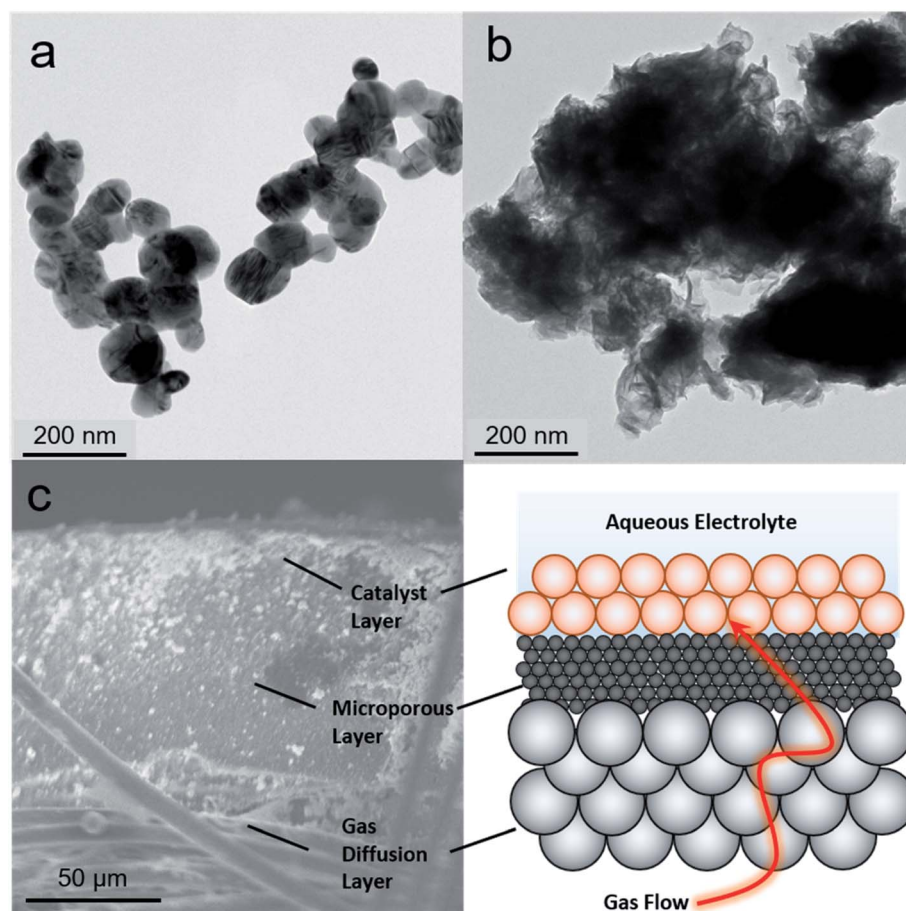


Fig. 2 The Cu (a) and CuO (b) catalyst particles were first characterized through transmission electron microscopy to probe their size and morphology. They were drop-cast onto a gas-diffusion electrode, which was characterized through scanning electron microscopy (c). The gas diffusion electrode consisted of several layers, illustrated with the graphic as a simplified representation. This electrode enabled gaseous reactants ( $\text{CO}_2$  in this case) to reach a solid electrocatalyst (Cu/CuO) and circumvent the limited solubility of  $\text{CO}_2$ .



that minimize the competing hydrogen evolution reaction at industrially relevant current densities (hundreds of  $\text{mA cm}^{-2}$ ).<sup>25,26</sup> The reaction cell was a modified one from those commonly employed in the field in order to minimize reaction volume and consequently maximize sensitivity for products. In particular, we employed an open cell design in which approximately 1 mL volume of electrolyte was used (Fig. S2†).

## Electrosynthetic studies

We employed 1 M KOH as an electrolyte for this work as highly alkaline electrolytes tend to minimize the hydrogen evolution reaction (HER) and thus favor  $\text{CO}_2$  reduction.  $\text{NH}_3$  was set to be a model nitrogen source. In the long term,  $\text{NH}_3$  would ideally be

replaced directly by  $\text{N}_2$  as an abundant feedstock, though at this stage, electrochemically activating  $\text{N}_2$  not yet a well-established reaction.<sup>27,28</sup> Formamide (Fig. 3a) and acetamide (Fig. 3b) primary amines were two likely C–N coupled products that could be formed from  $\text{NH}_3$  and  $\text{C}_1$  and  $\text{C}_2$  surface intermediates *via* several proton and electron transfer steps. In a gas-diffusion based electrochemical cell, both Cu and CuO featured an onset of catalytic current around 0 V vs. the reversible hydrogen electrode (RHE) and reached  $100 \text{ mA cm}^{-2}$  by  $-1.0 \text{ V}_{\text{RHE}}$  (Fig. S3a and b†). The addition of  $\text{NH}_4\text{OH}$  (present as mainly  $\text{NH}_3$  in alkaline solutions) to the electrolyte did not significantly alter the current density. Product quantification with gas chromatography (GC) and NMR revealed formate and  $\text{H}_2$  to be the two main products from the reaction (Fig. S4†). However, on

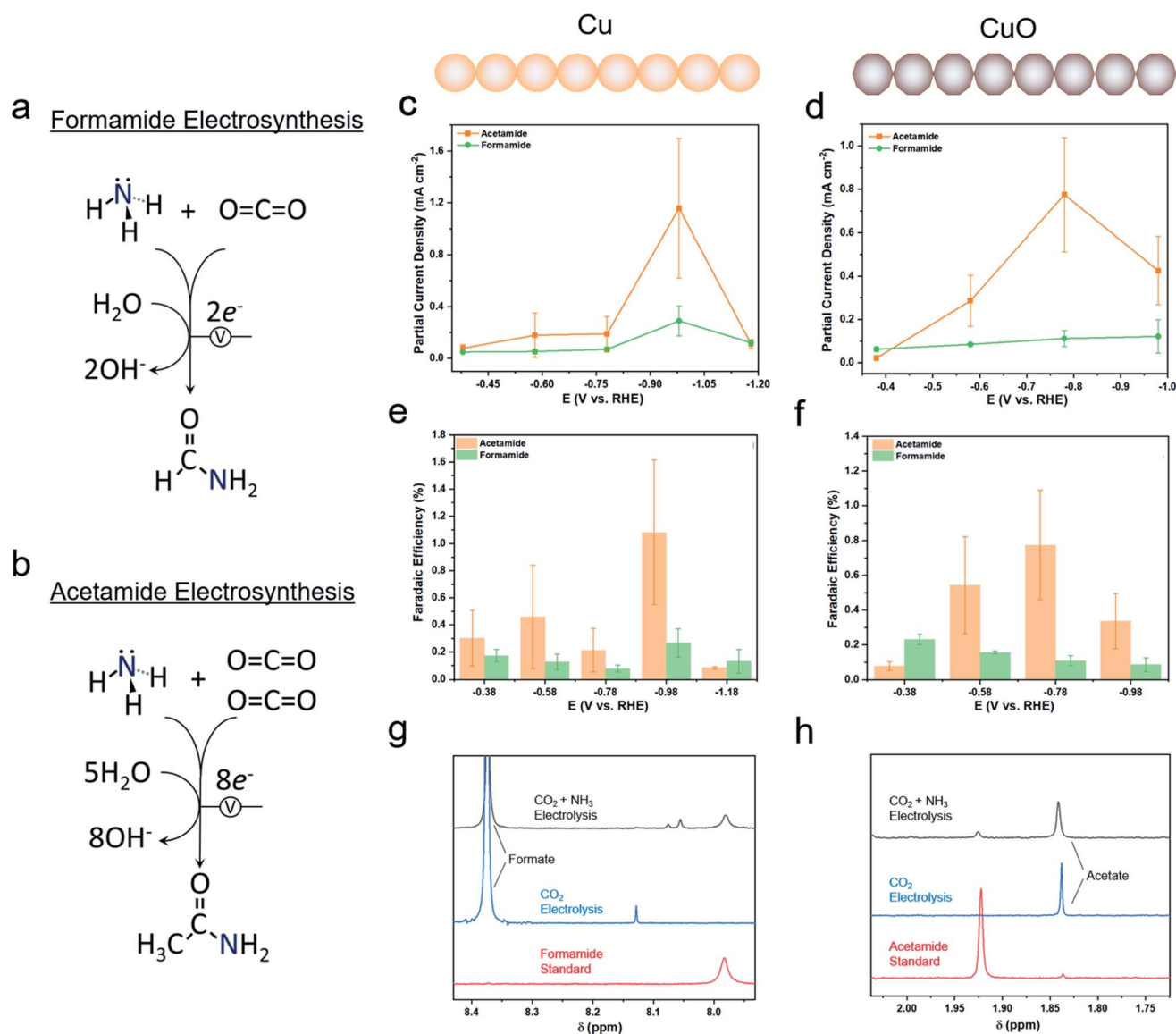


Fig. 3 The overall reaction is depicted for formamide involves 2 electrons and 1  $\text{CO}_2$  molecule (a) while acetamide electrosynthesis entails 8 electrons and 2  $\text{CO}_2$  molecules (b). In the gas diffusion electrode cell with a 1 M KOH electrolyte, 6SCCM  $\text{CO}_2$  flow, and the optimized concentration of  $\text{NH}_3$ , formamide (c) and acetamide (d) were quantified and their partial current densities derived from the resulting concentrations. The faradaic efficiencies for both products were similarly obtained for Cu (e) and CuO (f) catalysts. Representative NMR spectra are shown for formamide (g) and acetamide (h) from which the concentrations are quantified.



both Cu and CuO, formamide and acetamide were detected and were formed with partial current densities of ranging from 0.1 to 1.2 mA cm<sup>-2</sup>, depending on the applied potential (Fig. 3c and d). While the faradaic efficiency for their formation was rather modest, peaking at approximately 1% (Fig. 3e and f), this study constitutes the first report of their synthesis from CO<sub>2</sub> and NH<sub>3</sub> building blocks. In addition, performing the same measurements in a standard 3-electrode setup with the working electrode completely immersed in the aqueous phase did not result in any detectable C–N products, even after 24 h of electrolysis. This is likely due to a lower CO<sub>2</sub> concentration and lack of an alkaline environment that together promote a high degree of strongly bound C-based intermediates needed to couple with NH<sub>3</sub>. As a control experiment, CO<sub>2</sub> electrolysis alone only resulted in formate (Fig. 3g) and acetate (Fig. 3h) products that gave rise to NMR peaks in the range of interest. Interestingly, while the formate selectivity was very high (up to 90%) without NH<sub>3</sub>, NH<sub>3</sub> addition to the electrolyte decreased this value by a factor of 2–3 (Fig. S5†) while promoting hydrogen evolution. While this performance is not yet sufficient for economically competitive electrosynthesis, improving the initial system should certainly be feasible as one could point to the rapid maturation of CO<sub>2</sub> electrosynthetic technologies over the last decade.<sup>2</sup>

## Infrared spectroscopy

To extract a further level of insights into the formamide and acetamide electrosynthetic pathways, we turned to infrared

spectroscopy.<sup>29</sup> This technique measures the vibrational modes of species within the electrolyte and on the catalyst surface. Typically, measurements are carried out in difference mode, using the system at open circuit as a reference and subtracting this from the spectra under applied bias, thus detecting the appearance of new species (positive bands) and disappearance of others (negative bands). The spectroscopic measurements were carried out in an attenuated-total reflection (ATR) mode using a home-built spectroelectrochemical setup (Fig. 4a). Briefly, a thin layer of aqueous electrolyte (KOH or KOH + NH<sub>3</sub>) was on top of the diamond-coated ZnSe ATR crystal. The Cu catalyst layer/microporous layer/gas diffusion layer composite electrode was placed overtop so that the gas/liquid/solid boundary could be spectroscopically probed. In this configuration, the liquid was in static mode while the gas flowed above. The ability to probe this region was evident when measuring the difference spectrum between Ar flow and CO<sub>2</sub> flow in this configuration, which shows the presence of both gaseous and dissolved CO<sub>2</sub> and carbonate species (Fig. S12†).<sup>30</sup>

Under an argon flow with NH<sub>3</sub> and no CO<sub>2</sub>, the main spectral features corresponded mainly to that of water and to that of NH<sub>3</sub> (Fig. S13–S15†).<sup>30</sup> Under the same conditions but with CO<sub>2</sub> flowing in place of Ar, a new set of positive bands appeared which can primarily be assigned to carbonate and bicarbonate species coming from CO<sub>2</sub> reacting with the KOH electrolyte and changes in pH (Fig. S14†).<sup>31</sup> Such species been shown to spontaneously appear at the gas–liquid–solid interface in similar conditions with Raman measurements.<sup>32</sup> While spectral features in the 1800–2100 cm<sup>-1</sup> are noted where the C–O stretch

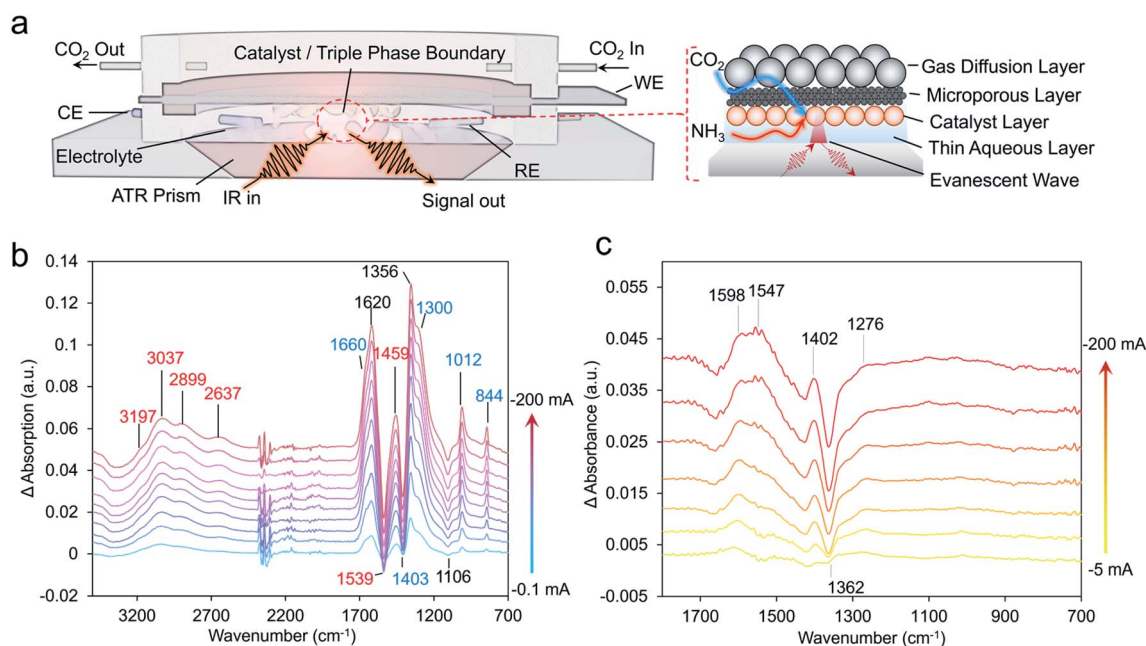


Fig. 4 Spectroelectrochemical setup enabling *operando* infrared spectroscopic probing of the electrosynthetic reactions in a gas-diffusion electrode cell (a). This setup used a thin electrolyte window with the GDE just overtop to probe both liquid, gas and solid phases. The gas was flowing through while the liquid was static. Using the spectrum at open circuit as the background, spectra under select operating current densities with CO<sub>2</sub> and NH<sub>3</sub> present were recorded (b). Subtracting out the bi(carbonate) contributions using the spectra at 1 mA cm<sup>-2</sup> as a background, enables the identification of additional species present on the CuO surface (c).



of  $^*CO$  is located, the inherent absorbance of our diamond-coated ATR crystal makes this region rather noisy rendering bands here more difficult to fit and explicitly assign.

In the presence of both  $CO_2$  and  $NH_3$ , new bands appeared in the region containing N–H bonds from the generation of  $NH_4^+$  (Fig. 4b).<sup>33</sup> As a method of validation, spectra were also acquired with  $^{15}NH_3$  instead of  $^{14}NH_3$  (Fig. S13†). Indeed, the isotope effect was noted *via* a red-shift around  $30\text{ cm}^{-1}$  for bands at both spectral regions. Interestingly, the intensity of the bicarbonate band at  $1300\text{ cm}^{-1}$  saturated very early with only  $CO_2$  present, but continually gained intensity under increasingly higher currents when  $NH_3$  was present (Fig. S13†). A possible explanation for this could be the presence of  $NH_3$  diminishes the concentration of  $CO_2$ /carbonate reactants near the interface at low currents.

Finally, as (bi)carbonate species dominate the IR spectra, we opted to subtract spectra of the catalyst systems operating at  $-1\text{ mA}$  from those at  $-200\text{ mA}$ , as the (bi)carbonate species are mostly saturated and those with smaller spectral contributions could be visualized (Fig. 4c). Indeed, for the CuO catalyst in the presence of  $CO_2$  and  $NH_3$ , the evolution of positive spectral features ( $1645$ ,  $1598$ ,  $1547$ ,  $1402$  and  $1096\text{ cm}^{-1}$ ) and negative bands ( $1660$ ,  $1425$  and  $1362\text{ cm}^{-1}$ ) were noted. While a fully unambiguous assignment at this stage is not yet possible, we note that these spectral features correlate well with those previously assigned to  $^*COO^-$  and  $^*COOH$  and these species are thus our tentative assignments.<sup>34–36</sup> The complete set of plausible band assignments is compiled in Table 1.

As formate is the dominant product in each of these systems, it would seem reasonable to have a substantial  $^*COO^-$  surface coverage and thus this is our tentative assignment. While the precise mechanism of formate electrosynthesis is still under debate, on copper surfaces, it has been argued through a combination of surface-enhanced Raman spectroscopy and DFT modelling that the all  $CO_2$  reduction pathways share a common first intermediate in a  $\mu_2$ ,  $-C$ ,  $-O$  bound  $CO_2^{*}$  that subsequently gets hydrogenated en route to formate or

protonated to  $^*COOH$  en route to  $CO$  and other  $C_2$  downstream products.<sup>37</sup> Thus, the observation of  $^*COO^-$  and  $^*COOH$  intermediates being the main ones on the surface would support our product distribution.

While this study used  $NH_3$  as a starting point for generating C–N bond containing products, ideally, the nitrogen source would be gaseous  $N_2$  as the technology matures and scales. To this end, we have explored as an intermediate step the co-reduction of nitrate and nitrite ions in place of  $NH_3$  to generate the same products. The experimental procedure was the same except that the nitrate/nitrite was in place of  $NH_3$  in the electrolyte, with optimized concentrations of  $NO_2/NO_3$  (Fig. S15 and S16†). Over a Cu catalyst at  $-0.98\text{ V vs. RHE}$ , formamide and acetamide were indeed formed, albeit at reduced faradaic efficiencies and partial current densities (Fig. 5). While a comprehensive set of electroanalytical and spectroscopic experiments is outside of the scope of this initial work, the results indicate that there is much to discover in optimizing the reduction pathways of both C and N sources en route to C–N bond formation. A likely reaction pathway that would explain our results would be the reduction of nitrate/nitrite on the electrode surface to  $NH_3$  or  $^*NH_2$ , which then couples with intermediates from  $CO_2$  reduction.

When using  $NH_3$  as the starting point we believe that it is not directly bound to the surface but rather located in the near-surface region (Fig. S18†). Its presence seems to hinder both hydrogen evolution and  $CO_2$  reduction, likely through inhibiting reactant diffusion to active sites on the catalyst. The tendency to enhance hydrogen evolution likely stems through an enhanced hindrance of  $CO_2$  diffusion as opposed to that of water molecules.

Finally, we moved to optimize the C–N product generation of our system. We first screened the parameters of KOH concentration, Cu catalyst loading, and cation identity (Fig. S21†). The biggest enhancement in acetamide generation came from a higher Cu loading ( $10\text{ mg cm}^{-2}$  vs.  $2\text{ mg cm}^{-2}$ ). We attribute this to the propensity of the catalyst layer to generate a higher amount of highly reduced  $C_2$  intermediates through the thicker Cu film (Fig. 6a and b). Further, a lower KOH concentration and a change from  $K^+$  to  $Cs^+$  also yielded selectivity enhancements for acetamide, possible due to a more favorable near-surface reaction environment to stabilize reaction intermediates en route to the  $C_2$  species that can couple with  $NH_3$ .<sup>39</sup> Interestingly, the activity enhancements were not realized for formamide. This can be rationalized as the factors necessary for favorizing a  $CO_2$  reduction pathway to  $C_2$  species as not being necessary for formamide, which is more dependent on the initial  $^*COO^-$  intermediate coupling with  $NH_3$ .

Then, with all three optimized parameters incorporated ( $0.1\text{ M CsOH}$  electrolyte,  $10\text{ mg cm}^{-1}$  Cu loading), we compared the faradaic efficiency and partial current densities for the optimized system to that of the original from Fig. 2 across the experimental potential range (Fig. 6c).

We believe that the formate and formamide electrosynthetic pathways are linked on the Cu surface in that they share a common intermediate. This belief is backed by their similarity in chemical structure. Considering that the formation of C–N

**Table 1** Peaks detected and plausible assignments from infrared experiments

Band position ( $\text{cm}^{-1}$ )	Species	Figure	Possible assignment
1660	$HCO_3^-$	4b	$\nu_{as}(-C-O)^{30}$
1356	$HCO_3^-$	4b	$\nu(-C-O)^{30}$
1300	$HCO_3^-$	4b	$\delta(C-OH)^{30}$
1012	$HCO_3^-$	4b	$\nu_{as}(C-OH)^{30}$
844	$HCO_3^-$	4b	$\nu_s(C-OH)^{38}$
1459	$CO_3^{2-}$	4b	$\delta(-NH)^{30}$
3197	$NH_4^+$	4b	$\nu(N-H)^{33}$
3037	$NH_4^+$	4b	$\nu(N-H)^{33}$
2899	$NH_4^+$	4b	$\nu(N-H)^{33}$
1459	$NH_4^+$	4b	$\nu(N-H)^{33}$
1645	$H_2O$	4c	$\delta(H-O-H)$
1598	$^*COO^-$	4c	$\nu_{as}(COO^-)^{37}$
1547	$^*COO^-$	4c	$\nu_{as}(COO^-)^{35}$
1402	$^*COO^-$	4c	$\nu_s(COO^-)^{35,36}$
1276	$^*COOH$	4c	$\nu COOH, OH\text{-deformation}^{35}$



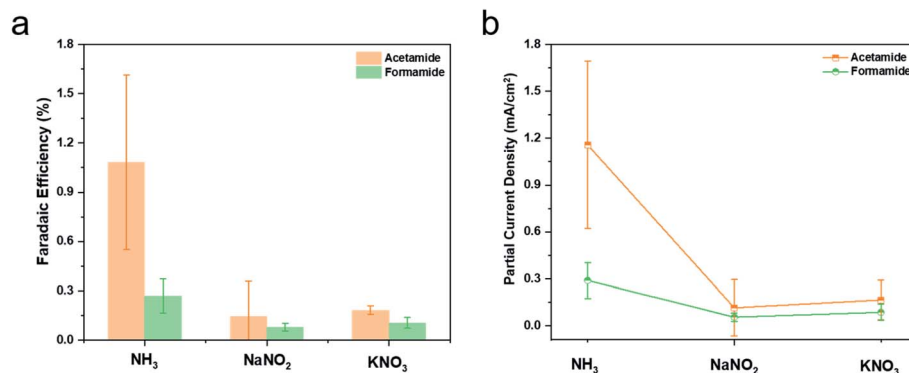


Fig. 5 In the equivalent reaction setup as in Fig. 3 with Cu catalysts and NO<sub>2</sub><sup>-</sup> (0.5 M) or NO<sub>3</sub><sup>-</sup> (1 M) in place of NH<sub>3</sub> at -0.98 V vs. RHE, formamide and acetamide were generated. The faradaic efficiencies (a) and partial current densities (b) for their generation were compared to those from using NH<sub>3</sub>.

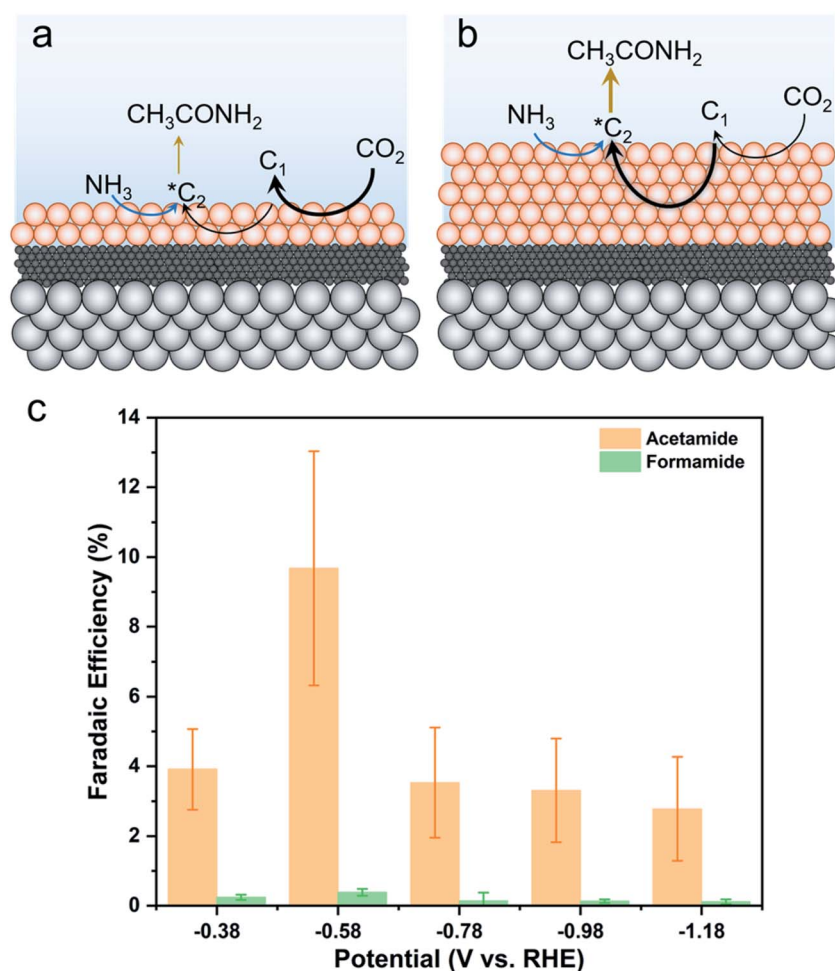


Fig. 6 A thicker catalyst loading was found to promote acetamide formation. Thin layers tend to form C<sub>1</sub> products at a greater rate (a) while increasing the layer thickness leads to further reduction and accumulation of C<sub>2</sub> intermediates that can be used for C–N bond formation (b). As such, an optimized Cu loading of 10 mg cm<sup>-2</sup> resulted in up to 10% acetamide selectivity (c).

containing products involves the nucleophilic attack of a carbon atom by the lone pair on the nitrogen atom of ammonia, an activated, yet exposed carbon species that could couple with

ammonia for formamide generation could be that of the  $\mu_2$ , -C, -O bound \*CO<sub>2</sub><sup>-</sup>.<sup>37</sup> There would then be a competition between hydrogenation of this species to produce formate or



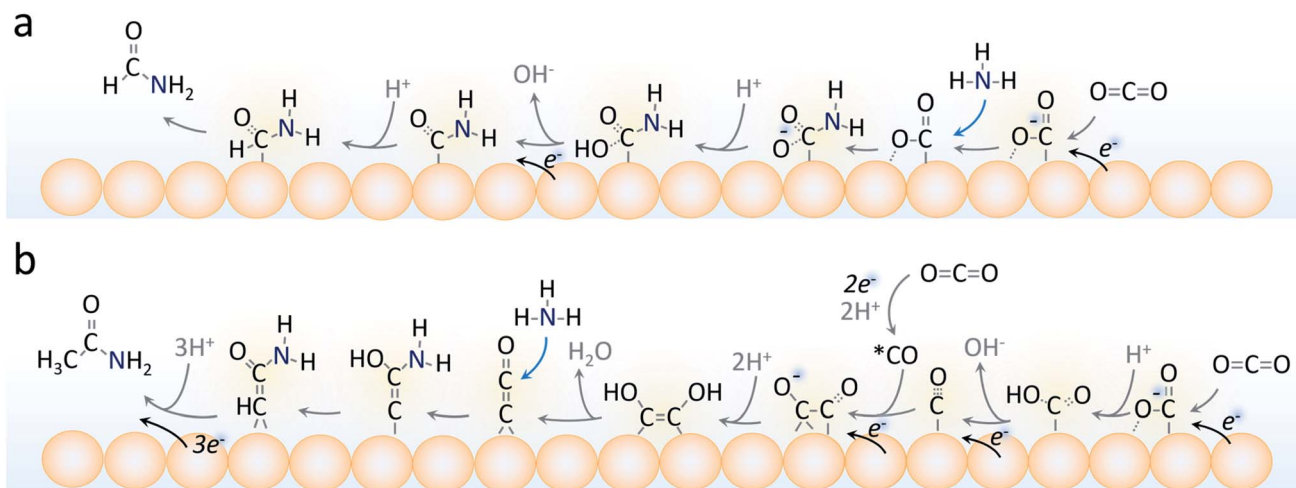


Fig. 7 Plausible surface reaction pathways in the electrochemical process of formamide (a) and acetamide (b) generation. Formamide generation depends on NH<sub>3</sub> coupling with the first reaction intermediate while acetamide generation requires highly reduced C<sub>2</sub> intermediates to be present on the catalyst surface. For simplicity, the donation of a proton to surface intermediates via H<sub>2</sub>O → OH<sup>-</sup> is depicted as →H<sup>+</sup>.

a nucleophilic attack to eventually form formamide (Fig. 7a). This branching point is also supported by the fact that using 1 M formate instead of CO<sub>2</sub> as the C-source in otherwise identical conditions did not lead to any detectable formamide and only a small amount of acetamide (Fig. S14†).

On the other hand, acetamide synthesis likely shares a reaction pathway with acetate and thus requires a C<sub>2</sub> intermediate to already be present.<sup>18</sup> The branching for this step also plausible occurs at the  $\text{*CO}_2^-$  intermediate, which is converted to  $\text{*CO}$ . The coupling of 2  $\text{*CO}$  molecules is thought to be a key step to generating C<sub>2</sub> products (acetate, ethanol, ethylene). The  $\text{*CCO}$  intermediate was recently proposed as a likely candidate for this through a DFT analysis of acetamide synthesis *via* CO and NH<sub>3</sub> building blocks and would be a plausible candidate for our work as well.<sup>18</sup> The middle carbon would thus be subject to nucleophilic attack by the NH<sub>3</sub> in this pathway where it diverges from the acetate pathway as previously postulated.<sup>18</sup> The enhancement of C<sub>2</sub> intermediates such as this with a thick catalyst layer is postulated to be the main driving factor for its generation.

The two pathways presented here are not so different than what occurs in enzymatic catalytic pockets, where an electron rich amine couples with an electron poor carbon<sup>12–14</sup> and one can imagine that generating on-surface catalytic pockets in a synthetic system to promote this reaction would lead to further enhancements of electrochemical selectivity. As the nucleophilic attack is by the ammonia nitrogen is a thermodynamically downhill step, reduction potential for both of these reactions is still dictated by that necessary to reduce CO<sub>2</sub>.

## Concluding remarks

While two new reaction pathways have been discovered in formamide and acetamide electrochemical synthesis using CO<sub>2</sub> and NH<sub>3</sub> building blocks, many avenues are now opened for further understanding and improving the efficiency of these reactions.

First, while we used commercially purchased Cu and CuO nanoparticles as a readily available model system, they feature a diversity of active sites, defects, (sub)surface oxygen species, and exposed crystallographic facets. It is entirely possible that each of these factors may influence the reaction like they do in the electrochemical synthesis of carbon-based products *via* CO<sub>2</sub> reduction. A rational way forward would be the precise study of well-defined copper catalysts in which the nature surface-active sites are known and with complementary theoretical modelling of likely reaction pathways on these surfaces. Further, it is not known if Cu is the only catalyst capable of carrying out this reaction and if formate-selective metals like Sn and Bi would thus be more effective at formamide synthesis. In addition, we have developed an *operando* infrared spectroscopic method for the first time that was used to help understand this reaction pathway but additional complementary techniques such as Raman and X-ray absorption would contribute immensely valuable pieces to this puzzle.<sup>40</sup>

This principal significance to this work is the electrochemical reaction discovery which we envision will accelerate the adoption of this methodology at large in both the academic and industrial domains. While NH<sub>3</sub> is used as the model nitrogen source and nitrate and nitrite as further examples, eventually, this may be replaced by N<sub>2</sub> in a fully sustainable nitrogen cycle. In general, the capacity to drive heteroatomic surface coupling reactions with renewable-electricity powered systems stands to open up an abundance of decentralized green synthetic routes in place of heavy-infrastructure requiring fossil fuel based thermochemical approaches. In parallel, there is much more fundamental chemistry to be discovered through the use of new interfaces, spectroscopic methodology, and catalytic systems.

## Data availability

Experimental data available upon reasonable request.





## Author contributions

N. K. and J. L. both designed the project, carried out experiments, processed data, contributed intellectual insights and wrote the manuscript.

## Conflicts of interest

None to declare.

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