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## Solvent-mediated outer-sphere CO<sub>2</sub> electro-reduction mechanism over the Ag111 surface†

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The electrocatalytic CO<sub>2</sub> reduction reaction (CO<sub>2</sub>RR) is one of the key technologies of the clean energy economy. Molecular-level understanding of the CO<sub>2</sub>RR process is instrumental for the better design of electrodes operable at low overpotentials with high current density. The catalytic mechanism underlying the turnover and selectivity of the CO<sub>2</sub>RR is modulated by the nature of the electrocatalyst, as well as the electrolyte liquid, and its ionic components that form the electrical double layer (EDL). Herein we demonstrate the critical non-innocent role of the EDL for the activation and conversion of CO<sub>2</sub> at a high cathodic bias for electrocatalytic conversion over a silver surface as a representative low-cost model cathode. By using a multiscale modeling approach we demonstrate that under such conditions a dense EDL is formed, which hinders the diffusion of CO<sub>2</sub> towards the Ag111 electrocatalyst surface. By combining DFT calculations and *ab initio* molecular dynamics simulations we identify favorable pathways for CO<sub>2</sub> reduction directly over the EDL without the need for adsorption to the catalyst surface. The dense EDL promotes homogeneous phase reduction of CO<sub>2</sub> *via* electron transfer from the surface to the electrolyte. Such an outer-sphere mechanism favors the formation of formate as the CO<sub>2</sub>RR product. The formate can undergo dehydration to CO *via* a transition state stabilized by solvated alkali cations in the EDL.

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## Introduction

Electrochemical conversion of CO<sub>2</sub> holds promise to help mitigate the carbon footprint of the production of fuels and chemicals.<sup>1</sup> The abundant CO<sub>2</sub> green-house waste gas is an attractive substrate to stabilize excess “electrons” generated from renewable energy *via* the CO<sub>2</sub> reduction reaction (CO<sub>2</sub>RR).<sup>2</sup> A wide range of electrocatalysts have been described so far for the CO<sub>2</sub>RR.<sup>3,4</sup> Depending on the catalyst employed the primary CO<sub>2</sub>RR product can be either formate or CO resulting from a 2e<sup>-</sup> reduction, or multi-electron transfer products such as alcohols and hydrocarbons. The electrocatalytic reduction of CO<sub>2</sub> to CO opens a path for carbon recycling within the established syngas chemistry infrastructure to produce fuels and chemicals.<sup>1</sup>

In an electrocatalytic cell, the CO<sub>2</sub>RR proceeds on the cathode side. The cathode material, its morphology and electrolyte properties collectively influence the electrocatalytic activity and selectivity at the solid–liquid interface.<sup>5,6</sup> Gold-based electrocatalysts have been reported to reduce CO<sub>2</sub> to CO with high activity and selectivity. Hori and co-workers used bulk Au to reduce CO<sub>2</sub> to CO with 87.1% faradaic efficiency (FE) at

–1.14 V (NHE) with a partial CO current of 5 mA cm<sup>-2</sup>.<sup>7</sup> At the same partial current, Ag showed a FE of 81.5% towards CO at –1.37 V (NHE).<sup>7</sup> The lower cost, and comparable selectivity and activity to Au make Ag an attractive electrocatalyst for the CO<sub>2</sub>RR.

The selectivity and activity of an electrocatalyst for the CO<sub>2</sub>RR are strongly influenced by the electrolyte and the local environment close to the cathode.<sup>5,6,8–11</sup> At potentials ( $\Phi_M$ ) below the potential of zero charge (pzc) ( $\Phi_{pzc}$ ) the negative charge density on the cathode surface increases, attracting more cations and resulting in the formation of an electrical double layer (EDL). The EDL influences the local electrochemical environment close to the cathode surface such as the interfacial pH and the structure of water at the interface.<sup>11–13</sup> The cations in the EDL also interact with the surface intermediates and tune the stabilization of transition states and adsorbates on the electrocatalyst surface.<sup>14,15</sup> However, at a higher cathodic bias, the EDL becomes very dense and compact, thus strongly hampering the mass transport of CO<sub>2</sub> to the electrocatalyst surface.<sup>16–18</sup>

The electron transfer (ET) from the cathode to the reagent is the key mechanistic step of any electrocatalytic conversion. ET can in principle proceed *via* two alternative mechanisms: inner- and outer-sphere ET (Fig. 1). The inner-sphere mechanism starts with the chemisorption of the reagent (CO<sub>2</sub>) to the catalyst surface that enables direct ET *via* overlapping orbitals (Fig. 1a). In the outer-sphere mechanism indirect ET from the catalyst

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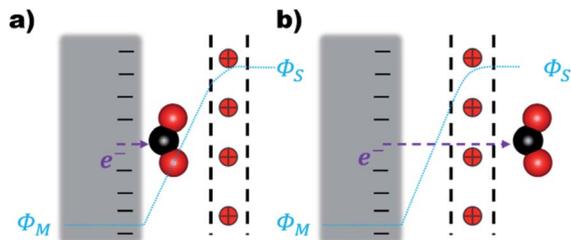


Fig. 1 (a) Inner-sphere and (b) outer-sphere electron transfer from the cathode to  $\text{CO}_2$ . The red positive charges within the black dotted lines denote the EDL. ( $\Phi_M$ ) and ( $\Phi_S$ ) are the surface and bulk solution phase potentials respectively.

surface to  $\text{CO}_2$  takes place through the electrolyte without the direct chemical interaction between the reagent and the electrocatalyst (Fig. 1b).

The inner-sphere mechanisms describing the electrocatalytic conversions in the framework of surface adsorbed species dominate the current literature.<sup>5,19–25</sup> Investigating the selectivity of the  $\text{CO}_2\text{RR}$  on various Ag facets, Bohra and co-workers proposed that the formation of formate species is self-inhibited on Ag surfaces resulting in improved selectivity to CO at low to moderate potentials, and to  $\text{H}_2$  at higher potentials.<sup>26</sup> Their work did not consider the effects from the EDL and the electrolyte explicitly. A realistic description of the reaction medium and conditions in modelling studies has been currently emphasized across the field of catalysis.<sup>27–40</sup> The importance of including an explicit representation of the EDL and accounting for the reaction conditions in mechanistic studies of the electrocatalytic  $\text{CO}_2\text{RR}$  has been emphasized in recent literature.<sup>14,15,41,42</sup>

However, most mechanistic studies assume facile mass transport of  $\text{CO}_2$  from the bulk phase to the surface *via* the EDL. Such an assumption is reasonable for hydrodynamic transport through a low concentration electrolyte but under the operando  $\text{CO}_2\text{RR}$  potentials, the EDL can get more condensed and strongly impact the mass transport of  $\text{CO}_2$ .<sup>16,17</sup> Under such conditions an outer-sphere ET in the homogeneous phase over the EDL is a plausible mechanism for the  $\text{CO}_2\text{RR}$ .

Herein, we have taken a multiscale operando modelling approach to investigate the possibility and the impact of homogeneous ET on the  $\text{CO}_2\text{RR}$  under realistic electrocatalytic conditions. The combination of classical molecular dynamics (CMD) and *ab initio* molecular dynamics (AIMD) simulations shows that  $\text{CO}_2$  can be favourably reduced to formate anions *via* outer-sphere ET over the dense EDL. The formate species can then convert to CO *via* a thermally activated dehydration reaction facilitated by the solvated cations within the EDL.

## Results and discussion

### Molecular structure of the EDL

To rationally construct an atomistic operando model of the cathode-electrolyte interface under the reaction conditions, the formation and structure of the EDL at the Ag111 surface was first investigated by classical molecular dynamics (CMD)

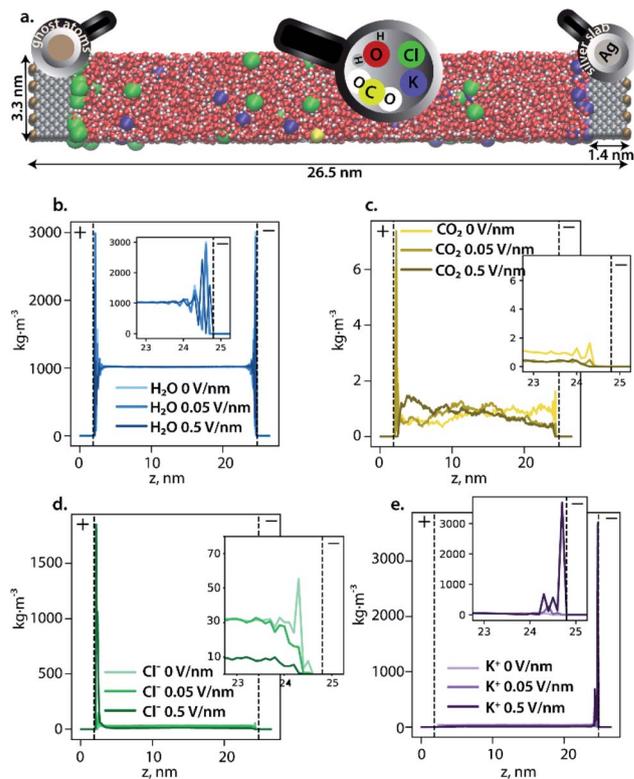


Fig. 2 (a) A snapshot of the model of the KCl electrolyte with  $\text{CO}_2$  confined between two silver slabs simulated at  $0.5 \text{ V nm}^{-1}$ . The surface on the left represents the anode, with positive ghost charges imposed behind the wall, while the right surface represents the cathode, with negative ghost charges imposed behind the wall. The color code is as follows: silver is grey, oxygen is red, hydrogen is white, potassium is violet, chlorine is green, carbon of  $\text{CO}_2$  is yellow, and the ghost atoms are light brown. (b–e) Density profile of  $\text{H}_2\text{O}$  (b),  $\text{CO}_2$  (c),  $\text{Cl}^-$  (d), and  $\text{K}^+$  (e), under increasing polarization conditions represented by the electric field at the surface of the electrode. At low or zero polarization most of the ions are present in the bulk phase. With increasing polarization, the respective densities of  $\text{K}^+$  at the cathode and  $\text{Cl}^-$  at the anode show a sharp increase leading to the formation of compact EDLs at the respective electrodes.

simulations. The electrocatalytic system was modelled as an aqueous electrolyte containing  $0.86 \text{ M KCl}$  and  $0.06 \text{ M CO}_2$ , confined between two Ag111 slabs (the model cathode and the anode) in a super cell of dimensions  $33.1 \times 37.2 \times 265.5 \text{ \AA}^3$  periodic in the  $x$  and  $y$  directions (see Fig. 2a). Complete details of the CMD model and simulations are presented in the ESI.† These simulations aimed to probe the formation of the EDL at the electrodes under different polarization conditions. The polarization conditions were mimicked by placing uniform distributions of point charges behind the Ag111 slabs resulting in negative (cathode) and positive (anode) surface charge densities on the electrolyte facing surfaces.

The CMD simulations of the extended electrocatalyst system representing the electrochemical cell revealed the formation of a dense EDL at the cathode as the polarization was increased (Fig. 2). The density of water oscillates within  $1 \text{ nm}$  of the cathode surface indicating the formation of ordered layers of solvation, while it was found to be constant at  $1 \text{ kg m}^{-3}$  in the



bulk phase. The simulations show a deeper penetration of water molecules into the outer Helmholtz plane (OHP; indicated by the dashed line at 1 Å from the electrode) of the cathode compared to the anode.  $K^+$  ions accumulated near the cathode while  $Cl^-$  anions accumulated at the anode and their respective concentrations in the EDL region increased with increasing surface polarization. This resulted in the compaction of the EDL and associated depletion of the  $CO_2$  near the electrocatalyst surface. The latter is fully covered by solvated alkali cations, which can be further considered as the reactive sites for  $CO_2$  activation instead of the bare metal surface.

### Outer-sphere $CO_2RR$

The outer-sphere reduction of  $CO_2$  over the solvated Ag111 surface was next investigated using periodic density functional theory (DFT) calculations (see the ESI† for details†). The reactive events were simulated using a smaller molecular model representing the reaction environment near the Ag111 cathode ( $4 \times 4 \times 5$  slab model). The initial static DFT calculations on the simplified models revealed the critical role of the EDL in the outer-sphere charge transfer eliciting indirect reduction of  $CO_2$ . Indeed, the interaction of  $CO_2$  with an aqueous solvation layer

on Ag111 is very weak and does not lead to notable perturbations of the adsorbed molecules ( $Ag111-(H_2O)_{24}-CO_2$ , Fig. 3a). The situation drastically changes upon the introduction of sodium ions and the formation of the EDL ( $Ag111-(Na^+)_4-(H_2O)_{24}-CO_2$ , Fig. 3b), which facilitates the reduction of  $CO_2$ . The  $CO_2$  molecule in this case adopts a bent configuration due to the partial charge transfer from the silver slab. The bent anionic  $CO_2$  moiety is stabilized by hydrogen bonding with the neighbouring  $H_2O$  molecules.

To better investigate the outer-sphere ET and the subsequent conversions of  $CO_2$  over the EDL, an extended fully solvated model was employed containing ( $Ag111-4Na^+-(H_2O)_{61}-CO_2$ , Fig. 2c) in combination with AIMD simulations (BLYP-PW-400 eV, VASP 5.5.4, for further details see the ESI†). The reactive environment was simulated with a 19 ps long AIMD simulation of  $CO_2$  in the solvated phase over the Ag111-EDL interface.  $CO_2$  was found to preferentially stay in the 3<sup>rd</sup> and 4<sup>th</sup> water layers of solvation at about 10 Å from the Ag111 surface. The average  $O=C=O$  angle was  $\sim 172^\circ$  during the runs. Two of the four  $Na^+$  cations forming the EDL were found at about  $\sim 3$  Å from the surface while the other two  $Na^+$  cations were located further away at  $\sim 5$  Å on the Ag111 surface (Fig. 3c).<sup>43</sup> The water molecules within the EDL close to the surface facet were found to show a limited mobility. They pointed their protons towards the metal surface during the simulations. In the absence of the EDL, the water molecules preferentially oriented with O moieties pointing towards the Ag111 surface (Fig. 2a). Next, constrained AIMD simulations were carried out on 11 intermediate states representing different stages of the outer-sphere ET  $CO_2RR$ . The  $O=C=O$  angle was chosen as the reaction coordinate ( $Q$ ), and it was varied from  $172^\circ$  to  $125^\circ$ . The resulting Gibbs free energy profile along with the representative snapshots of the relevant reactant and product configurations are presented in Fig. 4.

The constrained AIMD simulations revealed that upon bending, the  $CO_2$  moiety diffused closer to the EDL. The transition state was located between  $O=C=O$  angles of  $140^\circ$ – $145^\circ$  (Fig. 4). Releasing the constraint at  $\angle O=C=O = 140^\circ$  directly

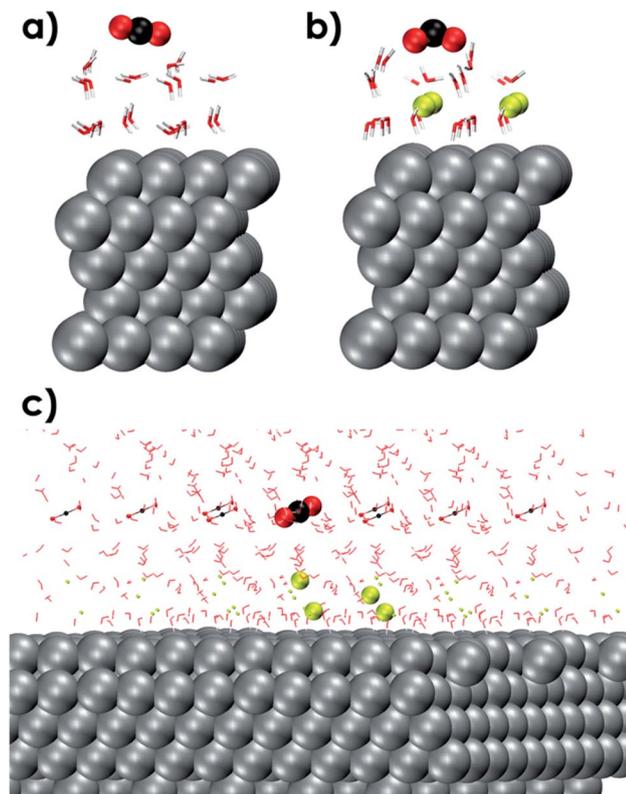


Fig. 3 DFT optimized molecular models of a  $4 \times 4 \times 5$  Ag111 slab with one  $CO_2$  molecule in vacuum over the (a) Ag111-water interface (b) Ag111-EDL interface EDL =  $(Na^+)_4(H_2O)_{24}$  and (c) fully solvated and periodic (in  $x$ ,  $y$  and  $z$  directions) system used for AIMD simulations composed of a  $4 \times 4 \times 5$  Ag111 slab,  $4Na^+$  cations and 61 water molecules. Periodic images have also been shown. The  $CO_2$  and  $Na^+$  species in the original simulation cell are shown as larger VdW spheres. Color code: C (black), Ag (silver), O (red), Na (Yellow), and H (white).

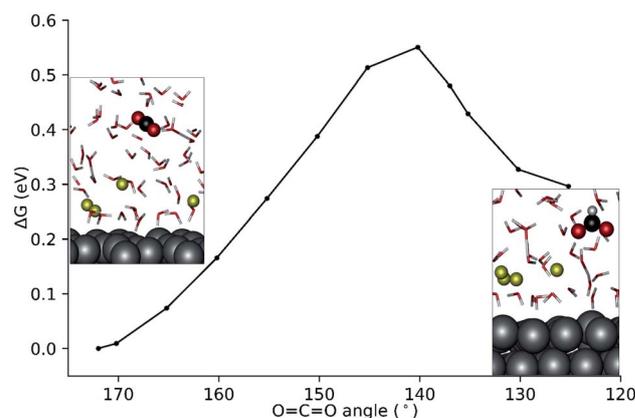


Fig. 4 Computed Gibbs free energy profile for the homogeneous reduction of  $CO_2$  to formate using the  $\angle O=C=O$  angle as the reaction coordinate.



results in the formation of the formate product. The activation free energy barrier for the outer-sphere  $\text{CO}_2\text{RR}$  is 0.55 eV with reference to the linear  $\text{CO}_2$  molecule.

Bader charge analysis at  $Q = 140^\circ$  (equilibrated supercell at  $\sim 18$  ps) revealed a net atomic charge of  $-0.80$  on the  $\text{CO}_2$  moiety. This is comparable to the Bader net atomic charge of  $-0.74$  units computed for the  $1e^-$  reduced  $\text{CO}_2$  radical in water (see the ESI†). Therefore, the bent  $\text{CO}_2$  moiety at  $140^\circ$  represents a  $1e^-$  reduced  $\text{CO}_2$  radical. Upon further bending the transient radical species accepts an  $\text{H}^+$  from the solvent simultaneously with the second ET to yield the formate product. A snapshot of the proton transfer at  $\angle \text{O}=\text{C}=\text{O} = 137^\circ$  is shown in the ESI.† Three water molecules coordinated to a  $\text{Na}^+$  in the EDL are actively involved during the proton transfer *via* H-bonding interactions. The bent  $\text{CO}_2$  moiety is strongly solvated forming 5 H-bonds.

The interaction with the EDL is critical for the reduction of  $\text{CO}_2$  and can be compared with cation mediated outer-sphere ET among species in the homogeneous phase (Fig. 5a).<sup>44</sup> In the homogeneous phase solvated alkali cations have been reported to mediate outer-sphere ET between two species.<sup>44</sup> AIMD simulations show that the solvated cations in the EDL can facilitate a similar outer-sphere ET between the cathode surface and  $\text{CO}_2$  (Fig. 5b and c). The critical role of the EDL in facilitating the  $\text{CO}_2\text{RR}$  was further highlighted by the slow-growth

approach (SGA) simulations that slowly bent the  $\text{CO}_2$  moiety when it was located far away from the EDL. In the absence of interaction of  $\text{CO}_2$  with the EDL, bending the  $\text{CO}_2$  moiety resulted in the formation of the  $\text{HCO}_3^-$  species by the nucleophilic attack of water suggesting the importance of the EDL in facilitating electroreduction (see the ESI†).

The formate species formed *via* the outer sphere ET mechanism can undergo dehydration to form CO, which is expected to be favoured by low pH conditions near the EDL.<sup>45</sup> Dehydration of formic acid/formate in acidic conditions is well-established chemistry. Consistently, AIMD simulations revealed similar free energy barriers for the dehydration of formate to CO in the solvated phase in the presence of the EDL (Ag111-4 $\text{Na}^+$ -( $\text{H}_2\text{O}$ )<sub>61</sub>- $\text{CO}_2$  system; 1.19 eV) and without the EDL (Ag111-( $\text{H}_2\text{O}$ )<sub>61</sub>- $\text{CO}_2$  system; 1.26 eV) (see the ESI†). The presence of the EDL provides a small stabilization (0.07 eV) to the dehydration TS. The dehydration of formate to CO is expected to be more favourable with larger cations such as  $\text{K}^+$  and  $\text{Cs}^+$  where the pH near the EDL is lower.<sup>45</sup>

Experimental results for the  $\text{CO}_2\text{RR}$  over Ag show that at high cathodic bias the hydrogen evolution reaction outcompetes the formation of CO. Jaramillo and co-workers showed that the partial current density for the formation of  $\text{H}_2$  exceeds that for CO at cathodic potentials below  $-1.3$  V (RHE).<sup>46</sup> The rate of formation of CO peaks around  $-1.1$  V (RHE) and then decreases as the potential is lowered. Further analysis revealed that the decrease in CO formation was due to mass transport limitations. Both formate and CO require  $\text{CO}_2$  to reach the cathode surface for the  $\text{CO}_2\text{RR}$  to proceed *via* an inner-sphere ET mechanism. Therefore, the rate of formate production is also expected to decrease around the same potential where CO production dips due to mass transport limitations. Contrastingly, the partial current density of formate, although always lower than CO and  $\text{H}_2$  kept growing as the cathodic bias was decreased.

An outer-sphere ET mechanism, which does not require mass transport of  $\text{CO}_2$  to the surface, can explain the increasing partial formate current density. At moderate to low cathodic bias,  $\text{CO}_2$  can reach the surface, and its adsorption is stabilized by the EDL<sup>14,15</sup> leading to the production of CO (kinetically favoured) and  $\text{HCOO}^-$  (less favoured). Thus, the current density for CO and  $\text{HCOO}^-$  both increase as the applied voltage is lowered. At high cathodic bias the current density switches from kinetic control to mass transport limitations leading to decreased CO production. We suggest that the mass transport limitations (at least partially) result from a condensed EDL rather than only solubility and diffusion of  $\text{CO}_2$  in the electrolyte.  $\text{CO}_2$  is therefore available in the region close to the EDL and gets reduced to formate *via* an outer-sphere ET mechanism, which explains the increasing formate current density.

To gain further insight into  $\text{HCOO}^-$  *versus* CO production we compare the free energy barriers for CO formation (inner-sphere ET) reported in the literature *versus*  $\text{HCOO}^-$  formation (outer-sphere ET) as computed by us. Based on the results reported by Chen and co-workers for the  $\text{CO}_2\text{RR}$  over the Ag111 surface in the presence of a model EDL, a free energy barrier of 0.52 eV can be estimated for surface-mediated CO formation at

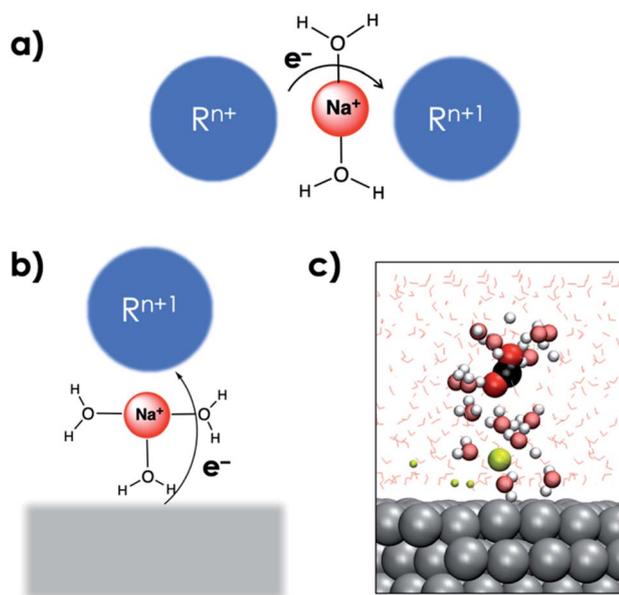
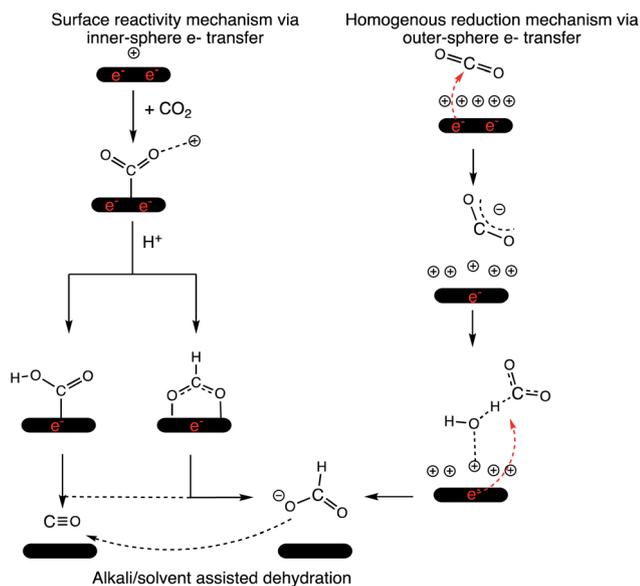


Fig. 5 (a) Schematic representation of alkali cation promoted outer-sphere ET between two species in the homogeneous phase. (b) Proposed schematic representation of alkali cation mediated outer-sphere ET from a cathode surface to a species in the homogeneous phase. (c) Snapshot of an AIMD trajectory with the  $\text{O}=\text{C}=\text{O}$  angle constrained at  $140^\circ$  showing the interaction between a solvated cation in the EDL and the solvated  $\text{CO}_2$  moiety analogous to the schematic depiction in (b). VdW representation is used for the  $\text{CO}_2$  moiety interacting with a solvated  $\text{Na}^+$  cation along with its first solvation shells which are shown in brushed metal colors. Other  $\text{Na}^+$  cations are shown as smaller spheres, and water molecules are shown *via* line representation. Periodic images have also been included.





**Scheme 1** Mechanism of the CO<sub>2</sub>RR over a cathode surface *via* inner- and outer-sphere electron transfer. 2e<sup>-</sup> that originate from the cathode surface and reduce the CO<sub>2</sub> moiety are shown in red for representative purposes. The cathode surface is maintained at a constant potential in the electrolyzer.

an applied external potential of  $-1$  V (SHE) at pH = 7.<sup>14</sup> The current AIMD-computed free energy barrier of 0.55 eV at  $-1.45$  V (SHE) ( $\sim 1$  V *versus* the PZC)<sup>47</sup> reflects a relatively higher barrier for the CO<sub>2</sub>RR *via* the outer sphere ET, explaining the lower partial current density for formate.

Therefore, we propose that the mechanism of 2e<sup>-</sup> reduction of CO<sub>2</sub> is dependent on the applied bias. Scheme 1 summarizes the outer- and inner-sphere 2e<sup>-</sup> mechanisms to produce CO and formate *via* the CO<sub>2</sub>RR. Jaramillo and co-workers also demonstrated the formation of >2e<sup>-</sup> reduction products of the CO<sub>2</sub>RR over Ag at high overpotentials<sup>46</sup> and an outer-sphere ET could potentially be involved in those mechanistic steps as well.

## Conclusions

We have explored the CO<sub>2</sub>RR over the Ag111 surface *via* an outer-sphere ET mechanism. Following a multiscale operando modelling strategy, we first simulated the multi-component electrolyte–cathode interface under various applied potentials. Investigation of density profiles of water, ions and CO<sub>2</sub> revealed the formation of a condensed EDL within 1 nm of the cathode surface composed of cations and ordered layers of solvation at high overpotentials. This finding motivated the development of a smaller periodic model of the cathode-electrolyte interface which was used to investigate the reactive events during the CO<sub>2</sub>RR at high cathodic potentials *via* AIMD simulations. AIMD simulations showed that an outer-sphere ET mechanism resulted in the formation of formate species over the EDL. The formate species was further shown to undergo alkali promoted dehydration to CO with a moderate free energy barrier of 1.19 eV. The presence of the EDL was found to be the key to promote an outer-sphere ET CO<sub>2</sub>RR mechanism.

The outer-sphere ET CO<sub>2</sub>RR is a plausible mechanism to produce formate and CO under high cathodic bias. A surface-based, alkali promoted CO<sub>2</sub>RR is likely still the dominant mechanism for the formation of CO. Our calculations show that an alternative reaction channel to reduce CO<sub>2</sub> is accessible in the presence of a dense EDL, and the reaction mechanism is a complex network of voltage dependent inner- and outer-sphere ET steps. Outer-sphere mechanisms should be further explored for heterogeneous electrocatalytic systems and can be especially relevant for electrocatalytic reduction of organic substrates which occur under high voltage conditions.

## Data availability

Data related to this publication is available *via* the 4TU database under the DOI: 10.4121/19142303.

## Author contributions

EAP conceived and supervised the project. VS carried out DFT and AIMD simulations. EK performed CMD simulations. All the authors discussed the results and wrote the manuscript.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

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