Fundamental Reaction Limitations for Formic Acid Oxidation

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ABSTRACT. Electro catalytic conversion of formic acid oxidation to CO₂ and the related CO₂ reduction to formic acid represent a potential closed carbon-loop based on renewable energy. However, formic acid fuel cells are inhibited by the formation of poisoning species during the reaction. Recent studies have elucidated how the binding of carbon and hydrogen on catalyst
surfaces promote CO\textsubscript{2} reduction towards CO and formic acid. This has also given fundamental insights to the reverse reaction, \textit{i.e.} the oxidation of formic acid. In this work, simulations on multiple materials have been combined with formic acid oxidation experiments on electrocatalysts to shed light on the reaction and the accompanying catalytic limitations. We found that: \textit{(i)} The desired principal reaction for efficient formic acid oxidation should progress through adsorbed carboxyl, "COOH, which should then be oxidized to CO\textsubscript{2}. \textit{(ii)} "H adsorbed on the surface results in "CO formation and poisoning through a chemical disproportionation step. \textit{(iii)} Catalysts are poisoned by formate and/or hydroxyl. Using these results, a framework for the reaction has been developed explaining the fundamental limitations and progressing our understanding.

**Introduction**

Tremendous efforts are currently going into out-phasing fossil fuels in favor of sustainable fuels.\textsuperscript{1} This is motivated by our need to close the carbon cycle\textsuperscript{2} as well as the decrease in renewable electricity prices, which simultaneously have paved the way for new fuel production routes.\textsuperscript{3} Electrocatalytic technologies propose direct electrification of chemical and fuel production. Examples include reduction of CO\textsubscript{2} towards HCOOH, CH\textsubscript{3}OH, C\textsubscript{2}H\textsubscript{5}OH and H\textsubscript{2}O towards H\textsubscript{2}.\textsuperscript{1} Efficient fuel consumption through fuel cells (FCs) also holds great potential.\textsuperscript{4} Liquid fuels such as formic acid and methanol have attracted a lot of attention due to their: Viable energy density \textit{per} mass- and volume, attractive handling/storage properties and potential uses in other non-fuel applications, \textit{e.g.} as a high value chemical building blocks for industry.\textsuperscript{3}

Some liquid fuels such as methanol, are notoriously limited in the oxidation toward CO\textsubscript{2} since the process goes through a CO intermediate.\textsuperscript{5} CO oxidation then becomes the limiting factor determining the performance of direct methanol FCs (DMFCs). Formic acid as liquid fuel behaves
differently; it has a CO$_2$-like structure with two hydrogens attached. This molecular structure predicates that the oxidation process only requires the removal of two hydrogen atoms. Consequently, formic acid oxidation should ideally circumvent the problem of CO-poisoning.

To gauge formic acid’s efficiency as a fuel we compare the single round trip efficiency of relevant closed-loop chemical compounds, i.e. hydrogen, formic acid, methanol and lithium batteries as seen in Table 1. Here we observe that the Li-battery storing and release of energy exhibits the highest efficiency followed by hydrogen. However, both Li-batteries and H$_2$ suffer from low energy density. Storing energy as methanol and formic acid is very similar in terms of the cost in electrolyzer energy. The major difference between formic acid and methanol arises when using the chemical in a fuel cell, where methanol is limited by CO oxidation.

Table 1. Estimated round-trip efficiencies ($\eta = \frac{U_{OBR} - U_{Fuel cell reaction}}{U_{OER} + U_{Electrolyzer reaction}} \times 100\%$) calculated using the difference in energy potentials. Common for hydrogen, formic acid, and methanol we use $U_{OER} = 1.6\ V_{RHE}$ and $U_{OBR} = 0.8\ V_{RHE}$. Here only the cost of syngas production is considered, not the full formation of methanol. **Typical charge/discharge efficiency.

<table>
<thead>
<tr>
<th></th>
<th>Energy stored (electrolyzer)</th>
<th>Energy released (fuel cell)</th>
<th>Round trip efficiency, (\eta)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hydrogen</td>
<td>$U_{HER} = -0.1\ V_{RHE}$</td>
<td>$U_{HOR} = 0.1\ V_{RHE}$</td>
<td>$\approx 41%$</td>
</tr>
<tr>
<td>Formic acid</td>
<td>$U_{CO_2\rightarrow HCOOH} = -0.8\ V_{RHE}$</td>
<td>$U_{FAOR} = 0.2\ V_{RHE}$</td>
<td>$\approx 25%$</td>
</tr>
<tr>
<td>Methanol*</td>
<td>$U_{CO_2\rightarrow CO + H_2} = -0.6\ V_{RHE}$</td>
<td>$U_{CO\rightarrow CO_2} = 0.65\ V_{RHE}$</td>
<td>$\approx 7%$</td>
</tr>
<tr>
<td>Li-battery**</td>
<td></td>
<td></td>
<td>$\approx 90%$</td>
</tr>
</tbody>
</table>
Depending on the $U_{FAOR}$, formic acid fuel cells can be considered an attractive alternative to methanol fuel cells. Methanol provides 6 protons per reacted molecule and formic acid only two and therefore methanol has a ~3 times higher volumetric energy density. However, the potential of a formic acid fuel cell is high; even a few hundred millivolts reduction in overpotential can allow formic acid to output energy than methanol per molecule.

Key to the formic acid oxidation is the direct link to the reverse electrochemical reaction, i.e. the CO$_2$ reduction reaction (CO$_2$RR). Combining these two reactions allows for a closed carbon-loop with formic acid working as a liquid energy storage media. CO$_2$RR to formic acid and the formic acid oxidation reaction (FAOR) can be written in the form:

(CO$_2$RR) \[ \text{CO}_2 + 2\text{H}^+ + 2\text{e}^- \rightarrow \text{HCOOH} \quad \Delta G_{\text{CO}_2 \rightarrow \text{HCOOH}}^0 = 0.12 \text{ eV per } \text{H}^+ + \text{e}^- \quad (1) \]

(FAOR) \[ \text{HCOOH} \rightarrow \text{CO}_2 + 2\text{H}^+ + 2\text{e}^- \quad \Delta G_{\text{FAOR}}^0 = -0.12 \text{ eV per } \text{H}^+ + \text{e}^- \quad (2) \]

Where $\Delta G^0$ is the thermodynamic potential per proton-electron pair of the reaction.

CO$_2$RR selectivity is highly dependent on the catalyst material used and the crystal orientation. $^{9-11}$ $^{12}$ Hori et al. showed that hydrogen is produced on Pt, Ru, Fe and Ni, carbon monoxide is produced on Au, Ag, Zn, Ga and Pd (which also produces similar amounts of H$_2$), and finally, formic acid is produced on Pb, In, Hg, Sn, Cd and Tl.$^{12}$ Importantly, hydrocarbons are uniquely produced on Cu.$^{12}$ Using simulations, we were able to classify the CO$_2$RR product distributions towards hydrogen, hydrocarbons, CO and formic acid due to the catalyst’s affinity towards adsorbed $^*$H and $^*$CO.$^{10}$ Interestingly, we hereto noted that the CO$_2$RR propensity towards formic acid generally appears to be more selective when weakening $^*$H binding, or in other words; CO is favored over HCOOH when catalyst binds $^*$H sufficiently strong without forming H$_2$. Moreover, from this study$^{10}$ we noted that the $^*$OOCH vs. the $^*$COOH intermediate cannot be used to
distinguish the CO or formic acid product formation. These findings are readily usable when considering the reverse reaction FAOR.

FAOR exhibits the highest intrinsic pure metal activity on Pt and Pd.\textsuperscript{8,13,14} However, the reaction is affected by high overpotentials and formation of various poisoning intermediates.\textsuperscript{15} The following observations can be made in the literature, as shown in Supporting information (SI) Figure S1: (i) FAOR onsets at low potentials does not necessarily correspond to high FAOR currents. (ii) Hysteresis between anodic and cathodic scans is a common occurrence. (iii) Pt(111) is more active than Pd(100) in the low overpotential region; however, interestingly, this relationship shifts at higher potentials. (iv) As an observation it is known that there is a difference for Pt and Pd with respect to the CO poisoning during FAOR.\textsuperscript{8} The ideal FAOR catalyst on the other hand should show reversible cyclic voltammetry (CV), high activity, low onset potential and stable currents, as illustrated in Figure S2.

To understand the mechanism of this important reaction numerous attempts have been made to map possible FAOR pathways,\textsuperscript{15–19} which we illustrate by an in-depth literature study in Figure 1. The motivation has often been the elucidation of the reaction mechanism along with the identification and circumvention of poisoning issues. Either through pathway engineering,\textsuperscript{16,17} changing electrolyte composition,\textsuperscript{18} or inclusion of sites with the ability to remove poisoning species.\textsuperscript{19–22} CO-poisoning from partial HCOOH oxidation is often considered the principal culprit\textsuperscript{20} and various works suggest CO formation can be avoided utilizing a single/dual-site catalyst.\textsuperscript{23–26} However, catalysts such as Au-Pt,\textsuperscript{16} Pt-Hg/C\textsuperscript{27} and Pd-Hg/C\textsuperscript{28} exhibit limited catalytic improvement over their pure metal counterparts, for some overview see Figure S4 in SI.
Figure 1. Literature study highlighting all the conceived FAOR reaction pathways during potential cycling. Historically, FAOR has been split into the direct (gray) and indirect (green and yellow) pathways. Further, partial FAOR and catalyst oxidation forming unwarranted surface blocking have been suggested, e.g. CO$_2$H species (purple), CO (red) and hydroxide/oxides (blue). Even CO$_2$RR induced CO formation by applying too cathodic potential (brown) have been suggested. Recently, formate (yellow) in various arrangements has gained attention as potential catalyst poisons.

In this work, we address the fundamental questions in FAOR:

(i) Does FAOR proceed through *COOH or *OOCH?

(ii) How is CO formed during FAOR?

(iii) Can *OOCH act as both a spectator and a poisoning species?

(iv) Why do only few catalysts work for FAOR?
While Figure 1 treats all the proposed FAOR reaction paths uncritically, Figure 2 summarizes the minimum number of considerations we believe is required to understand the FAOR process qualitatively.

**Figure 2.** Presentation of the minimum reaction elements required to understand FAOR activity and the poisoning events potentially occurring during CVs, based on Figure S3 in the SI.

The important aspects for FAOR highlighted in Figure 2 are *(i)* the principal reaction through \(^*\)COOH, *(ii)* the disproportionation reaction of \(^*\)COOH reacting with \(^*\)H to form CO, *(iii)* formate poisoning via \(^*\)OOCH and *(iv)* oxidation of the catalysts. Considerations of the reactions *(i)* and *(iv)* is of common occurrence in FAOR literature. While in this work, we consider and analyze *(ii)* and *(iii)* based on FAOR’s strong link to the CO\(_2\)RR.\(^{10}\) To the best of our knowledge, the disproportionation reaction *(ii)* is only anecdotally mentioned in the comprehensive work of Sun *et al.* on Pt.\(^{29}\) In contrast, it has been widely accepted in CO\(_2\)RR literature that protonating \(^*\)COOH leads to CO.\(^{43}\) For CO\(_2\)RR it has been a paradox that formate (\(^*\)OOCH) binds stronger than carboxyl (\(^*\)COOH). Hence, while carboxyl describes the onset potential for CO formation the
formate species does not correlate with any CO$_2$RR reaction activity and is consequently considered a CO$_2$RR poison.$^{44}$

To probe the scientific questions, we use a combination of experimental and simulation tools. For experiments we use cyclic voltammetry (CV) and chronoamperometric (CA). For simulations we use density functional theory (DFT) calculations on the binding of carboxyl, *COOH, formate bidentate, *OOCH, and hydrogen, *H.

**Results and Discussion**

![Graphs](image)

**Figure 3.** Electrochemical data recorded at room temperature of equal loading nanoparticle catalysts: Pt/C (*navy blue*), Pd/C (*crimson*), Pt-Hg/C (*blue*), Pd-Hg/C (*red*) and Pt-Bi/C (*green*) systems on glassy carbon in Ar-saturated 0.1 M HClO$_4$ with 0.1 M HCOOH at 1600 rpm, unless
otherwise stated. (a) FAOR CVs at 10 mV/s. (b) Base CVs in Ar-saturated 0.1 M HClO₄ at 400 rpm and 50 mV/s. (c) CA at 0.55 V\textsubscript{RHE} for 30 min. All measurements were repeated at least three times (shade represents standard deviation), IR-compensated and post-corrected, for experimental details see SI and Figure S5-S8.

Figure 3 summarizes experimental electrochemical data of Pt/C, Pd/C, Pt-Hg/C, Pd-Hg/C and Pt-Bi/C. Here we investigate only known and active FAOR catalyst, and both extended surfaces as Pt/C and Pd/C, but also single site catalyst Pt-Hg/C and Pd-Hg/C and the noteworthy very active Pt-Bi/C system. Figure 3a shows the formic acid oxidation CV during rotation and 10 mV/s scan-rate on the five catalysts. Figure 3b depicts the base CVs exhibiting suppressed hydrogen underpotential deposition (H\textsubscript{UPD}) on the Pt-Bi/C, Pt-Hg/C and Pd-Hg/C compared to the Pd/C and Pt/C counterparts. Figure 3c displays the formic acid oxidation CA at 0.55 V\textsubscript{RHE} for 30 min, illustrating the loss in activity at this potential due to the formation of poisoning or blocking species. For Pt/C and Pt-Hg/C, an apparent hysteresis is seen in the oxidation between the forward and backward scans of Figure 3a indicating an irreversible change in the catalyst going to low potentials. Interestingly, taking a combined view on Figure 3a,c shows that forming single-sites of Pt through Hg alloying\textsuperscript{27,28} tend to improve the FAOR onset. In contrast, Pd-based catalysts generally do not exhibit any hysteresis. The Pt-Bi/C system exhibit the highest FAOR current with least hysteresis, but also with the highest overpotential. Additional relevant electrochemical studies can be found in the SI, represented through Figures S10-S14. To make sense of all of these observations we turned to DFT.
Figure 4. DFT calculated binding energies for metal (111) fcc (light-blue crosses) and single-site catalyst. MNC-based (black points), Pt atom in Au(111) denoted Pt;Au(111) (cyan triangle), PdHg₄ (black/red square) and PtHg₄ (black/blue square). (a) *COOH vs. *H. (b) *OOCH vs. *H. (c) *OOCH vs. *COOH, here the dashed line shows the diagonal indication the affinity towards formate bound through carbon or oxygen. Here its assumed ΔECOOH ≈ ΔEHH. Further, we used CO₂ and H₂ for references when calculating *COOH, ΔECOOH is 0.12 eV per electron.

Figure 4a maps out the DFT calculated *COOH vs. *H binding energies of the most relevant metals and single-site catalysts, such as MNCs, PtHg₄, PdHg₄ and single Pt atoms in Au, Pt;Au(111). For metals and the single-site-catalysts linear scaling appears: ΔE∗COOH = ΔE∗H + b, where b is about 0.29 eV as previously observed for metals and 0.0 eV for single-site catalyst, respectively. Besides the scaling, a vertical- and a horizontal line indicating HUPD (ΔEHUPD) and formic acid’s thermodynamic equilibrium potential have been included in Figure 4a,b. Here its assumed that ΔECOORD ≈ ΔGCOORD, as thermodynamic corrections and water stabilization are expected to cancel out for the *COOH intermediate. The fundamental scaling between *COOH and *H, together with the potential adsorption of *H on the surface, leads to the possibility of disproportionation towards CO. This entails that crossing the scaling and the H_UPD line represents a fundamental limit for the binding of *COOH at 0.2 V_RHE for metals to avoid disproportionation.
Figure 4b displays the DFT calculated binding energies of *OOCH vs. *H. Conversely to Figure 4a, there is no apparent scaling between formate bidentate and adsorbed hydrogen (however the oxygen bond for *OOCH scales with *OH). Essentially, Figure 4b shows that the FAOR pathway cannot follow the *OOCH intermediate. Comparing the data of Figure 4b with literature one can observe that metal catalysts, such as Cu and Ag exhibiting a short distance to the equilibrium potential line - meaning that they should be active - are not active for the FAOR in experimental investigations (also shown in Figure S10). In short, it is unlikely that *OOCH is an important reaction intermediate. Instead, given that the *OOCH scales with *OH, the *OOCH binding can be considered a probe of the oxidation affinity of the catalyst, *i.e.* having a strong *OOCH binding results in a lower oxidation potential.

Figure 4c shows the *OOCH vs. *COOH binding for the catalyst, with a dashed line indicating the affinity towards carbon or oxygen bound formate. Depending on the catalyst we can see whether formate bidentate or carboxyl is favored. In this regard there is a small caveat, that *OOCH scales with *OH binding, as discussed above. Figure 4c further reveals that most pure metal catalyst, *e.g.* Ir46 or Au47, should perform poorly as FAOR catalyst in agreement with the literature. Only two outliers exist for this interpretation, that is Pd13 and Rh48. However, Pd *OOCH binding is too close to that of *COOH disallowing specific conclusions on that element. Moreover, consulting Rh’s Pourbaix diagram49 the divergence from our hypothesis is explained by the formation of conductive oxides forming in the FAOR relevant pH- and potential conditions.

The type of analysis illustrated in Figure 4 is an powerful tool able to identify which catalyst suffers from disproportionation and, by virtue of the scaling-relations,10 it gives fundamental insights into why literature historically has shown no significant FAOR activity below ~0.2 V_{RHE} for a metal catalysts. By trusting Figure 4a as a the principal design guide governing FAOR
activity, it even suggests that, to achieve FAOR activities below 0.2 \( V_{\text{RHE}} \), one needs to consider single-site-based catalyst as the \(^*\text{COOH} \ vs. \ ^*\text{H} \) scaling is shifted \textit{ca.} 0.29 eV weaker compared to metal-based catalysts. Implicitly Figure 4a states that hysteresis between anodic and cathodic sweeps in FAOR CVs arise due to this type of disproportionation occurring at low potential arising from the \( \text{H}_{\text{UPD}} \) and the consequent creation of \(^*\text{CO} \) poisoning species. From this insight, one would expect that disproportionation is mitigated in the CVs by simply staying above \( \text{H}_{\text{UPD}} \) potentials.

**Figure 5.** FAOR CVs at 10 mV/s, 1600 rpm and room temperature in Ar-saturated 0.1 M HClO\(_4\) with 0.1 M HCOOH at different potential limits. Ohmic drops were post-corrected. (a) Pt/C. (b) Pt-Hg/C. (c) Pd/C. Note, increasing lower potential limit minimizes the hysteresis in the CVs, lowering the upper potential limit generally decreases the activity.
Figure 5 shows the performance of Pt/C, Pt-Hg/C and Pd/C cycled with varying potential limits. The potential range from 0.00-1.05 V_{RHE} reveals that Pt/C and Pt-Hg/C are poisoned in anodic sweeps. Changing to potentials, ranging from 0.40-1.05 V_{RHE} for the Pt/C and 0.35-1.05 V_{RHE} for the Pt-Hg/C, significantly increases the anodic activity. Hence, this allows us to prove that H_{UPD} mediated disproportionation account for poisoning through CO on Pt catalysts. For Pd/C in Figure 5c, decreasing the lower potential limit has no influence on the almost non-existing FAOR hysteresis. In this context, it is important to note that Pd is well-known to form Pd-hydride phases below 0.2 V_{RHE}, i.e. at potentials relevant for both FAOR and CO_{2}RR. In relation to CO_{2}RR, we also note that Pd-hydride, leads to a high faradaic efficiency towards formate, whereas at higher overpotentials CO and H_{2} will dominate.

Concerning Figure 5, one could erroneously assume that lowering the upper potential limit would not affect the CVs, while staying above the CO oxidation potential. This is however not the case, cycling 50 times from 0.35-1.00 V_{RHE} reveals some form of deactivation of both Pt/C, Pt-Hg/C and Pd/C. We do not know what the origin of this deactivation is. Various studies reported in the literature suggests different reasons including deactivation to irreversible metal oxidation due to insufficient surface reduction or accumulation of either *OOC*, *COH*, *OCOH* or *CO* species. Most interesting is that in situ Fourier-transform infrared spectroscopy (FTIR) work has shown that Pt, contrary to Pd, continuously form CO above H_{UPD} potential during FAOR.

To gauge how the FAOR is affected in the potentials regions above H_{UPD} we conducted pulsed voltammetry inspired by Clavilier et al. In this type of pulsed voltammetry experiments, each potential investigated is separated by a surface re-initialization (at 1.05 V_{RHE}) cleaning the surface.
for all poisons through surface oxidation. The impact from dissolution at this oxidizing potential should be minimal.\textsuperscript{58,59}

\begin{figure}[h]
\centering
\includegraphics[width=\textwidth]{image.png}
\caption{2 s pulsed voltammetry and CVs at 10 mv/s at room-temperature at 1600 rpm in Ar-saturated 0.1 M HClO\textsubscript{4} with 0.1 M HCOOH. (a) Pt/C. (b) Pd/C. Measurements repeated three times.}
\end{figure}

Figure 6 shows pulsed voltammograms and corresponding CVs for Pt/C and Pd/C samples. It appears that only Pt/C’s activity is by the pulses, indicative of poisoning at potentials above those associated with the H\textsubscript{UPD} induced disproportionation to CO. In contrast, Pd/C does not show any changes from the pulsing. Essentially, these results suggests that the *COOH vs. *OOCH formate map is a poor descriptor for identifying, which catalysts are likely to be poisoned by non-CO species during FAOR. We believe this warrants a careful mention, to discourage future misinterpretations: It seems catalyst favoring *COOH over *OOCH performs better during FAOR above H\textsubscript{UPD} potentials but assigning this to an effect of surface poisoning by *OOCH is neither supported from our experiments (additionally see Figure S9) or the literature\textsuperscript{42,56}. 
In general, in a selective two-electron reaction a single intermediate binding energy should be sufficient to describe the reaction, as it is the case of hydrogen evolution or oxidation reaction (HER or HOR). Contrary, selective multi-electron reactions require more intermediates, as it is the case for oxygen evolution and oxygen reduction reaction (OER and ORR). Unfortunately, the traditional tool to gauge activity — the theoretical volcano plot — applicability is challenged when considering hetero-selective reactions. Given that the FAOR is a two-electron reaction, it should ideally have only one important intermediate. However, due to the unwanted side reactions—the disproportionation towards CO—this is not the case. From a theoretical perspective, this means that a FAOR volcano can be drawn, but never truly probed in experiments, as the volcano will encompass regions where competitive disproportionation dominates. Instead, we can probe the energetic conditions allowing for \(^*\text{COOH}\) adsorption/desorption while avoiding H\text{UPD}, which allow us to draw an active catalyst area, at various potentials where selective FAOR towards CO\(_2\) occur, see Figure 7.
Figure 7. Energetic considerations of FAOR. (a-c) *COOH vs. *H binding energy maps, with the scaling of the metals (light-blue) and single-sites (black) and the platinum catalyst (light-blue cross). The horizontal and vertical line denotes FAOR and HOR equilibriums when applying a potential. Both lines moves correspondingly when applying a potential, outlining an area where active catalyst avoid CO disproportionation. (d-f) Corresponding free energy diagrams of (a-c) for FAOR and HOR for the optimal metal catalyst using the metal scaling (black/light-blue) and the platinum catalyst (light-blue) when applying a different potential. In (a-f) the potentials selected are the FAOR equilibrium potential (-0.12 V RHE), HOR equilibrium potential (0.0 V RHE) and the limiting potential on Pt (0.17 V RHE).

Figure 7 combines the fundamental findings limiting the FAOR. Figure 7a-c show the *COOH vs. *H binding energy where the horizontal and vertical line depict the FAOR and HOR equilibrium at three relevant potentials, respectively. Once potentials above the equilibriums are applied, an
energetic area (shaded red) defining the conditions in which active and selective catalysts can be found. Using the metal scaling, Figure 7d-f shows the corresponding free energy diagrams for FAOR and HOR, for the optimal metal catalyst and Pt at relevant potentials. Three potentials were deemed relevant, that is; the FAOR equilibrium potential (-0.12 V_{RHE}), the RHE potential (0.0 V_{RHE}) and the limiting potential at Pt (0.17 V_{RHE}).

Figure 7a-c highlights that there are three limiting potentials important when considering FAOR electrocatalysis: (i) The weak binding of \(^*\text{COOH}\) binding governing the possible FAOR onset. (ii) The strong binding of \(^*\text{H}\), imposing possible CO disproportionation and/or HER. (iii) Active FAOR limitations imposed by the applied potential in relation to, too strong binding of \(^*\text{COOH}\), governing \(^*\text{COOH}\) retention on the catalyst surface during FAOR.

Figure 7d-f suggest that the optimal metal catalyst found on the \(^*\text{COOH}\) vs. \(^*\text{H}\) scaling, although having a \(^*\text{COOH}\) binding facilitating FAOR at relevant potentials, will be limited by having \(^*\text{H}\) on the surface until \(\sim 0.0 \text{ V}_{\text{RHE}}\). This can interpreted as: \(^*\text{H}\) will always bind to the surface when the energy level of \((\text{H}^++e^-)\) is above \(^*\text{H}\). Hence, a working catalyst should favor \(^*\text{H}\) binding weaker than \((\text{H}^++e^-)\) not to be limited by disproportionation. This interpretation form the basis of a theoretical FAOR overpotential, \(\eta=0.12 \text{ eV}\). Figure 7 also suggests that platinum generally is limited due to being too reactive toward hydrogen and consequently explain all the Pt-based FAOR literature discussed here and in the SI.

A final note should be made concerning the analysis in Figure 4 and 7. The interpretation and data strongly suggest that a single-site Co-MNC catalyst should not be restricted by \(^*\text{H}\), whilst having a nearly ideal \(^*\text{COOH}\) binding value. In total Co-MNCs should work as an efficient catalyst for FAOR near its equilibrium potential, assuming the Co-MNC is stable under acidic conditions.
**Conclusion**

In conclusion we have shown the fundamental limitations of FAOR and used this to derive the underlying volcano. We have observed that for an ideal catalyst, the FAOR equilibrium potential should be above its corresponding H$_{\text{UPD}}$ potential in order to avoid the disproportionation. Additionally, such catalyst should have an optimal $^\ast$COOH binding and is suggested to generally favor $^\ast$COOH over $^\ast$OOCH. We found that $^\ast$COOH and $^\ast$H binding scale on both metal and single-site catalysts. This creates the fundamental limitation of either too weak $^\ast$COOH resulting in a high overpotential or limiting potential due to H$_{\text{UPD}}$ mediated disproportionation on the surface. The carbon–hydrogen scaling is indeed a fundamental limitation, analog to the $^\ast$OH and $^\ast$OOH scaling for oxygen evolution and reduction. Experimentally, we show that a good performing FAOR catalyst should have the attributes of; (i) a close to the fundamental derived onset, (ii) no hysteresis between anodic- and cathodic CV scans and (iii) high and stable FAOR CA currents above the derived onsets fundamental limits. Attributes (i-iii) can all be achieved considering the $^\ast$COOH vs. $^\ast$H map or volcano, Figure 4a and 5b, respectively.

**ASSOCIATED CONTENT**

**Supporting Information.** The Supporting Information is available free of charge on the ACS Publications website at DOI: XXXXXXXX. Relevant aspects of the computational model and methods and the experimental methodology are included herein. (PDF). Additionally, atomic structures and analysis scripts are available on the webpage: https://chem.ku.dk/research_sections/nanochem/theoretical-electrocatalysis/
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The manuscript was written through contributions of all authors. All authors have given approval to the final version of the manuscript. †These authors contributed equally. K.D.J. conceptualized experiments with the aid of M.A., M.E.-E., J.D. and D.Z. and conducted the majority of the experiments with the aid of M.R. and I.P. A.B. together with J.R. conceptualized the necessary
simulations and conducted the majority of the computational simulations with the aid of R.L. Finally, J.R, M.A., M.E.-E., R.L., A.B. and K.D.J. all participated in the discussions of reaction mechanism and helped develop this mechanistic study. A.B. and K.D.J. jointly drafted the manuscript.

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ABBREVIATIONS

CA, chronoamperometry; CV, cyclic voltammogram/voltammetry; CO$_2$RR, CO$_2$ reduction reaction; DFAFC, direct formic acid fuel cell; DMFC direct methanol fuel cell; DFT, density functional theory; fcc, face centered cubic; FAOR, formic acid oxidation reaction; FC, fuel cell; GCHE, generalized computational hydrogen electrode; HER, hydrogen evolution reaction; H$_{\text{UPD}}$, hydrogen underpotential deposition; MNC, metal-nitrogen-carbon; OER, oxygen evolution reaction; ORR, oxygen reduction reaction; RHE, reversible hydrogen electrode; RDE, rotating disk electrode; SI, supporting information;

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TOC