Correlations between experiments and simulations for formic acid oxidation

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ABSTRACT. Electrocatalytic 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 site-blocking species during the formic acid oxidation reaction. Recent studies have elucidated how the binding of carbon and hydrogen on catalyst surfaces promote CO₂ reduction towards CO and formic acid. This has also given fundamental insights to the reverse reaction, 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 correlate data on different catalysts to show that (i) formate, which is the proposed formic acid oxidation intermediate, has similar binding energetics on Pt, Pd and Ag, while Ag does not work as catalyst, and (ii) *H adsorbed on the surface results in *CO formation and poisoning through a chemical disproportionation step. Using these results, the fundamental limitations can be revealed and progress our understanding of the mechanism of the formic acid oxidation reaction.

Introduction

Tremendous efforts are currently going into out-phasing fossil fuels in favor of sustainable fuels.¹ This is motivated by our need to close the carbon cycle² and pave the way for new fuel production routes.³ Electrocatalytic technologies can in the future possibly allow direct electrification of chemical and fuel production. Examples include reduction of CO₂ towards CO, HCOOH, C₂H₄, C₂H₅OH and H₂O towards H₂.¹ Efficient fuel consumption through fuel cells (FCs) also holds great potential.⁴ Liquid fuels such as formic acid and methanol have attracted a lot of attention due to their: Viable energy density per mass- and volume, attractive handling/storage properties and
potential uses in other non-fuel applications, e.g. as a high value chemical building blocks for industry.\(^3\)

Some liquid fuels such as methanol, are notoriously limited in the oxidation toward CO\(_2\) since the process goes through a CO intermediate.\(^5\) 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.\(^6\)

Table 1. Estimated round-trip efficiencies \(\eta = \frac{U_{OER} - U_{Fuel \ cell \ reaction}}{U_{OER} + U_{Electrolyzer \ reaction}} \times 100\%\) calculated using the difference in energy potentials. Common for hydrogen, formic acid, and methanol we use \(U_{OER} = 1.6 \text{ V}_{\text{RHE}}\) and \(U_{ORR} = 0.8 \text{ V}_{\text{RHE}}\). Here only the cost of syngas production is considered, not the full formation of methanol. **Typical charge/discharge efficiency.
| Energy stored (electrolyzer) | Energy released (fuel cell) | Round trip efficiency, $\eta$ |
|-----------------------------|-----------------------------|-------------------------------|
| Hydrogen                    | $U_{HER} = -0.1 \text{ V}_{RHE}$ | $U_{HOR} = 0.1 \text{ V}_{RHE}$ | $\approx 41\%$ |
| Formic acid                 | $U_{CO_2\rightarrow HCOOH} = -0.8 \text{ V}_{RHE}$ | $U_{FAOR} = 0.2 \text{ V}_{RHE}$ | $\approx 25\%$ |
| Methanol*                   | $U_{CO_2\rightarrow CO+H_2} = -0.6 \text{ V}_{RHE}$ | $U_{CO\rightarrow CO_2} = 0.65 \text{ V}_{RHE}$ | $\approx 7\%$ |
| Li-battery**                |                           |                               | $\approx 90\%$ |

Depending on the $U_{FAOR}$, direct formic acid fuel cells (DFAFCs) 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 more energy than methanol per molecule.

Key to understanding the limitation of formic acid oxidation is the direct link to the reverse electrochemical reaction, i.e. the CO$_2$ reduction reaction (CO$_2$RR).$^7,^8$ CO$_2$RR to formic acid and the formic acid oxidation reaction (FAOR) can be written in the form:

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

$$\text{(FAOR)} \quad \text{HCOOH} \rightarrow \text{CO}_2 + 2\text{H}^+ + 2e^- , \quad \Delta G^0_{\text{FAOR}} = -0.12 \text{ eV per H}^+ + e^- \quad (2)$$

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.$^9-^{12}$ 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 (with limited amounts of H$_2$), and finally, formic acid is produced on Pb, In, Hg, Sn, Cd and Tl with almost 100% faradaic efficiency.$^{11}$ Importantly, hydrocarbons are uniquely produced on Cu.$^{11}$ Using simulations, we were able to classify the
CO₂RR product distributions towards hydrogen, hydrocarbons, CO and formic acid due to the catalyst’s affinity towards adsorbed *H and *CO.¹⁰ Interestingly, we noted that the CO₂RR appears selective towards formic acid when weakening *H binding (i.e. when there is no *H on the catalyst surface). Moreover, from this study¹⁰ we noted that the *OOCH vs. the *COOH intermediate cannot be used to distinguish the CO or formic acid product formation. Where previous works used *OOCH as a descriptor for formic acid oxidation and *COOH as a descriptor for CO production, and even recent discussion for CO₂ reduction highlights the possible formic acid formation through *COOH.¹³ These findings and discussion are readily usable when considering the reverse reaction FAOR, which involves similar reaction intermediates to CO₂RR and vice versa.

FAOR exhibits the highest intrinsic activity on Pt and Pd¹⁴–¹⁹ However, the reaction is affected by high overpotential and formation of various poisoning and site-blocking intermediates.²⁰ The following observations are reported in the literature, as also shown in Supporting information (SI) Figure S1: (i) FAOR onsets currents at low potentials does not necessarily correspond to high FAOR currents. (ii) Hysteresis between anodic and cathodic scans is a common occurrence, showing a higher current in cathodic scans in than anodic ones, which shows that the reaction rate depends on the prehistory. Typically, this “memory effect” is related to unwanted reactions which form species that block the surface, i.e. poisoning species. (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 FAOR⁸. 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{20–24} which we illustrate by the literature study overview in Figure 1. A dual-pathway mechanism for formic acid oxidation is established by the community:\textsuperscript{25,26} The direct pathways, which leads to the desired final product of CO\textsubscript{2} through the formate adsorption,\textsuperscript{27} and another path where adsorbed CO, the poisoning species, is formed. The CO formation can be thought of as being formed through a so-called chemical disproponation reaction, where an activated formic acid intermediate reacts with a hydrogen to form water and CO (e.g. $^*$COOH +$^*$H \rightarrow $^*$CO + H\textsubscript{2}O). However, the nature of the reactive intermediate in the direct pathway is still under strong debate and it is not given that it is the most stable intermediate (e.g., formate) should also be the reaction intermediate. The community has focused on elucidating the reaction mechanism and attempted to circumvent of poisoning issues: through pathway engineering,\textsuperscript{21,22} changing electrolyte composition,\textsuperscript{23} or inclusion of sites with the ability to remove poisoning or site-blocking species.\textsuperscript{24,28–30} CO-poisoning from partial HCOOH oxidation is often considered the principal culprit\textsuperscript{28} and various works suggest CO formation can be avoided utilizing a single/dual-site catalyst.\textsuperscript{31–34} However, catalysts such as Au-Pt,\textsuperscript{21} Pt-Hg/C\textsuperscript{35} and Pd-Hg/C\textsuperscript{36} exhibit limited catalytic improvement over their pure metal counterparts, for some overviews see Figure S4 in SI. In this work, we will take a new view on all possible pathways for working catalysts. We carry out electrochemical characterization on Pd/C and Pt/C catalyst as well as Pt-Hg/C,\textsuperscript{35} Pd-Hg/C\textsuperscript{36} and Pt-Bi/C,\textsuperscript{37} following the protocols reported in the literature for the preparations of these catalysts. Furthermore, we try to validate the possible reaction path for not/low performing catalysts (e.g., Ag) for formic acid oxidation.
Figure 1. Literature study highlighting all the conceived FAOR reaction pathways during potential cycling. Historically, FAOR has been split into the direct (gray)\textsuperscript{39,40} and indirect (green and yellow)\textsuperscript{16,39,41,42} pathways. Further, partial FAOR and catalyst oxidation forming unwarranted surface blocking have been suggested, e.g., CO\textsubscript{x}H\textsubscript{y} species (purple),\textsuperscript{20,25,43} CO (red)\textsuperscript{44} and hydroxide/oxides (blue)\textsuperscript{38}. Even CO\textsubscript{2}RR induced CO formation by applying too cathodic potential (brown)\textsuperscript{40} have been suggested. Recently, formate (yellow)\textsuperscript{28,39,45,46} in various arrangements has gained attention as potential catalyst site-blocking agents.

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

(i) Is the FAOR activity correlated with the \textsuperscript{*}COOH or the \textsuperscript{*}OOCH intermediate?

(ii) How is CO formed during the FAOR?

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

**Results and Discussion**

**Figure 2.** 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 HClO₄ 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<sub>RHE</sub> 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. Note, Pt-Bi/C was unstable at higher potentials (see SI) hence a reduced potential range was utilized.

Figure 2 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. These investigated catalysts are synthesized from the same starting materials, i.e. premade Pd or Pt catalyst which we then modified by (electro)deposition of Hg or Bi at room temperature. Consequently, it is reasonable to expect that no significant impact of the catalysts’ surface area arises due to the modification, allowing for catalyst comparison. However, verifying this expectation using standard electrochemical surface area (ECSA) evaluation (e.g. utilizing *CO-stripping or hydrogen under potential deposited, H-UPD, charges) is not possible as these species do not adsorb specifically on Hg and Bi modified catalysts.

Figure 2a shows the formic acid oxidation CV during rotation and 10 mV/s scan-rate on the five catalysts. Figure 2b depicts the base CVs exhibiting suppressed hydrogen underpotential deposition (H_{UPD}) on the Pt-Bi/C, Pt-Hg/C and Pd-Hg/C compared to the Pd/C and Pt/C counterparts. Figure 2c displays the formic acid oxidation CA at 0.55 V_{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 2a indicating an irreversible change in the catalyst going to low potentials. Interestingly, taking a combined view on Figure 2a,c shows that forming single-sites of Pt through Hg alloying^{35,36} 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-S13. To explain these observations in Figure 2 we turned to DFT.
**Figure 3.** DFT calculated binding energies for metal fcc(111) (*light-blue crosses*) and single-site catalyst. MNC-based (*black points*), Pt atom in Au(111) denoted Pt1Au(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 \( \Delta E_{FAOR}^0 \approx \Delta G_{FAOR}^0 \). Further, we used CO2 and H2 for references when calculating *COOH, \( \Delta E_{FAOR}^0 \) is 0.12 eV per electron.

Figure 3a maps out the DFT calculated *COOH vs. *H binding energies of the most relevant model metal (111) facets and single-site catalysts, such as MNCs, PtHg4,35 PdHg436 and single Pt atoms in Au, Pt1Au(111)21 (for computational details see SI). *Via* simulations we study both metals and the single-site-catalysts as they both have a linear scaling: \( \Delta E_{*\text{COOH}} = \Delta E_{*\text{H}} + b \). However, \( b \) is about 0.29 eV as previously47 observed for metals and 0.0 eV for single-site catalysts, respectively. This allows for a fundamental destabilization of *H vs. *COOH at the single-site catalyst motifs as compared to metals, which is beneficial for improved FAOR by limiting the disproportionation reaction. Besides the scaling, a vertical- and a horizontal line indicating HUPD (\( \Delta E_{HUPD} = 0 \) for \( 1/2\text{H}_2 \leftrightarrow *\text{H} \)) and formic acid’s thermodynamic equilibrium potential have been included in Figure 4a,b. Here its assumed that \( \Delta E_{FAOR}^0 \approx \Delta G_{FAOR}^0 \), as thermodynamic corrections and water stabilization are expected to cancel out for the *COOH intermediate. The challenge to understand the implications of Figure 3a is that one needs to compare *H as a reduction from H2O and *COOH as oxidation from HCOOH. This means that *H stabilizes with negative potential, while *COOH stabilizes with positive potential during FAOR. Essentially, from Figure 3a it can be inferred that materials to the left of approximately \( \Delta E_{HUPD} \) are limited by disproportionation at low potentials. While materials to the right are limited by weak binding *COOH. Most importantly, the fundamental scaling between *COOH and *H matter as adsorption of *H on the surface leads to the
possibility of disproportionation towards CO. In conclusion, the single-atom catalyst scaling crosses the FAOR thermodynamic potential at values significantly above $\Delta E_{\text{Hupd}}$, which suggests that single-atom catalyst may work for FAOR near the FAOR equilibrium potential without carrying out the disproportionation reaction.

Figure 3b displays the DFT calculated binding energies of $^*\text{OOCH}$ vs. $^*\text{H}$. Conversely to Figure 3a, there is no apparent scaling between formate bidentate and adsorbed hydrogen (however the oxygen bond for $^*\text{OOCH}$ scales with $^*\text{OH}$). However, notable there would be a scaling between $^*\text{OOCH}$ and (bi)-carbonate (HCO3/C03), which also would have surface binding through oxygen to the surface. Carbonate species could be formed over time due to the equilibrium with CO2 and hence poisoning the surface, revealing a decaying activity as observed in Figure 2c for all working FAOR catalysts. Investigating $^*\text{OOCH}$ as a function of $^*\text{H}$ is important as $^*\text{OOCH}$ has been suggested as an important reaction intermediate. Figure 3b shows that the binding energy of the $^*\text{OOCH}$ intermediate on Ag is slightly too strong, but very close to the thermodynamic potential and similar to Pt and Pd. Essentially, if $^*\text{OOCH}$ was the important reaction intermediate for FAOR, then Ag should be working as FAOR catalyst at very low potential, and furthermore Ag does not suffer from disproportionation (as Ag $^*\text{H}$ binding far exceeds $\Delta E_{\text{Hupd}}$). However, Ag is experimentally shown to be inert for the FAOR. The role of formate in the reaction is clearly a puzzle. For FAOR surface-enhanced IR absorption on Pt has experimentally confirmed formate on the surface27 above 0.7 V_RHE and further for CO2RR in situ surface-enhanced Raman spectroscopy on Ag has experimentally confirmed formate on the surface.48 As Pt converts formic acid to CO2 and Ag converts CO2 to CO, there seems no relation between the experimental observation of formate on the surface as reaction intermediate. We note that $^*\text{OOCH}$ 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 3c shows the *OOCH vs. *COOH binding for the catalyst, with a dashed line indicating the affinity towards carbon or oxygen. Depending on the catalyst we can see whether formate bidentate or carboxyl is favored. The working catalyst is Pt, which is well above the diagonal and Pd which is at the diagonal, while poorly working FAOR catalysts e.g. Ir or Au are slightly below the diagonal. Catalysts having stronger *OOCH binding vs. *COOH (below the diagonal), are basically oxidized before they can carry out FAOR.

The type of analysis illustrated in Figure 3 is a powerful tool able to identify which catalyst suffers from disproportionation, poisoning or oxidation. By virtue of the *COOH vs. *H scaling-relations, it gives fundamental insights into why literature historically has shown no significant FAOR activity at the thermodynamic potential for a metal catalysts. Implicitly Figure 3a shows disproportionation occurring at low potential arising from the H\textsubscript{UPD} and the consequent creation of *CO poisoning species. The creation of *CO leads to hysteresis between the anodic and cathodic sweeps in FAOR CVs. From this insight, one would expect that disproportionation is mitigated in the CVs by simply staying above H\textsubscript{UPD} potentials.
Figure 4. 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. Note, that after cycling in the different potential limits returning to the full range of 0.025 to 1.05 V$_{RHE}$ re-initializes the FAOR response i.e. no irreversible changes arise due to dissolution or sintering.

Figure 4 shows FAOR CVs of Pt/C, Pt-Hg/C and Pd/C cycled with varying potential limits. The potential ranges from 0.00-1.05 V$_{RHE}$ reveals that Pt/C and Pt-Hg/C are poisoned in anodic sweeps. While, 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 indicate 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}, which then competes with \(^1\)H adsorbed, \textit{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\(^{11}\).

Concerning Figure 4, 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\(^{53,54}\) due to insufficient surface reduction or accumulation of either \(^*\)OOCH, \(^*\)COH, \(^*\)OCOH or \(^*\)CO species. Interesting is that \textit{in situ} Fourier-transform infrared spectroscopy (FTIR) work\(^{57,58}\) has shown that Pt, contrary to Pd, continuously form CO above \(H_{UPD}\) potential during FAOR.

Finding that \(^*\)H limits FAOR activity at low potential \textit{via} disproportionation, allows one to hypothesize about the Pt-Bi and Pt-Hg systems, which perform better than Pt on two different perspectives. Pt-Bi shows no hysteresis and is active at higher overpotential as compared to Pt. The absence of a hysteresis in the CV indicates that Pt-Bi somehow circumvent the disproportionation reaction, potentially by blocking the surface for \(^*\)H at low potential as onset for Pt-Bi is higher than on Pt. Pt-Hg on the other hand is more active in the negative sweep, particular at lower potential, but does show clear hysteresis. Further we hypothesize that the Pt single-sites in Pt-Hg destabilizes \(^*\)H shifting the onset to lower potentials and increasing the activity at low potential, simply due to a limited disproportionation reaction.
To gauge how the FAOR is affected in the potentials regions above $H_{UPD}$ we conducted pulsed voltammetry inspired by Clavilier et al.\textsuperscript{59} 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{60,61}

![Figure 5](image.png)

**Figure 5.** 2 s pulsed voltammetry and CVs at 10 mv/s at room-temperature at 1600 rpm in Ar-saturated 0.1 M HClO$_4$ with 0.1 M HCOOH. (a) Pt/C. (b) Pd/C. Measurements repeated three times.

Figure 5 shows pulsed voltammograms and corresponding CVs for Pt/C and Pd/C samples. Interestingly Pt/C becomes more active when pulsing, which is in contrast to Pd/C that does not show any changes from the pulsing. This experiment challenges *OOCH species as site-blocking species. Since Pd has stronger relative *OOCH to *COOH binding than Pt, and it should hence be on Pd where activity was affected by blocking of *OOCH species. One view for this observation
could be given in the recent work by *Koper and coworkers*, who noted that formate adsorption is important for formic acid oxidation, not as an active intermediate, but more as a self-protector against CO poisoning.

**Figure 6.** FAOR data on a Cu, Ni and Ag wire at 50 mV/s at room-temperature Ar-saturated 0.1 M HClO$_4$ with 0.1 M HCOOH taken at 50 mV/s.

Beyond Pd and Pt catalyst, in Figure 6, we tested Cu, Ni and Ag wires as FAOR catalyst. The experiments revealed that Ni indeed seems very active to oxidize either HCOOH or Ni. However, going to potentials above 0.2 V$_{RHE}$ tended to bring out a yellowish tinge in the electrolyte, and by looking into Ni's *Pourbaix* diagrams it appears to readily dissolve as Ni$^{2+}$ at 0.15 V$_{RHE}$ at pH 1, consequently making it a poor FAOR catalyst (unless there is some very narrow window that Ni is stable enough to oxidize HCOOH without dissolving). Similarly, Cu shows no FAOR activity only the well-known Cu oxidation current at potentials larger than 0.2 V$_{RHE}$ is observable. Ag is perhaps active towards FAOR but again this occur close to Ag dissolution potentials suggesting it to be a rather poor FAOR catalyst.
Most interesting when combining the observation of similar intermediate binding energetics in Figure 3, we identified that Ag has similar binding energetics of formate as Pd and Pt, whereas Cu and Ni do not follow the energetic of Pd and Pt. Hence Cu and Ni can be used as test catalysts far from known working catalysts. However, as tested here in Figure 6 there appears no significant FAOR on Ag, Cu and Ni.

**Conclusion**

In conclusion we have correlated the FAOR activity with simulated DFT binding energies of *COOH, *OOCH and *H across multiple metal catalysts. We have observed that for an ideal catalyst, the FAOR equilibrium potential should be above its corresponding H\textsubscript{UPD} potential in order to avoid the disproportionation. We found that *COOH and *H binding scale on both metal and single-site catalysts. This creates the fundamental limiting potential due to H\textsubscript{UPD} mediated disproportionation on the surface. The carbon – hydrogen scaling is indeed a fundamental limitation, analog to the *OH and *OOH scaling for oxygen evolution and reduction. Experimentally, we show that a good performing FAOR catalyst should have the attributes of: (i) an onset 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. Interestingly, this works concludes on the direct relation between FAOR and CO\textsubscript{2}RR; *H in combination with *COOH forming CO in both FAOR and CO\textsubscript{2}RR.

**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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ABBREVIATIONS

CA, chronoamperometry; CV, cyclic voltammogram/voltammetry; CO₂RR, CO₂ 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_{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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