Recent advances in formic acid electro-oxidation: from the fundamental mechanism to electrocatalysts

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Direct formic acid fuel cells have attracted significant attention because of their low fuel crossover, high safety, and high theoretical power density among all the proton-exchange membrane fuel cells. Much effort has been devoted to the study of formic acid oxidation, including the reaction processes and electrocatalysts. However, as a model reaction, the anodic electro-oxidation process of formic acid is still not very clear, especially regarding the confirmation of the intermediates, which is not helpful for the design and synthesis of high-performance electrocatalysts for formic acid oxidation or conducive to understanding the reaction mechanisms of other small fuel molecules. Herein, we briefly review the recent advances in investigating the mechanism of formic acid electro-oxidation and the basic design concepts of formic acid oxidation electrocatalysts. Rather than an exhaustive overview of all aspects of this topic, this mini-review mainly outlines the progress of this field in recent years.

1. Introduction

Environmental pollution and the pending energy crisis are two major problems that need to be solved urgently. Due to the outstanding advantages of cleanliness and high energy density, proton-exchange membrane fuel cells (PEMFCs) can effectively aid alleviating the above-mentioned problems. However, although PEMFCs have been studied intensively since the 1960s, inherent limitations still exist. For example, the anode has the following prominent problems: unsatisfactory safety, high fuel crossover, high cost of catalysts, low durability, and a great loss of power density and cell efficiency due to the polarization and slow reaction kinetics of organic small molecular fuels in the electro-oxidation process. Recent studies have demonstrated that formic acid can be used as a suitable anodic fuel to overcome the safety problems and fuel crossover issues. The main reasons for this are the following. First, formic acid can be used as an additive in food and medicine and is not flammable, and therefore it is

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safe to store and transport.\textsuperscript{20,21} Second, formic acid can be ionized into its formic acid root, which is mutually repulsive with the sulfonic acid root of Nafion film, thereby partially hindering the crossover of formic acid and thus reducing the probability of producing mixed potential at the cathode. The low formic acid crossover allows using a high concentration of formic acid as fuel, which compensates for the lower energy density of formic acid compared to other liquid fuels.\textsuperscript{12-14} Moreover, the theoretical open circuit voltage (OCV) of direct formic acid fuel cells (DFAFCs) is 1.48 V, which is higher than other existing PEMFCs.\textsuperscript{15,16} Such a high voltage can bring high power density, which enhances the PEMFC’s practical application in the fields of electric vehicles, portable electronic equipment, and others.\textsuperscript{7,18}

At present, the cost, durability and catalytic activity of the catalysts are mutually restrictive problems that commonly exist in almost all PEMFCs.\textsuperscript{15,20} Therefore, the design and synthesis of high-performance electrocatalysts are one area that represents one of the most critical research content aspects in fuel cells.\textsuperscript{21,22} The effective design and preparation of an excellent electrocatalyst for formic acid oxidation should be based on the following two aspects. Firstly, the reaction mechanism of formic acid oxidation should be explored, including the reaction path, reaction intermediates, species adsorption states, and adsorption configurations. Based on the understanding of the reaction mechanism, the compositions and surface structures of the electrocatalysts can then be rationally designed and adjusted. Although formic acid is a relatively simple type of fuel molecule among the commonly used organic fuels for fuel cells, the electro-oxidation process is still complicated. During the past decades, much work has been done to elucidate the reaction mechanism. To this end, single-crystal catalysts with different surface planes have been used as model catalysts for formic acid. Meanwhile, different \textit{in situ} spectroscopic techniques combined with electrochemical methods have been widely used to detect the surface adsorbed species during formic acid oxidation, with an aim to elucidate the possible reaction mechanism.\textsuperscript{23}

In recent years, with the development of \textit{in situ} and \textit{operando} techniques, great advances have been achieved in the above-mentioned two aspects of formic acid oxidation, \textit{i.e.}, the reaction mechanism and electrocatalysts. However, a detailed summary of the advance in the reaction mechanisms and electrocatalysts related to formic acid electro-oxidation (FAEO) is insufficient but imperative.\textsuperscript{24,25} Consequently, in this mini-review, the work and progress on the electrocatalytic oxidation mechanism, electrocatalyst design, and synthesis for formic acid oxidation in recent years are summarized. Furthermore, the fundamental chemistry of DFAFCs and FAEO are also discussed. Finally, a brief conclusion is provided. We hope this review is helpful for readers interested in this field to understand the current research situation of formic acid electro-oxidation.

## 2. Fundamental electrochemistry of DFAFCs and FAEO

PEMFC is an energy conversion device that converts chemical energy from broken bonds in small fuel molecules into electrical energy.\textsuperscript{26,27} For PEMFCs, the energy density, power density, and cycles are the important performance-evaluation parameters.\textsuperscript{2} Energy density is defined as the amount of energy stored and released by devices of the same mass/volume, so the theoretical energy density of DFAFCs can be described as:

\[
\text{Energy \ density} = \frac{n \times F \times \text{OCV} \times (\text{MW})^{-1}}{\text{weight or volume}}
\]

where \(n\) is the number of electrons transferred, \(F\) is the Faraday constant, \(\text{OCV}\) is the open circuit voltage, \(\text{MW}\) is the molecular weight of the fuel, and \(\text{weight or volume}\) is the weight or volume of the device. DFAFCs can provide fundamental information of FAEO. As illustrated in Fig. 1a, for formic acid oxidation on a Pt/C catalyst, the oxidation current peak I at \(\text{O}_{2}\) (vs. RHE) corresponded to the direct oxidation of HCOOH to CO2. Obviously, because the surface of Pt/C has a relatively constant coverage of CO poisoning species below 0.85 V (vs. RHE), only a very low current density could be observed. Meanwhile, the peak II at \(\approx 1.05\) V (vs. RHE) was assigned to the electro-oxidation of CO intermediates accumulated in the indirect oxidation pathway. Anodic sweep is often used to evaluate the activity of the used catalyst, and the intensity ratio of peak I and peak II is used to determine the pathway of FAEO on catalysts, \textit{i.e.}, a direct or indirect pathway. On the other hand, based on the oxidation currents in positive-going and negative-going scans in CV measurements, the CO anti-poisoning ability of the catalyst could be qualitatively evaluated.\textsuperscript{29,30} By comparing the CVs of formic acid oxidation on Pt/C and Pd/C (Fig. 1b and c), Pt/C showed higher catalytic activity, while Pd/C showed better anti-poisoning ability. On the other hand, both Pt/C and Pd/C have poor electrochemical stability. From Fig. 1d, the Ir/CN showed relatively higher stability than Pt/C and Pd/C, although it showed a higher overpotential for HCOOH oxidation.\textsuperscript{23} From this example, a simple CV curve can provide abundant information about the HCOOH oxidation on different catalysts.

In addition to CV, CO stripping (Fig. 1e) is generally employed to assess the electrochemical active surface area (ECSA) and the catalytic active sites,\textsuperscript{21} and CA (Fig. 1f) is often used to examine the electrochemical stability.\textsuperscript{31} Meanwhile, Tafel plots (Fig. 1g) can reveal the reaction rate and mechanism during the catalytic process, where a smaller Tafel slope suggests a faster charge-transfer kinetics.\textsuperscript{13} Moreover, EIS (Fig. 1h) is always used to investigate the charge-transfer...
3. Reaction mechanism of FAEO

Since formic acid is a kind of simple organic small molecule, it is suitable for use as a model molecule to study the reaction mechanism of the electro-oxidation of organic molecule fuels (HCOOH, CH₃OH, C₂H₅OH, etc.) on different electrodes. Although simple, the electro-oxidation of HCOOH has a complicated reaction processes and various possible routes to generate the CO₂ product, depending on the different electrode surface structures, as shown in Fig. 2a. Parsons et al. proposed the widely recognized two-pathway mechanism as early as 1973. That is, formic acid dehydrogenates in two steps and simultaneously electrons transfer to the electrode to form CO₂ and H⁺. That is, formic acid dehydrogenates in two steps and simultaneously electrons transfer to the electrode to form CO₂ and H⁺. The potential: HCOOH $\rightarrow$ active intermediate $\rightarrow$ CO₂ + 2H⁺ + 2e⁻. In the indirect pathway, formic acid first dehydrates to form CO, which is further oxidized to CO₂ at a higher potential: HCOOH $\rightarrow$ CO_ads + H₂O_ads $\rightarrow$ CO₂ads $\rightarrow$ CO₂ + 2H⁺ + 2e⁻. The product of CO₂ and the intermediate of CO and their transformation can be directly observed by using in situ attenuated total reflection-Fourier transform infrared spectroscopy (ATR-FTIR). As shown in Fig. 2b, for HCOOH oxidation on Pt black catalyst, in addition to the clear absorption signal of the CO₂ product at 2345 cm⁻¹, the peaks at 1869 cm⁻¹ can be assigned to the poisonous CO intermediate. Recently, Wang et al. reported an in situ electrochemical shell-isolated nanoparticle-enhanced Raman spectroscopy (EC-SHINERS) technique that could monitor the electro-oxidation processes of CO to CO₂. As shown in Fig. 2c, for CO absorbed on a Pt monolayer, the intensity of the band at 472 cm⁻¹, corresponding to the Pt–C stretching vibration, decreased drastically as the potential increased and then disappears after 0.6 V, suggesting the desorption and oxidation of CO at 0.6 V. Meanwhile, two peaks appeared when the potential rose up to 0.9 V, one broad band at 570 cm⁻¹, and another sharp peak at 330 cm⁻¹. These two peaks were attributed to the Pt–O stretch with the top and bridge configurations of CO₂. It should be noted that due to the complicated process, the possible oxidation mechanism of formic acid is under debate and the related research is still underway.

3.1 Active intermediate identification

Willsau et al. first demonstrated the direct pathway of HCOOH oxidation by combining the electrochemical differential mass spectrometry technique with the isotopic method. Usually, COH, CHO, –HCOO, and –COOH were considered as active intermediates until Osaka’s research group detected the bridge-bonded formate (HCOO₅⁻) with two oxygen atoms adsorbed on the platinum electrode surface by attenuated total reflection surface-enhanced infrared absorption spectroscopy (ATR-SEIRAS). At the beginning, adsorbed formate (HCOO₅⁻) was regarded as a reactive intermediate in the direct pathway through ATR-SEIRAS. Then objections were raised. Chen et al. proposed that bridge-bonded formate...
(HCOO\textsubscript{m}) was not a reactive intermediate but a triple path for FAEO. It was also predicted that in a direct pathway, the weakly adsorbed HCOOH\textsubscript{ads} precursor (HCOOH\textsuperscript{m}) is directly oxidized to CO\textsubscript{3} and the adsorbed formate (HCOO\textsubscript{m}) acts as a site-blocking spectator. These conclusions were based on the results of the ATR-FTIR technique with a thin-layer electro-oxidation of formic acid on commercial Pd black in 0.25 M HCOOH + 0.5 M H\textsubscript{2}SO\textsubscript{4} solution at different potentials from −0.20 to 0.60 V, with the reference spectra acquired at −0.25 V. 200 interferograms (resolution: 8 cm\textsuperscript{−1}) were collected and combined for each spectrum. Reproduced from ref. 18. Copyright 2018, Elsevier. (c) In situ electrochemical shell-isolated nanoparticle-enhanced Raman spectroscopy (EC-SHINERS) spectra of CO electro-oxidation on Au(111) @Pt monolayer in 0.1 M HClO\textsubscript{4} saturated CO. Reproduced from ref. 43. Copyright 2018, Wiley-VCH.

3.2 Poisoning intermediate identification

It is widely accepted that adsorbed carbon monoxide (CO\textsubscript{ads}) is the main poisonous species during the formic acid oxidation. Very early, infrared spectroscopy studies identified that CO\textsubscript{ads} is a poisoning intermediate in the indirect pathway of FAEO.\textsuperscript{56}

The above determinations of the active intermediates were from spectral results or classical theoretical calculations. More recently, the identity of the reactive intermediate has been studied by different methods. By using a combined experimental and computational approach, Vilaplana\textit{et al.} proposed that the reactive intermediate is monodentate-adsorbed formate (HCOO\textsuperscript{m}) and the presence of adjacent adsorbates inhibits the transition from HCOO\textsubscript{m} to HCOO\textsubscript{m}.\textsuperscript{57} Consistently, through DFT computations, Huang’s group found that HCOO\textsubscript{m} was the main active intermediate with HCOO\textsuperscript{−} as the main precursor, and HCOO\textsubscript{m} was the site-blocking species with HCOOH as the precursor.\textsuperscript{58} In recent years, based on single-crystalline electrochemistry and in situ electrochemical Fourier transform infrared spectroscopy (EC-FTIR), Chen and co-workers proposed that in the direct pathway of FAEO, formate is the site-blocking species instead of the active intermediate.\textsuperscript{59} By combining ATR-SEIRAS with ultraviolet (UV) reflectance measurements, another interesting observation made by Hartl \textit{et al.} showed that the galvanostatic electro-oxidation of formic acid involves a complex autocatalytic loop.\textsuperscript{60} In the process, all of the species mentioned above play roles, and the results come from the oscillations during the galvanostatic oxidations. On the other hand, bimetal catalysts have been used recently in discussing the mechanism of FAEO. For example, through DFT calculations, Meng \textit{et al.} proposed that the *COOH intermediate plays different roles for palladium atoms and palladium ensembles on the surface of copper; the presence of Pd ensembles mainly results in CO\textsubscript{2} and isolated Pd atoms can catalyze HCOOH dissociation to form CO.\textsuperscript{61} Additionally, Yang \textit{et al.} suggested that for the M\textsubscript{core}@Pd\textsubscript{shell} (M = Cu, Au, Co, Ni, Ag, Al) core@shell bimetal catalysts, the M@Pd (M = Au, Co, Ni, Ag), especially Ag, are highly selective for CO\textsubscript{2} formation that goes through the COOH or HCOO intermediate, depending on the types of core metal. Meanwhile, among the studied catalysts, Al@Pd is favorable for CO formation via the *COOH intermediate instead of the HCO\textsuperscript{−} intermediate.\textsuperscript{62}
And therefore, catalysts with high activity for CO electro-oxidation can efficiently remove the surface adsorbed CO and thus can significantly promote the FAEO.\(^{14}\) Just like with the identification of reactive intermediates, bridge-bonded formate (HCO\(_2\))\(^*\), dimeric formic acid, carboxylate (−COOH), CO\(_2\), and other species were once thought to be the precursors to CO.\(^{57,65-67}\) As for the formation origination of CO\(_{ads}\), Cuesta \textit{et al.} proposed that the presence of adjacent platinum atoms would lead to the formation of CO\(_{ads}\). This view has been widely accepted, and many efforts have been made to construct discontinuous platinum sites to mitigate the poisoning of platinum.\(^{14,66,67}\) Except for FAEO, palladium is less reactive than platinum in most electrochemical reactions.\(^{68}\) The reasons for this are not yet known, perhaps because palladium has a more specific affinity for hydrogen, allowing the rapid dehydrogenation of formic acid molecules on the palladium surface.\(^{13,21,69,70}\)

Meanwhile, the severe inactivation of palladium is often attributed to agglomeration, surface oxidation state change, component migration and loss, and CO-like toxic species gathering on the palladium surface. For CO-like toxic species, there is considerable evidence that CO is produced by the reduction of the product CO\(_2\) at a low oxidation potential, but there are other unknown reasons for the inactivation of palladium catalysts.\(^{71-73}\)

To reveal the mechanism of FAEO, electrochemical methods cannot provide direct information about the reaction parameters or intermediates, nor can they provide information about the electrode/solution interface at the molecular level. On the other hand, there is a gap between the density functional theory (DFT) simulated environment and the real electrode/solution interface. Therefore, it is necessary to combine some other technologies to achieve \textit{in situ} characterizations, just like differential electrochemical mass spectrometry (DEMS), \textit{in situ} infrared spectroscopy (\textit{in situ} IR), surface-enhanced Raman spectroscopy (SERS), and so on.

For the identification of intermediates, the reported conclusions are quite different due to the differences in the experimental apparatus and conditions, and the systems selected by researchers. Based on the current research, perhaps the most well-accepted two-pathway mechanism would be a good place to start, by using the same experimental method to study various forms of monometals and alloys in various specific systems, which involves single atoms, diatoms, clusters, monolayer atoms, nanoparticles, and films composed of different index facets. After a systematic study and summary, a cross comparison should be carried out to find the commonalities and the differences. Moreover, some possible intermediates during the FAEO have a short life and the present instruments may not be sensitive enough to detect them. Therefore, developing new devices, improving existing instruments, detecting breakdown products that are specific to the short-lived intermediates, and finding suitable probe molecules are very important for clearly and correctly revealing the reaction mechanism of FAEO. One step further, there are two main solutions to the problem of electrocatalyst poisoning caused by carbon monoxide: one is the introduction of oxyphilic active sites to rapidly oxidize carbon monoxide, and the other is to eliminate certain surface sites to inhibit carbon monoxide production.

### 4. Electrocatalysts for FAEO

The electrocatalysts for formic acid oxidation need to meet the following requirements: safety, low cost, high electrocatalytic activity, high tolerance to poisoning species, high electrical conductivity, high physical and electrochemical stability without particle agglomeration, surface oxidation state change, component migration and loss, carrier corrosion, and so on.\(^{6,16}\)

Base on the above desired requirements, the basic design concept of electrocatalysts for formic acid oxidation is to regulate the electronic structure and surface structure, which complement each other. However, the electronic structure effect should be considered first when designing the catalysts. In other words, the activation energy of the FAEO reaction could be reduced by selecting suitable electrocatalytic materials. The effect of the surface structure on the reaction rate and mechanism should be considered after selecting a suitable electrocatalyst. Currently, the surface structure manipulation includes the chemical structure (composition and valence), geometric structure (shape and form), surface atoms arrangement, electronic structure, and others. Among these, surface ligand effects, lattice strain effects, and synergistic effects are often used to explain the enhancement of catalytic activity, and meanwhile ensemble effects and third-body effects are often used to analyze the reasons that inhibit the formation of carbon monoxide.\(^{18,20,24,25,74}\)

#### 4.1 Pt- and Pd-based electrocatalysts for formic acid oxidation

Both platinum- and palladium-based materials are the traditional and the most widely used catalysts for the electro-oxidation of formic acid, but their research challenges are a little different.\(^{6,9}\) For platinum-based materials, inhibition of the indirect pathway and carbon monoxide adsorption are the core problems.\(^{9,21,75-77}\) However, improving the stability is the most critical issue for palladium-based catalysts.\(^{13}\) Up to now, a lot of efforts have been made by researchers to resolve the above-mentioned problems.

For example, controlling the shape and exposing high-index facets (Fig. 3a and b) with a low coordination number, open structure, and high surface energy can significantly enhance the catalytic activity and stability for FAEO.\(^{2,5-7}\) Through chemical vapor deposition (CVD) with an alloying-dealloying shape-regulating process, Huang \textit{et al.} synthesized tetrahexahedron (TTH)-shaped Pt–M (M = Sb, Bi, Pb, or Te), Pd–Bi nanocrystals on carbon black, and TTH-shaped Pd/C–Bi catalyst. The synthesized catalysts with high-index \{210\} planes (Fig. 3a) showed higher electrocatalytic activity and stability than commercial Pd/C and Pd/C catalysts for FAEO. With the prepared catalysts, the FAEO followed a direct reaction pathway. In addition to the foreign metal modification, the enhanced catalytic performance was mainly attributed to the high-index facets of the nanocatalysts.\(^{7}\) It should be noted that not all
metals with high-index facets will promote the catalytic performance for the electro-oxidation of formic acid. It has been found that the catalytic properties of metal nanocrystals are related to the surface structures. For Pt and Pd single-crystals, the catalytic activities of their basal planes for formic acid oxidation follow the order: Pt (110) > Pt (111) > Pt (100) and Pd (100) > Pd (111) > Pd (110).

For high-index facets of Pt and Pd nanocrystals, there is still no accurate conclusion due to the large number of studies carried out in different systems. Tetrahedron Pd \{100\} facets have been studied in detail and the oxidation current density of FAEO was found to have the following order: Pd (730) < Pd (830) < Pd (310) < Pd (1030) < Pd (1130) < Pd (100). Furthermore, El nabawy et al. found that the open facets of Au, Ag, Cu, Pt, and Pd required less over-potentials for the FAEO when compared to their close-packed facets, and the opposite was true for Ni, Ir, and Rh metals.

The catalytic performance for the electro-oxidation of formic acid can also be improved by taking advantage of the interaction between the metal and a suitable substrate. For example, Wang et al. reported the atomic dispersion of Pt on mesoporous sulfur-doped carbon (meso-S-C) with Pt loading as high as up to 10% by a \( \text{H}_2 \) thermal reduction process (Fig. 3c-e). The prepared hybrid showed enhanced catalytic performance for the FAEO. It was also found that the formation of Pt clusters in Pt/meso-S-C could reduce the catalytic performance. However, both of the prepared samples exhibited higher catalytic activity and stability than the commercial Pt/C. The authors ascribed the improved catalytic properties to the large surface area and mesoporous structure containing abundant and easily accessible sulfur-containing sites to accommodate a high mass percentage of Pt atoms. In fact, the isolated Pt sites and high Pt atom utilization were favorable for following a direct pathway. Ye et al. prepared Pd/cubes-CeO\(_2\) composite via a chemical reduction method by using ethylene glycol as a reducing agent. The presence of CeO\(_2\) could increase the metal dispersion and the ECSA. Meanwhile, with Pd dispersed on cubic ceria with a polar (100) surface and oxygen vacancies, the high degree of oxygen storage in CeO\(_2\) and the fast oxygen mobility at the noble metal–ceria interface could effectively resist Pd from being oxidized and could remove CO or other related poisons efficiently. In another work, Wang et al. used a microwave-assisted ethylene glycol method to synthesize a Pd–Fe\(_2\)P/C catalyst in which Pd nanoparticles were deposited onto an Fe\(_2\)P–carbon hybrid support. The fast charge-transfer kinetics and the appropriate down-shift of the d-band center of the Pd–Fe\(_2\)P/C catalyst significantly enhanced the catalytic performance for FAEO. The performance improvement was mainly attributed to the contribution from Fe\(_2\)P in the composite since Fe\(_2\)P can make the adsorbed CO and other toxic intermediates be oxidized at extremely low potentials.

In addition, constructing special nanostructures with a high specific surface area, edge areas, and surface defects can also
enhance the catalytic activity and stability for the FAEO. For instance, Ding et al. synthesized perpendicular Pd nanotube arrays (P-PdNTA) via a dual template electrodeposition method. Meanwhile, the massive Pd nanoparticles served as building blocks for the mesoporous walls of the nanotubes. The perpendicular and mesoporous structure provided a large electrochemical active surface area and active sites that were not easily aggregated. Additionally, there were anisotropic substructures on the catalyst surface as step edges and lattice defects (grain boundary, twin, etc.). These unique structures could accelerate the transportation, adsorption, and oxidation of formic acid. Recently, a one-pot hydrothermal synthesis of porous half-shells Pd−Pt in the presence of urea was reported by Tang and co-workers. Here, urea was used as a guiding surfactant to produce NH₃ and CO₂, which could act as a bubble template for building a hollow or porous structure. The enhanced catalytic activity and stability for FAEO was attributed to the high porosity, large electrochemical active surface area, fast mass transfer, abundant edge atoms, and a mass of crystal defects, such as distortions, a twin boundary, and atomic holes, as shown in Fig. 3f–i. Our group used a one-phase synthesis method to prepare Pt₁Au₂₄(SC₁₂H₂₅)₁₈ nanoclusters, which had a similar geometry with the Au₂₅ nanoclusters. It was found that although Au₂₅ nanoclusters exhibited very poor electrocatalytic activity for FAEO, a single Pt atom doping in the center of the Au₂₅ cluster could significantly enhance the catalytic performance. In the Pt₁Au₂₄(SC₁₂H₂₅)₁₈ nanocluster, the single Pt atom could decrease the CO production while the gold atoms could protect the catalyst from CO poisoning. Therefore, the electro-oxidation of formic acid followed a direct pathway with COOH* as the preferred reactive intermediate.

Finally, alloying Pt or Pd with other transition metals could change the d-band center, thus changing the interaction between the adsorbed species and the surface of the electrocatalysts. In such studies, high-angle annular dark field scanning transmission electron microscopy (HAADF-STEM), X-ray powder diffraction (XRD), and energy-dispersive X-ray spectroscopy (EDX) are commonly used methods to investigate the structure of alloys. For example, Duchesne et al. reported a colloidal method to prepare a series of platinum–gold (Pt–Au) nanoparticles by using ethylene glycol as a reducing agent. In these nanoparticles, different contents of platinum were modified on the surface of gold, and these NPs were categorized in terms of their surface structure (Fig. 4a): Au with single-atom Pt sites (Pt₇Au₉ and Pt₉Au₇), Au with single-atom and few-atom Pt sites (Pt₁₇Au₉₃), and Au-core/Pd-shell (Pt₇Au₂₂ and Pt₃₃Au₁₉) nanoparticles. It was found that the lower the number of platinum atoms, the better the electrocatalytic performance for formic acid oxidation. The characterization results showed that the high catalytic activity and stability were derived from the optimized elemental composition, low coordination numbers of Pt atoms, and the isolated Pt atoms preventing the catalyst from self-poisoning via CO generation. The most critical role was played by single-atom Pt surface sites surrounded by Au atoms. In addition, Luo et al. synthesized intermetallic PtSnBi nanoparticles (Fig. 4b–f) via a one-pot sequential complexing–reducing–ordering process. In this work, Bi played a key role in the formation of 2D nanoplates with an hcp crystalline structure and in suppressing the dehydration path during HCOOH oxidation. On the other hand, Sn could promote the CO oxidation on Pt sites by reacting with H₂O to form Sn–OHads. Therefore, the high FAEO catalytic activity of the PtSnBi nanoplates originated from the improved catalytic selectivity with direct oxidation from HCOO* to CO₂, rather than the anti-poisoning ability. Even further, Yang et al. used a seeded growth method with a self-assembly process to prepare a sea urchin-like Aucore@Pdshell electrocatalyst (Fig. 4g and h). The prepared material had a high ECSA with more exposed active sites and an obvious change of the d-band center of Pd. Meanwhile, the mismatched lattice constants between the Au core and Pd shell resulted in lattice strain. These factors could adjust the adsorption strength of formic acid molecules, intermediate species, and hydroxide radicals on Pd, which optimized

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**Fig. 4** (a) Synthesis of platinum–gold (Pt–Au) electrocatalysts. Reproduced from ref. 75. Copyright 2018, Nature. (b–d) TEM image, EDX spectrum, and XRD pattern of Pt₄₅Sn₂₅Bi₃₀ nanoplates. The inset in (d) shows the unit cell of intermetallic Pt₅₀Snₐ₂Bi₂₅. In which green, purple, and yellow spheres represent Pt, Sn, and Bi atoms, respectively. (e) Aberration-corrected HAADF-STEM image of a typical hexagonal Pt₄₅Sn₂₅Bi₃₀ nanoplate. (f) High-resolution HAADF-STEM image and the corresponding EDX maps of the representative area in Pt₄₅Sn₂₅Bi₃₀ nanoplate, and the schematic illustration of the atomic arrangement in the nanoplate. Reproduced from ref. 14. Copyright 2019, Wiley-VCH. (g and h) The reaction process of Au₅₆@Pd₄₄DCS, Au₇₁@Pd₂₉DCS, and Au₅₆@Pd₄₄DCS, and TEM image of Au₅₆@Pd₄₄DCS. Reproduced from ref. 13. Copyright 2020, Elsevier.
### Table 1  Pt- and Pd-based catalysts for formic acid electro-oxidation

| Electrocatalyst | Preparation | Electrolyte | Peak potential (vs. RHE) | Peak current density | Current density after stability test | Ref. |
|----------------|-------------|-------------|---------------------------|----------------------|--------------------------------------|-----|
| **Controlling the shape and exposing high-index facets** | | | | | | |
| Pt-M (M = Ni, Co, Cu) (NPs, [210]) | Colloidal synthesis and CVD | 0.5 M H₂SO₄ + 0.5 M HCOOH | ~0.55 V | 13.25 mA cm⁻² (Pt THH NPs) | 0.5 mA cm⁻² (Pt THH NPs) | 5 |
| Pt-M (M = Sb, Bi, Pt, Te), Pd-Bi, Pt/C-Bi (NPs, [210]) | CVD with alloying-dealloying | 0.5 M H₂SO₄ + 0.5 M HCOOH | ~0.55 V | 13.25 mA cm⁻² (Pt-Bi), 1.44 A mg⁻¹ (Pt-Bi) | 0.44 mA cm⁻² (Pt-Bi) (0.5 V, 7200 s) | 7 |
| PtH₂ (nanocrystal) | Treating commercial Pd black | 0.5 M H₂SO₄ + 0.25 M HCOOH | ~0.28 V | 5.12 mA cm⁻² (0.5 V, 3600 s) | 1.06 A mg⁻¹ (0.34 V, 400 s) | 18 |
| **Metals supported on suitable substrates** | | | | | | |
| Pt-Fe₂P/C (NPs-metal phosphide/C) | Microwave-assisted ethylene glycol method | 0.5 M H₂SO₄ + 0.5 M HCOOH | ~0.45 V | 1526 mA mg⁻¹ (0.45 V, 3600 s) | 108.24 mA mg⁻¹ (0.44 V, 7200 s) | 8 |
| Pt/C (NPs-metal oxide) | H₂ thermal reduction | 0.5 M H₂SO₄ + 0.5 M HCOOH | ~0.4 V | 11.38 mA cm⁻² (0.4 V, 3600 s) | 1060 A g⁻¹ (0.44 V, 7200 s) | 11 |
| PdSn/SnO₂/TiC (triple junction structure) | NaBH₄ reduction method | 0.5 M H₂SO₄ + 0.5 M HCOOH | ~0.45 V | 0.23 mA cm⁻² (0.3 V, 3600 s) | 0.029 mA cm⁻² (0.3 V, 3600 s) | 16 |
| Pt/S-C (single atom/meso S-C) | H₂ thermal reduction | 0.5 M H₂SO₄ + 0.5 M HCOOH | ~0.85 V | 3.5 A mg⁻¹ (0.1 A mg⁻¹, 0.4 V, 7200 s) | 2215 mg⁻¹ (0.4 V, 10,000 s) | 83 |
| Pd/BG-CN (NPs/hybrid of B-N-C) | H₂ thermal reduction | 0.5 M H₂SO₄ + 0.5 M HCOOH | ~0.45 V | 31.7 mA cm⁻² (0.44 V, 10,000 s) | 400 A g⁻¹ (0.4 V, 3600 s) | 88 |
| Pd/NRGO@SiO₂ (nanoclusters/N-C@SiO₂) | NaBH₄ reduction method | 0.5 M H₂SO₄ + 0.5 M HCOOH | ~0.36 V | 2.37 A mg⁻¹ (0.1 A mg⁻¹, 0.25 V, 3600 s) | 532.7 mg⁻¹ (0.2 V, 3600 s) | 89 |
| Pd-Mo₂N/rGO (NPs-metal nitride/C) | Assembly-immobilization + NaBH₄ reduction method | 0.5 M H₂SO₄ + 0.5 M HCOOH | ~0.45 V | 0.95 mA mg⁻¹ (89 A mg⁻¹) (3000 s) | 182 mA mg⁻¹ (0.25 V, 3600 s) | 90 |
| **Special nanostructures** | | | | | | |
| Pd (periodically ordered mesoporous membranes) | Template electrodeposition method | 0.5 M H₂SO₄ + 0.5 M HCOOH | ~0.49 V | 3.6 mA cm⁻² (3.34 A mg⁻¹) | 3.65 A mg⁻¹ (3.65 A mg⁻¹) | 6 |
| Pd (nanotube arrays/ITO) | Dual template electrodeposition method | 0.5 M H₂SO₄ + 0.5 M HCOOH | ~0.5 V | 4.41 mA cm⁻² (3.65 A mg⁻¹) | 3.65 A mg⁻¹ (3.65 A mg⁻¹) | 19 |
| Pd/Pt (porous half-shells) | Hydrothermal one-pot synthesis | 0.5 M H₂SO₄ + 0.5 M HCOOH | ~0.07 V | 318 mg⁻¹ (70 mg⁻¹, 0.2 V, 3500 s) | 3.7 A mg⁻¹ (89 A mg⁻¹) (0.95 mg⁻¹) | 22 |
| Pt₃Au₂/MCNTs (nanocluster/C) | One-phase synthesis | 0.1 M HClO₄ + 0.5 M HCOOH | ~0.6 V | 318 mg⁻¹ (70 mg⁻¹, 0.2 V, 3500 s) | 3.7 A mg⁻¹ (89 A mg⁻¹) (0.95 mg⁻¹) | 86 |
| Pd 2D nanoframes (low-dimensional Pd nanocrystals) | Seeded growth method and solvothermal synthesis | 0.5 M H₂SO₄ + 0.5 M HCOOH | ~0.45 V | 258.27 mg⁻¹ (80 mg⁻¹, 3000 s) | 192 A g⁻¹ (0.4 V, 3500 s) | 92 |
| Pd nanosheets (perforated crystalline/amorphous heterostructures) | Carbon oxide reduction method | 0.5 M H₂SO₄ + 0.5 M HCOOH | ~0.46 V | 413.89 A g⁻¹ (192 A g⁻¹) (0.4 V, 3500 s) | 192 A g⁻¹ (0.4 V, 3500 s) | 93 |
| Pd tetrahedrons | One-pot synthesis | 0.5 M H₂SO₄ + 0.5 M HCOOH | ~0.42 V | 413.89 A g⁻¹ (192 A g⁻¹) (0.4 V, 3500 s) | 192 A g⁻¹ (0.4 V, 3500 s) | 93 |
| **Alloys** | | | | | | |
| Pd-M (M = Ru, Pt, Cu, Au, Ag) (NPs) | Successive reduction method | 0.5 M H₂SO₄ + 0.5 M HCOOH | ~0.4 V | 5.12 mA cm⁻² (Pd-Cu/C) | 0.4 mA cm⁻² (Pd-Cu/C) (0.26 V, 2000 s) | 3 |
### 4.2 Non-Pt and non-Pd electrocatalysts for formic acid oxidation

Although Pt and Pd are the best electrocatalysts so far for formic acid oxidation, their high cost and limited reservation largely hinder their wide application in fuel cell catalysis. Therefore, developing Pt- and Pd-free electrocatalysts for FAEO is paramount, crucial, and has become a hot research topic in recent years.

Many studies have found that downsizing metal nanoparticles can bring about excellent catalytic properties that conventional catalysts don’t possess.\(^\text{10,87}\) Li et al. reported iridium single atoms supported on nitrogen-doped carbon (Ir\(_1$/CN) by pyrolyzing a metal precursors@MOF composite at high temperature.\(^\text{31}\) The electrocatalytic performance of the Ir\(_1$/CN for formic acid oxidation was higher than those of commercial Pt/C, Pd/C, Ir/C and Ir\(_1$/Ir\(_n$/CN catalysts with mixed single atoms and nanoparticles. The enhanced catalytic activity and stability stemmed from the more favorable COOH* pathway and the easily activated O–H bond of the generated COOH* on the Ir–N\(_x/ active centers. Meanwhile, the isolated Ir sites, high Ir atom utilization, and the interaction between Ir and the highly electronic conductive CN support were also important factors for the improved catalytic activity. However, Pt\(_1$/CN and Pd\(_1$/CN showed no electrocatalytic activity for the oxidation of formic acid, which broke with the stereotype. In another study, Xiong et al. synthesized atomically dispersed Rh on N-doped carbon (SA-Rh/CN) via a high-temperature pyrolysis of the metal precursors@Zn-MOF composites, while Zn ion sites were substituted by Rh atoms (Fig. 5a–g).\(^\text{84}\) The obtained SA-Rh/CN exhibited promising catalytic properties for formic acid electro-oxidation, but SA-Pd/CN and SA-Pt/CN synthesized by the same procedure and the as-prepared Rh nanoparticles on carbon black (Rh/C) were found to be inert toward FAEO. It was proposed that the SA-Pd/CN and SA-Pt/CN lost their catalytic abilities for FAEO due to their weak bonding ability to the intermediates. The high catalytic activity and stability of SA-Rh/ CN arose from the interaction between the Rh and CN support to form RhN\(_x/ the isolated metal atoms with high metal utilization, and the porous structure with the large surface area. This study also demonstrated that CO is difficult to generate and easy to remove. Meanwhile, the SA-Rh/CN was thermodynamically and kinetically favorable for formic acid oxidation via a formate route, and the C–H cleavage of HCOO* was the rate-determining step.\(^\text{26}\)

Transition metals, such as platinum and palladium, contain unoccupied d orbitals and unpaired d electrons. Therefore, when the catalysts containing transition metals contact with the reactant molecules, a variety of chemical adsorption bonds with various characteristics will be formed, thus achieving the purpose of molecular activation and a reduction of the activation energy for complex reactions. However, to obtain a high

| Electrolyte | Preparation | Electrode | Catalysts | Peak potential (vs. RHE) | Peak current density | Current density a vs. C0 | Reference |
|-------------|-------------|-----------|-----------|-------------------------|---------------------|-------------------------|-----------|
| 0.1 M HClO\(_4/ + 0.1 M HCOOH | Solved growth method and self-assembly process | Au@Pd (core with shell) | PdFe/C (ordered intermetallic) | ~0.47 V | 23.33 A m\(^{-2}/ | 1.71 mA mg\(_{\text{Pd}}/ | 1.5 A mg\(_{\text{Pt}}/ | 75 |
| 0.1 M HCOOH | Sequential complexing–reducing–ordering processes | PdFe/C (nanomeshes) | Pd-Fe-Pt (2D nanomeshes) | ~0.55 V | 14.5 A mg\(_{\text{Pt}}/ | 3.79 mA mg\(_{\text{Pd}}/ | 14.1 A mg\(_{\text{Pt}}/ | 96 |
| 0.1 M HCOOH | Wet chemical reaction | Pt–Au/C (NPs/C) | Pt–Au/C (NPs/C) | ~0.55 V | 9.06 A mg\(_{\text{Pt}}/ | 0.15 A mg\(_{\text{Pt}}/ | 0.015 A mg\(_{\text{Pd}}/ | 95 |
| 0.1 M HClO\(_4/ + 0.1 M HCOOH | Conventional ultrasonic-assisted method | Pt–Au/C (NPs/C) | Pt–Au/C (NPs/C) | ~0.55 V | 3.79 mA mg\(_{\text{Pt}}/ | 0.15 A mg\(_{\text{Pt}}/ | 0.17 mA mg\(_{\text{Pd}}/ | 95 |
| 0.1 M HClO\(_4/ + 0.1 M HCOOH | Conventional ultrasonic-assisted method | Pt–Au/C (NPs/C) | Pt–Au/C (NPs/C) | ~0.55 V | 3.79 mA mg\(_{\text{Pt}}/ | 0.15 A mg\(_{\text{Pt}}/ | 0.17 mA mg\(_{\text{Pd}}/ | 95 |
| 0.1 M HClO\(_4/ + 0.1 M HCOOH | One-pot synthesis method | Pt–Au/C (C–NPs/Pt) | Pt–Au/C (C–NPs/Pt) | ~0.55 V | 14.5 A mg\(_{\text{Pt}}/ | 3.79 mA mg\(_{\text{Pt}}/ | 14.1 A mg\(_{\text{Pt}}/ | 96 |
| 0.1 M HClO\(_4/ + 0.1 M HCOOH | Conventional ultrasonic-assisted method | Pt–Au/C (NPs/C) | Pt–Au/C (NPs/C) | ~0.55 V | 3.79 mA mg\(_{\text{Pt}}/ | 0.15 A mg\(_{\text{Pt}}/ | 0.17 mA mg\(_{\text{Pd}}/ | 95 |
| 0.1 M HClO\(_4/ + 0.1 M HCOOH | One-pot synthesis method | Pt–Au/C (C–NPs/Pt) | Pt–Au/C (C–NPs/Pt) | ~0.55 V | 14.5 A mg\(_{\text{Pt}}/ | 3.79 mA mg\(_{\text{Pt}}/ | 14.1 A mg\(_{\text{Pt}}/ | 96 |
| 0.1 M HClO\(_4/ + 0.1 M HCOOH | Conventional ultrasonic-assisted method | Pt–Au/C (NPs/C) | Pt–Au/C (NPs/C) | ~0.55 V | 3.79 mA mg\(_{\text{Pt}}/ | 0.15 A mg\(_{\text{Pt}}/ | 0.17 mA mg\(_{\text{Pd}}/ | 95 |
| 0.1 M HClO\(_4/ + 0.1 M HCOOH | One-pot synthesis method | Pt–Au/C (C–NPs/Pt) | Pt–Au/C (C–NPs/Pt) | ~0.55 V | 14.5 A mg\(_{\text{Pt}}/ | 3.79 mA mg\(_{\text{Pt}}/ | 14.1 A mg\(_{\text{Pt}}/ | 96 |
| 0.1 M HClO\(_4/ + 0.1 M HCOOH | One-pot synthesis method | Pt–Au/C (C–NPs/Pt) | Pt–Au/C (C–NPs/Pt) | ~0.55 V | 14.5 A mg\(_{\text{Pt}}/ | 3.79 mA mg\(_{\text{Pt}}/ | 14.1 A mg\(_{\text{Pt}}/ | 96 |
| 0.1 M HClO\(_4/ + 0.1 M HCOOH | Conventional ultrasonic-assisted method | Pt–Au/C (NPs/C) | Pt–Au/C (NPs/C) | ~0.55 V | 3.79 mA mg\(_{\text{Pt}}/ | 0.15 A mg\(_{\text{Pt}}/ | 0.17 mA mg\(_{\text{Pd}}/ | 95 |
| 0.1 M HClO\(_4/ + 0.1 M HCOOH | One-pot synthesis method | Pt–Au/C (C–NPs/Pt) | Pt–Au/C (C–NPs/Pt) | ~0.55 V | 14.5 A mg\(_{\text{Pt}}/ | 3.79 mA mg\(_{\text{Pt}}/ | 14.1 A mg\(_{\text{Pt}}/ | 96 |
| 0.1 M HClO\(_4/ + 0.1 M HCOOH | Conventional ultrasonic-assisted method | Pt–Au/C (NPs/C) | Pt–Au/C (NPs/C) | ~0.55 V | 3.79 mA mg\(_{\text{Pt}}/ | 0.15 A mg\(_{\text{Pt}}/ | 0.17 mA mg\(_{\text{Pd}}/ | 95 |
| 0.1 M HClO\(_4/ + 0.1 M HCOOH | Conventional ultrasonic-assisted method | Pt–Au/C (NPs/C) | Pt–Au/C (NPs/C) | ~0.55 V | 3.79 mA mg\(_{\text{Pt}}/ | 0.15 A mg\(_{\text{Pt}}/ | 0.17 mA mg\(_{\text{Pd}}/ | 95 |
| 0.1 M HClO\(_4/ + 0.1 M HCOOH | Conventional ultrasonic-assisted method | Pt–Au/C (NPs/C) | Pt–Au/C (NPs/C) | ~0.55 V | 3.79 mA mg\(_{\text{Pt}}/ | 0.15 A mg\(_{\text{Pt}}/ | 0.17 mA mg\(_{\text{Pd}}/ | 95 |
catalytic performance, the structure and composition of the materials need to be optimized, such as by reducing the size of nanoparticles with high-index facets, selecting suitable supports, and adjusting the pore size, effectively improving the structural stability, efficiently reducing the dissolution of non-precious metals during the electrochemical reactions, and applying the conclusions obtained from the study of model catalysts such as metal single-crystal surfaces to the actual catalyst systems. It should be pointed out that currently there are very few high-performance non-Pt and non-Pd electrocatalysts for FAEO. Therefore, further detailed research and investigations are urgently needed.

5. Conclusions and outlook

As a typical electrocatalytic reaction in liquid fuel cells, studies on formic acid electro-oxidation have attracted great attention, including the reaction mechanism and designing high-performance electrocatalysts. However, many problems and challenges still need to be resolved before formic acid electro-oxidation systems offer sufficient performance to satisfy basic research and practical application demands. The details are as follows: (1) to reveal the reaction mechanism and identify the active intermediates are still challenging. In addition to the traditional electrochemical methods and density functional theory (DFT) calculations, developing high-resolution and ultrafast in situ or operando spectroscopic techniques is necessary to detect the intermediates and analyze the reaction mechanism of formic acid oxidation. (2) Based on the reaction mechanism, rationally designing and adjusting the structures of electrocatalysts are an ideal strategy to explore high-performance catalysts for formic acid oxidation. Both the thermodynamics and kinetics should be taken into account to improve the current density and to reduce the overpotential of formic acid electro-oxidation. In addition, investigations on effective non-Pt and non-Pd electrocatalysts are significantly
important to explore cost-efficient anode catalysts for fuel cells. (3) Finally, the costs, durability, and activity should be balanced reasonably so as to realize the commercialization of direct formic acid fuel cells.

Conflicts of interest

There are no conflicts to declare.

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