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Highly Dispersed Pt Nanoparticles Root in Single-Atom Fe Sites in LDHs toward Efficient Methanol Oxidation

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Abstract

Active and durable electrocatalysts for methanol oxidation reaction are of critical importance to the commercial viability of direct methanol fuel cell, which has already attracted growing popularities. However, current methanol oxidation electrocatalysts fall far short of expectations and suffer from excessive use of noble metal, mediocre activity, and rapid decay. Here we report the Pt anchored on NiFe-LDHs surface hybrid for stable methanol oxidation in alkaline media. Based on the high intrinsic methanol oxidation activity of Pt nanoparticles, the substrates NiFe-LDHs further enhanced anti-poisoning ability and maintained unaffected stability after 200,000 s cycle test compared to commercial Pt/C catalyst. The use of NiFe-LDHs is believed to play the decisive role to evenly disperse Pt nanoparticles on their surface using single atomic dispersed Fe as anchoring sites, making full use of abundant OH groups and subsequent facilitating the oxidative removal of carbonaceous poison on neighboring Pt sites. This work highlights the specialty of NiFe-LDHs in improving the overall efficiency of methanol oxidation reaction.

Keywords: Layered double hydroxides; Methanol oxidation; Single-atom; Pt electrocatalysts

1. Introduction

Growing concerns regarding the climate change and regional energy disputes have arouse attention worldwide in finding renewable and clean energy to ensure the sustainable development of society [1,2]. Direct methanol fuel cells (DMFCs) have long been considered as a promising power conversion tool by virtue of their higher energy-conversion rate, wide working temperature range, no pollution emissions, portability and safety of raw materials [3–6]. However, it is still challenging to realize the mass commercialization, mainly originating from high overpotential at an anode (where methanol is oxidized, requiring hundreds of mV as

overpotential [7–9]), unaffordable cost associated with precious metal electrocatalysts (for example, Pt) and low operation durability of the anode operation [10–12] (carbonaceous species poisoning). In consideration of high cost and scarcity, exposing more accessible surface area by reducing size of Pt is an essential and a direct approach for achieving the commercialization of DMFCs, however, the limited durability is always observed due to the aggregation of nanoparticles (NPs) or the poisoning of noble metals [13–16]. Hence, desirable supporting materials with large surface area that can anchor and disperse the Pt NPs are necessary to be developed and optimized. In recent years, many materials have been utilized

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as supporting materials for loading of Pt-based nanocatalysts, including metal oxides and nanocarbon materials [11,17–19], while only limited improvements in durability have been achieved so far, presumably due to the limited capability of the supports to assist in the oxidative removal of CO on adjacent Pt sites, and the weak interaction between Pt and supports fails preventing the aggregation of Pt during methanol oxidation [20–22].

In seeking optimal supports, which can anchor and stabilize the noble metal NPs, and facilitate the removal of CO during methanol oxidation on noble metal NPs, two-dimensional metal hydroxides have been intensively investigated. On the one hand, the metal hydroxides possess high specific surface area and rich sites for noble metal anchoring, on the other hand, the metal hydroxides shall supply abundant OH during methanol oxidation to benefit the oxidation of CO [23–25]. For example, Li and Cai et al. successively reported the use of Ni(OH)₂ to disperse Pt and Pd NPs for methanol oxidation, showing enlarged electrochemical surface area and excellent working stability, which have proven to improve the overall methanol oxidation efficiency [11]. While the combination of noble metal with metal hydroxides is attracting and proving effective in methanol oxidation, further advancements such as facilitating the generation of OH_{ads} to improve the intrinsic activity and selectivity, achieving high dispersion of noble metals via strong support-metal interaction, are still vacant based on unitary metal hydroxides. Layered double hydroxides (LDHs), in which partial M²⁺ ions in laminates are replaced by M³⁺ ions via isomorphous substitution, while all the M³⁺ will be isolated from each other by electrostatic repulsion force, have shown potential in addressing above problems [26,27]. Firstly, fast generation of OH_{ads} which can be used to oxidize adsorbed intermediates of methanol oxidation, on LDHs can be facilitated via compositional modulation, this process has already shown in oxygen evolution reaction [28–31]. Secondly, the partial substitution of M²⁺ by M³⁺ results in unbalanced charge configuration, which will directionally adsorb noble metal precursors to specific sites and form strong electrostatic attraction interaction between M³⁺–O(H)–Pt [32], resulting in the uniformly dispersed and stable decoration of anchored noble metal. Thirdly, OH[–] transfer is favorable on LDHs with large surface area (generally over 120 m²·g^{–1}), which can relieve the local pH evolution around the surface of electrode and protect the electrode from reconstruction, especially at high current density [32,33].

In this work, Pt NPs with acknowledged methanol oxidation activity and NiFe-LDHs with excellent OH_{ads} generation capability are combined together by a two-step synthetic process, in which the single-atomic dispersed Fe³⁺ ions in LDHs laminates serve as anchoring sites for Pt precursors, ensuring the highly dispersion of Pt. Electrochemical characterizations for activity, selectivity and stability jointly confirm the positive effect from NiFe-LDHs in facilitating methanol oxidation, avoiding CO poison or Pt aggregation by showing the onset potential at –0.64 V vs. SCE and no obvious decay after the cycle test for 200,000 s. This work shall inspire the deep understanding in characteristic of LDHs, and optimization of LDHs with noble metal NPs toward efficient methanol oxidation.

2. Results and discussion

The Pt/NiFe-LDHs electrocatalysts were synthesized by an electrostatic interaction-based self-assembly strategy, which is illustrated in Fig. 1. Firstly, the NiFe-LDHs nanosheets were synthesized by co-precipitation method. After adding H₂PtCl₆ into LDHs suspensions (LDHs nanosheets were dispersed and stirred for over 72 h), PtCl₆^{2–}/LDHs composite was formed as a result of the electrostatic attraction between PtCl₆^{2–} and LDHs nanosheets, as shown in Fig. 1. Finally, the addition of sodium borohydride reduced the PtCl₆^{2–}/NiFe-LDHs hybrids to Pt/NiFe-LDHs hybrids [34]. A low concentration of H⁺ plays two roles: firstly, H⁺ can react with surface OH groups to open unsaturated coordinated Fe sites for PtCl₆^{2–} adsorption; secondly, H⁺ would controllably etch (by the modulation of H₂PtCl₆ addition) LDHs nanosheets to induce defective metal hydroxides nanostructure in the Pt/NiFe-LDHs hybrids. These defects are also believed to play a decisive role in promoting the transformation from adsorbed OH[–] species to OH_{ads} at applied potentials and subsequent oxidative removal of carbonaceous poison on neighboring Pt sites [35,36].

The morphology and nanostructure of NiFe-LDHs were characterized by scanning electron microscopy (SEM, Figure S1) and transmission electron microscopy (TEM, Figure S2). It can be observed that the NiFe-LDHs have a flake-like morphology with lateral sizes in a range of 50–100 nm. XRD patterns of NiFe-LDHs are shown in Figure S3. The diffraction peaks at 2θ = 11.4°, 23.0°, 34.4°, 39.0°, 46.0°, 59.9° and 61.3° are attributed to the (003), (006), (012), (015), (018), (110) and (113) planes, which are in accordance with that reported in the earlier literatures [37].

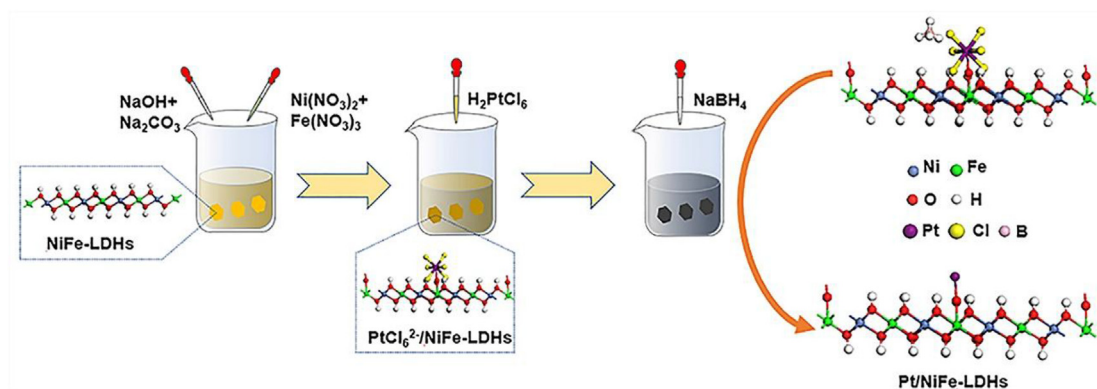


Fig. 1. Schematic illustration showing the preparation process of the Pt/NiFe-LDHs.

Fig. 2a and b shows the TEM images of Pt/NiFe-LDHs and commercial Pt/C, respectively. Notably, the Pt NPs loaded on NiFe-LDHs nanosheets show high uniformity and smaller particle size compared with the Pt on carbon support, highlighting that the electrostatic interaction between Pt NPs and single-atomic dispersed Fe^{3+} in LDHs laminates makes a real difference in dispersing the Pt NPs, and by the first-principles calculations (Figure S4) we confirm that the adsorption energy of Pt on Fe^{3+} sites ($\text{Pt}-\text{O}-\text{Fe}^{3+}$) is lowered than that on Ni^{2+} sites ($\text{Pt}-\text{O}-\text{Ni}^{2+}$). The high-resolution TEM (HR-TEM) images exhibit an interplanar spacing of 0.22 nm and 0.25 nm for the (111) plane of the Pt NPs and (012) plane of the LDHs,

respectively (Fig. 2c). The XRD patterns of Pt/NiFe-LDHs electrocatalyst are presented in Fig. 2d and S5, and the diffraction peaks locate at 2θ values of 39.8° , 46.2° and 67.5° , corresponding to (111), (200) and (220) planes, respectively, of a pure face-centered cubic structure of Pt [38]. Similar LDHs diffraction peaks prove that the crystalline structure of NiFe-LDHs has not been altered upon Ni:Fe ratios or the deposition of Pt NPs, which can also be observed from the SEM images in Figure S6, confirming the successful synthesis of the Pt/NiFe-LDHs electrocatalysts.

Cyclic voltammetry (CV) was used to study the electrochemical performance of the electrocatalysts in MOR. Data in Fig. 3a display an intense anodic

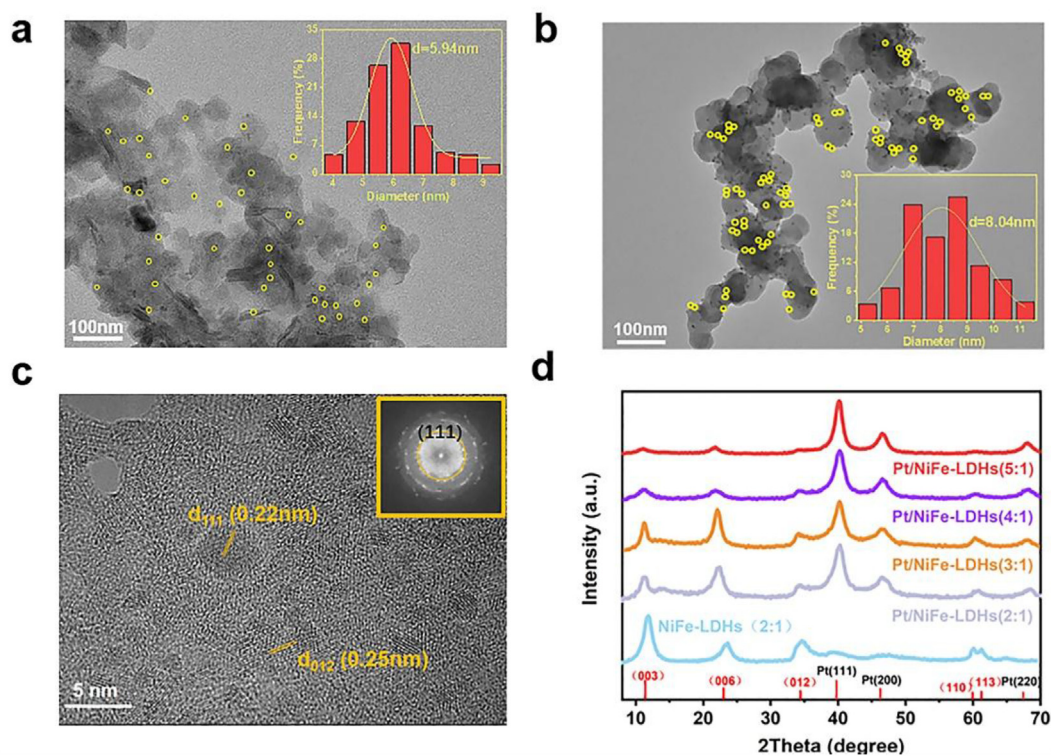


Fig. 2. TEM images of (a) the Pt/NiFe-LDHs and (b) the commercial Pt/C, (c) HRTEM image and (d) XRD patterns of the Pt/NiFe-LDHs.

peak in the forward scan and a minor anodic peak in the reverse scan. The anodic peak in the forward scan is ascribed to the oxidation of methanol to CO_2 , while another one in the reverse scan is ascribed to the oxidized removal of the CO and other carbonaceous species generated on Pt in the forward scan [39]. In addition, the ratio of forward-to-backward oxidation current, I_f/I_b , is an important parameter evaluating the electrocatalyst tolerance to CO and other poisoning carbonaceous species. A higher value demonstrates that the oxidative removal of the poisoning carbonaceous species on the electrocatalyst surface is more favorable. Fig. 3a and b study and compare the methanol oxidation reaction (MOR) activity and selectivity of Pt/NiFe-LDHs with different Ni:Fe ratios, from which a conclusion can be brought that the Pt/Ni₃Fe₁-LDHs have the best electrocatalytic ability on MOR according to the exceptional current (3.41 mA), high I_f/I_b (2.84) and low onset potential (-0.64 V vs. SCE). Since OH_{ads} species are both actively involved in MOR and oxygen evolution, the results in this work agree with previous research that Ni₃Fe₁-LDHs have advantages in generating OH_{ads} [26], which are helpful in facilitating the oxidative removal of the poisoning

carbonaceous species. The proposed MOR mechanism on Pt/NiFe-LDHs is shown in Figure S7. Firstly, atomic-dispersed Fe sites in LDHs help to disperse anchored Pt nanoparticles, resulting in the more exposed electrochemical surface area of Pt. Secondly, the oxidation of methanol can always lead to the tight adsorption of CO on the surface of Pt, while the OH_{ads} as generated on the surface of LDHs would facilitate the fast removal of CO species, resulting in long-term durability. CO stripping experiments were also used to probe the strong resistance of Pt/NiFe-LDHs to CO poisoning. As shown in Figure S8, the CO oxidation peak of Pt/NiFe-LDHs has a negative shift compared to that of Pt/C, and the integrated area for CO oxidation is also smaller on Pt/NiFe-LDHs than on Pt/C, indicating that the adsorption of CO on Pt/NiFe-LDHs is weaker than that on Pt/C.

Fig. 3c and d further study the influence of loading Pt contents on the MOR performance of the hybrid electrocatalysts. Obviously, Pt(16%)/Ni₃Fe₁-LDHs exhibited an optimal performance with the highest current of about 15 mA, locating at the apex of Pt content-MOR activity volcano plot. At the beginning of the MOR, the methanol will be adsorbed on Pt as the active site, so the increased

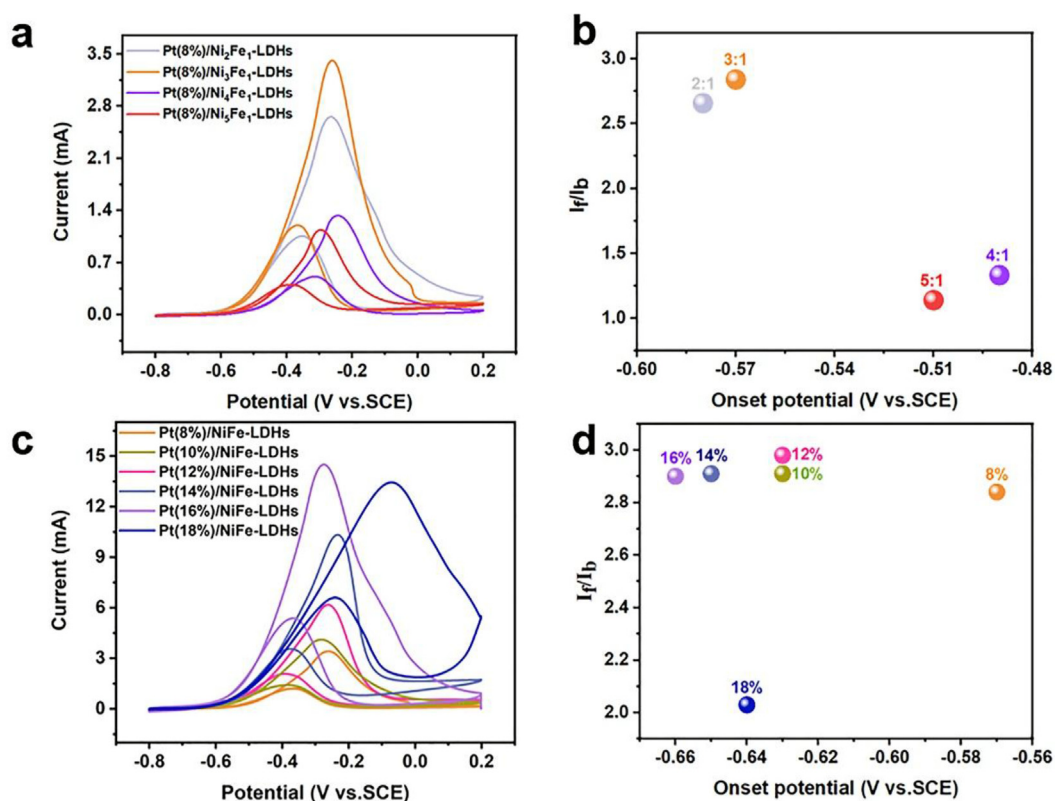


Fig. 3. (a) CV curves, (b) I_f/I_b and onset potential of Pt(8%)/NiFe-LDHs at different Ni:Fe ratios measured in N_2 -saturated $1 \text{ mol}\cdot\text{L}^{-1}$ CH_3OH solution containing $1 \text{ mol}\cdot\text{L}^{-1}$ KOH at a scan rate of $50 \text{ mV}\cdot\text{s}^{-1}$, (c) CV curves, (d) I_f/I_b and onset potential of Pt/Ni₃Fe₁-LDHs loaded by different amounts of H_2PtCl_6 measured in N_2 -saturated $1 \text{ mol}\cdot\text{L}^{-1}$ CH_3OH solution containing $1 \text{ mol}\cdot\text{L}^{-1}$ KOH at a scan rate of $50 \text{ mV}\cdot\text{s}^{-1}$.

content of Pt will be favor for the adsorption of methanol, resulting the growing MOR current, however when the content of Pt is too high, the aggregation of Pt NPs is inevitable, thereby reducing its active specific surface area and leading to excessive adsorption of carbonaceous by-products during MOR, finally causing the declines of MOR activity and selectivity. Therefore, the MOR activity of the Pt(16%)/Ni₃Fe₁-LDHs stands out among all composites.

Lately, the Pt valence state, which is tightly related to the adsorption energies of methanol and CO intermediates on Pt during MOR, was studied by X-ray photoelectron spectroscopy (XPS) in Figure S9. Firstly, the Pt 4f_{7/2} binding energies of the Pt/NiFe-LDHs are down-shifted compared to that of commercial Pt/C, which indicates the change in the electron density of Pt due to electrostatic interaction in Pt–O–Fe³⁺. Furthermore, the ratio between Pt²⁺/Pt⁰ decreases in Pt/NiFe-LDHs compared with that in Pt/C, verifying the drop of Pt valence state in Pt/NiFe-LDHs. The increase of electro density of Pt can possibly weaken the adsorption of reaction intermediates, here is CO, on the electrocatalyst surface, and in turn enhance the selectivity of MOR [40–43]. The high-resolution Ni 2p spectrum (Figure S10a) shows two peaks with binding energies at 856.4 and 874.1 eV correspond to the Ni²⁺ 2p_{3/2} and Ni²⁺ 2p_{1/2}, respectively, which is in good agreement with a previous report [44]. The Fe 2p signal is composed of two peaks (Figure S10b), belonging to Fe 2p_{1/2} and Fe 2p_{3/2} binding energies of the Fe oxidation state (Fe³⁺) [45]. Compared the valence states of Ni and Fe between the NiFe-LDHs and Pt/NiFe-LDHs, we could find that there is no difference between the nickel element, while the binding energy of iron decreases slightly from 712.95 eV to 712.6 eV after Pt anchoring. This result can be understood as since Pt⁴⁺ is preferentially reduced by the sodium borohydride [46], the adjacent Fe³⁺ is also partially reduced, further confirming the strong interaction between Pt and Fe by establishing Pt–O–Fe³⁺ bond on NiFe-LDHs surface.

The mass activity of Pt-based catalysts is usually used for evaluating the MOR catalytic properties, which is calculated by the peak current per milligram Pt. According to Fig. 4a, the mass activity of the Pt(16%)/Ni₃Fe₁-LDHs toward MOR is as high as 999 mA·mg_{Pt}⁻¹, which is 1.23 times higher than that of commercial Pt/C (781 mA·mg_{Pt}⁻¹) and 1.22 times higher than that of benchmark Pt/Ni(OH)₂ (820 mA·mg_{Pt}⁻¹). Then, we performed 1200 cycles CVs to investigate the frequent switch on/off durability of the electrocatalyst during MOR. Also exhibited in Fig. 4a, the peak current density

degenerates with prolonged cycling time. This phenomenon could be ascribed to the continuous generation of intermediates and their poisoning impacts on electrocatalyst during MOR, also including the detachment of Pt NPs from the substrate or the aggregation of Pt NPs. The peak current densities for commercial Pt/C, Pt/Ni(OH)₂ and Pt(16%)/Ni₃Fe₁-LDHs after the 1200th cycle were about 48.78 mA·mg_{Pt}⁻¹, 350.25 mA·mg_{Pt}⁻¹, 608 mA·mg_{Pt}⁻¹, respectively, which is 6.2%, 42.7% and 60% of the first cycle, respectively, highlighting the positive effect of NiFe-LDHs in facilitating the oxidation of methanol, accelerating the removal of carbonaceous by-products, and anchoring the Pt NPs against aggregation.

The continuous working stability is also an important index to evaluate the catalytic performance of MOR electrocatalysts. By using amperometric *i-t* method, we firstly characterized the short-term stability of electrocatalyst. As shown in Fig. 4b, three electrocatalysts display a current density decline in the initial period, perhaps resulting from electrocatalyst poisoning by chemisorbed carbonaceous species formed during the methanol oxidation. At the end of stability test, the current density of Pt(16%)/Ni₃Fe₁-LDHs electrocatalyst was higher than those of commercial Pt/C and Pt/Ni(OH)₂, further confirming the excellent working stability of Pt(16%)/Ni₃Fe₁-LDHs.

Then the long-term operation durability of these two electrocatalysts were characterized and are compared. At the end of continuous operation for 50,000 s, negligible activities (almost no MOR working current densities) maintained for commercial Pt/C (Figure S11). In stark contrast, Pt(16%)/Ni₃Fe₁-LDHs still delivered a current density of 145 mA·mg_{Pt}⁻¹, accounting for 33% of initial working current density. To our surprise, we found that Pt(16%)/Ni₃Fe₁-LDHs could still exhibit the initial MOR activity in a fresh electrolyte after working for 50,000 s, while Pt/C could not recover to the initial activity (merely recovered to around 30% as shown in Figure S11). We repeated the cycling up to four times for a total of 200,000 s, and the full activity was recovered every time (Figure S11). So, the drop of the MOR working current density of Pt(16%)/Ni₃Fe₁-LDHs may be mainly induced by the changes of pH and composition of electrolyte (generated CO₂ being dissolved in KOH, resulting in the formation of K₂CO₃ and drop of electrolyte pH), rather than the severe reconstruction or the aggregation of Pt NPs in Pt(16%)/Ni₃Fe₁-LDHs.

To supply experimental evidence for the greatly enhanced durability of the Pt(16%)/Ni₃Fe₁-LDHs as compared to that of the commercial Pt/C in MOR, we further monitored their morphological

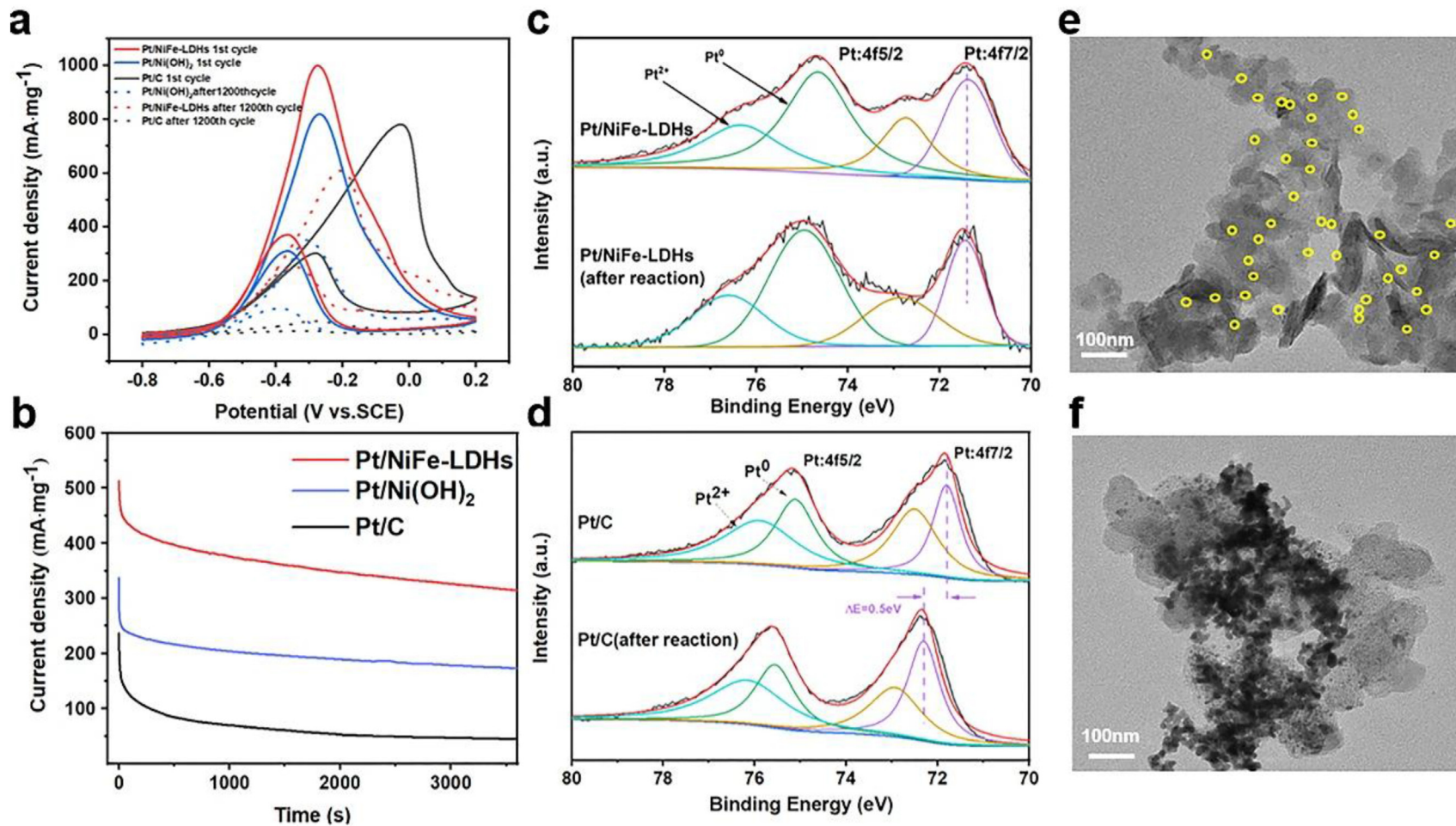


Fig. 4. (a) CV and (b) chronoamperometric (CA) curves of Pt/NiFe-LDHs, Pt/C and Pt/Ni(OH)₂ at different cycle numbers measured in N₂-saturated 1 mol·L⁻¹ CH₃OH solution containing 1 mol·L⁻¹ KOH at a scan rate of 50 mV·s⁻¹; Pt 4f XPS spectra of Pt/NiFe-LDHs, (c) and Pt/C, (d) after MOR reaction; TEM images of Pt/NiFe-LDHs (e) and Pt/C (f) after MOR reaction.

and compositional changes after the durability test. XPS spectra were used to monitor the evolution of the element composition and electronic states of Pt(16%)/Ni₃Fe₁-LDHs and Pt/C during the long-term stability test. The results indicate that the electronic states of Ni and Fe in Pt(16%)/Ni₃Fe₁-LDHs were well maintained (Figure S12), also negligible shifts of Pt binding energy can be observed in Fig. 4c. In contrast, the Pt binding energies of the commercial Pt/C were up-shifted 0.5 eV after reaction (Fig. 4d). The Pt²⁺/Pt⁰ ratio as compared in Fig. 4c exhibits that no obvious Pt oxidation could be observed in Pt(16%)/Ni₃Fe₁-LDHs, while a growing value of Pt²⁺/Pt⁰ ratio could be detected in Pt/C, indicating the transformation from Pt⁰ to Pt²⁺ during MOR, so it is reasonable that the activity of Pt/C decayed after prolonged working hours. Moreover, it is found that the Pt NPs in Pt(16%)/Ni₃Fe₁-LDHs maintained their morphology and dispersion well during the 200,000 s stability test (Fig. 4e). In the case of the commercial Pt/C, an obvious aggregation and ripening of Pt NPs can be observed (Fig. 4f).

The interactions between Pt NPs and NiFe-LDHs have merits in providing sufficient OH_{ads} for the oxidation of methanol and carbonaceous by-products, requiring low Pt loading by maximizing the available surface area of Pt NPs, anchoring Pt NPs via Fe³⁺-O-Pt bond against Pt aggregation during MOR procedure. In real DMFC devices, it would be highly desirable for MOR electrocatalysts with low Pt loading, high activity, and reactivated during operation breaks by simply replacing the electrolyte by fresh ones, the electrocatalyst reported in this work suits well to the requirements.

3. Conclusions

In summary, uniform and small Pt nanoparticles were loaded on the NiFe-LDHs nanosheets and used as a high-efficient MOR electrocatalyst. The presence of NiFe-LDHs dramatically promoted the electrocatalytic performance of Pt for the MOR in an alkaline solution by evenly dispersing Pt nanoparticles on their surface via Fe-O(H)-Pt bonding, promoting the generation of OH_{ads} and subsequent oxidative removal of carbonaceous poison on neighboring Pt sites. Through the synergistic effect between Pt and NiFe-LDHs, our electrocatalyst retained an exceptional mass activity of 145 mA·mg⁻¹ even after 50,000 s chronoamperometric measurement, and it could rework functionally in replaced fresh electrolytes. Finally, the resulting Pt/NiFe-LDHs electrocatalyst exhibited higher electrocatalytic activity (onset potential at -0.64 V vs. SCE) and better anti-poisoning ability comparing to commercial Pt/C

catalyst. This study implies that the Pt/NiFe-LDHs electrocatalyst has great potential applications in methanol fuel cell, which are essential for the clean development of society.

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层状金属氢氧化物中铁位点辅助分散铂纳米颗粒用于 高效甲醇氧化

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摘要

高活性和耐用性的甲醇氧化电催化剂对于直接甲醇燃料电池的商业可行性至关重要, 然而, 目前的甲醇氧化电催化剂与预期相去甚远, 存在贵金属用量过多、活性平庸、衰减快等问题。在这里, 我们报告了锚定在镍铁层状金属氢氧化物 (NiFe-LDHs) 表面 Pt 纳米颗粒复合材料, 用于在碱性介质中稳定电催化甲醇氧化。基于 Pt 纳米颗粒的高固有甲醇氧化活性, 与商业 Pt/C 催化剂相比, 基底材料 NiFe-LDHs 在 200,000 s 循环测试后进一步增强了 Pt 的抗中毒能力和稳定性。NiFe-LDHs 层板上单原子分散的 Fe 作为锚定位点将 Pt 纳米颗粒均匀分散在其表面, 进一步充分利用了层状金属氢氧化物表面丰富的 OH 基团, 促进邻近 Pt 位点上毒化中间体的氧化去除。这项工作突出了 NiFe-LDH 在提高甲醇氧化反应整体效率方面的特殊性, 为其他甲醇氧化电催化的设计和应用提供了指导。

关键词: 层状金属氢氧化物; 甲醇氧化; 单原子; Pt 电催化剂