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Facile synthesis of 3D Pd-P nanoparticle networks with enhanced electrocatalytic performance towards formic acid electrooxidation

Jingfang Zhang, You Xu and Bin Zhang*

3D Pd-P nanoparticle networks (NNs) have been successfully synthesized using a facile one-step soft-template-assisted method. The as-prepared Pd-P NNs exhibit markedly improved activity and stability towards formic acid electrooxidation over Pd NNs, commercial Pd/C and Pd-P nanoparticle aggregates (NAs).

Direct formic acid fuel cells (DFAFCs) have attracted considerable research interest as a new generation of environment-friendly power source due to their high energy density, limited fuel crossover, high electromotive force and low operating temperature. The rapid development of DFAFCs in the past decade has stimulated the intensive research in the design and synthesis of excellent electrocatalysts for formic acid electro-oxidation reaction (FAER). Pd-based composites are considered as effective electrocatalysts towards FAER for their lower cost and higher activity compared to Pt-based catalysts. However, pure Pd catalysts suffer from a fatal drawback of slow deactivation during FAER due to either the oxidation of Pd surface or the catalyst poisoning. Thus, it is desirable to develop new Pd-based electrocatalysts with higher activity and extended durability. Recent studies have indicated the doped non-metal element P with abundant valence electrons can improve the electrocatalytic activities and CO tolerances effectively. However, the development of synthetic methodology to control the morphology and structure of such P doped Pd-based nanostructures, which are linked to the properties with a greater versatility, imposes important challenges. In particular, metal-based nanoparticle networks (NNs) are considered to be promising electrocatalysts candidates because the combination of net-like feature and nanoparticle blocks can achieve both the construction of self-supported electrocatalysts and the retain of nanoparticle’s high activity. Moreover, the NNs can not only increase the active area accessible to the reactant molecules, but also provide effective electron mobility in the solid ligaments as well as efficient mass transfer in the catalytic reactions. Furthermore, abundant defect sites and low-coordination atoms on the nanoparticle surfaces are favorable active sites for organic small molecule electro-oxidation.

Herein, we reported a facile and efficient one-step soft-template-assisted strategy to prepare 3D Pd-P alloy NNs with sub-10 nm nanoparticles as building blocks. The as-prepared Pd-P NNs are found to exhibit excellent electrocatalytic activity and stability towards FAER.

The 3D Pd-P alloy NNs were prepared through the hydrothermal reaction of Na$_2$PdCl$_4$ and NaH$_2$PO$_2$ in the presence of HCHO as reducing agent and Brij 58 as soft template (see experimental details in ESI†). The as-obtained samples were firstly characterized by using scanning electron microscopy (SEM). As shown in Fig. 1a, the samples with 3D NNs could be prepared on a large scale. The high-magnification SEM image (Fig. 1b) further confirms the network-like feature composed of many nanoparticles. The energy dispersive X-ray (EDX)
spectroscopy (Fig. 1c) also indicates that the as-prepared networks are composed of Pd and P with the atomic ratio of 7.5:1 (Pd to P). In the absence of NaH$_2$PO$_2$ with other conditions remain unchanged, the as-produced samples are pure Pd NNs (ESI, Fig. S1†). X-ray diffraction (XRD) patterns in Fig. 1d show that both Pd-P NNs and Pd NNs exhibit diffraction peaks of a face centered cubic crystal structure (JCPDS 05-0681). In comparison to pure Pd NNs, it is found that the corresponding diffraction peaks of Pd-P NNs shift to the higher 2θ degrees, suggesting the lattice contraction due to the partial substitution of Pd by P. In addition, the diffraction peaks of the Pd-P NNs are much broader than those of the Pd NNs, showing that the average particle size of Pd-P NNs is much smaller than that of Pd NNs. According to the Scherrer formula$^{4}$, the average sizes of Pd-P and Pd NNs are calculated to be 3.3 nm and 7.7 nm, respectively. In other words, the addition of P reduces the particle size of Pd-P NNs.

The surface composition of Pd-P NNs was further analyzed by X-ray photoelectron spectroscopy (XPS) (ESI, Fig. S3† and Fig. 3). In the P 2p spectrum of Pd-P NNs (Fig. 3a), the P 2p peaks at 129.9 and 133.1 eV are assigned to elemental state phosphorus and oxidized phosphorus, respectively. In the Pd 3d spectrum of Pd-P NNs (Fig. 3b), the Pd 3d signal of Pd-P NNs is fitted to two pairs of doublets with a spin-orbit separation of 5.3 eV: Pd 3d$_{5/2}$ (341.2 eV), Pd 3d$_{3/2}$ (335.9 eV) and Pd 3d$_{5/2}$ (342.2 eV), Pd 3d$_{3/2}$ (336.9 eV), which can be assigned to Pd$^0$ and Pd$^{II}$, respectively. The energy of Pd$^0$ and Pd$^{II}$ in Pd-P NNs positively shift ca. 0.4 eV compared to that of pure Pd.$^5$ This may be due to the fact that the interacting role of Pd with P makes positively charged Pd donate electrons and thus it leads to a decrease of the 3d electron density of Pd atom in Pd-P NNs.$^5$

Fig. 3 XPS spectra of P 2p (a) and Pd 3d (b) of Pd-P NNs.

Fig. 2 (a) and (b) TEM images of Pd-P NNs. Insert in (a): SAED pattern. (c) HRTEM image of Pd-P NNs. The areas marked in Fig. 2c1-2 are many defects with vacancy (by red circles), dislocation (by red squares), as well as several atomic steps and corners (by red lines). (d) Representative TEM image of Pd-P NNs and corresponding EDX mapping images of Pd (red) and P (green) elements.
which is about 1.39 and 3.62 times than that of Pd NNs (2.32 mA cm$^{-2}$) and Pd/C (0.89 mA cm$^{-2}$). Meanwhile, the corresponding mass activity of three catalysts were also tested (ESI, Fig. S8a and Table S1†). This indicates that Pd-P NNs exhibit markedly enhanced electrocatalytic activity toward FAER. The electrochemical stability of Pd-P NNs was studied by chronoamperometry at 0.1 V (Fig. 4c; ESI, Fig. S8b†). It is found that the Pd-P NNs exhibit a slower current decay and much higher current densities than Pd NNs and Pd/C catalysts, indicating a higher tolerance to the intermediate poisoning species generated from FAER, thus better catalytic ability of Pd-P NNs toward FAER. In addition, the electrochemical durability of the Pd-P NNs was evaluated. As shown in Fig. S9 (ESI†), the CV measurements show a loss of 12.5% for Pd-P NNs and 45.0% for Pd/C in ECSAs after 1000 potential cycles, revealing that the durability of Pd-P NNs is also better than the Pd/C catalyst. While serious aggregation was observed for Pd/C, the morphology of the Pd-P NNs was still largely maintained after durability tests. Meanwhile, the sintering-induced mass activity loss was studied (ESI, Fig. S10†). The activities Pd-P samples with different atomic ratios are compared (ESI, Fig. S11†). The optimized atomic ratio is 7.5:1 (Pd to P). CO stripping measurements were applied to exhibit the antipoisoning ability of catalysts. In Fig. 4d and Fig. S12 (ESI†), the peak potential (0.69 V) of CO oxidation on the Pd-P NNs is obviously more negative than that of the Pd NNs (0.73 V) and commercial Pd/C (0.74 V). XPS results suggest that the incorporation of P into Pd lowers the 3d electron density of Pd and makes formic acid be easily oxidized through the direct pathway, which partly accounts for the high catalytic activity. Furthermore, the advantages of 3D Pd-P NNs over Pd-P nanoparticles aggregates (named Pd-P NAs here; ESI, Fig. S7d and S13†) can be confirmed on the basis of a series of electrochemical measurements (ESI, Fig. S14†). The catalytic enhancement of Pd-P NNs may be attributed to the exposure of their intrinsic active sites and larger ECSA provided by 3D networks, as well as the synergetic effects between Pd and P.

In summary, we have demonstrated a facile one-step soft-template-assisted synthesis of 3D Pd-P NNs with sub-10 nm nanoparticles as building blocks. It is found that the as-prepared Pd-P NNs exhibit markedly improved activity and stability towards FAER over Pd NNs, commercial Pd/C and Pd-P NAs, which may be associated with the unique 3D networks and the electronic effects between Pd and P. Importantly, this work will provide a new synthetic strategy to construct other P-doped multimetallic networks, which might be promising candidates for advanced catalytic applications.

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Notes and references
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