<table>
<thead>
<tr>
<th><strong>Title</strong></th>
<th>Triangular Ag–Pd alloy nanoprisms : rational synthesis with high-efficiency for electrocatalytic oxygen reduction</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Author(s)</strong></td>
<td>Xu, Lin; Luo, Zhimin; Fan, Zhanxi; Zhang, Xiao; Tan, Chaoliang; Li, Hai; Zhang, Hua; Xue, Can</td>
</tr>
<tr>
<td><strong>Date</strong></td>
<td>2014</td>
</tr>
<tr>
<td><strong>URL</strong></td>
<td><a href="http://hdl.handle.net/10220/24203">http://hdl.handle.net/10220/24203</a></td>
</tr>
<tr>
<td><strong>Rights</strong></td>
<td>© 2014 The Royal Society of Chemistry. This is the author created version of a work that has been peer reviewed and accepted for publication in Nanoscale, published by Royal Society of Chemistry. It incorporates referee’s comments but changes resulting from the publishing process, such as copyediting, structural formatting, may not be reflected in this document. The published version is available at: [Article DOI: <a href="http://dx.doi.org/10.1039/C4NR03600J">http://dx.doi.org/10.1039/C4NR03600J</a>].</td>
</tr>
</tbody>
</table>
Triangular Ag-Pd alloy Nanoprisms: Rational Synthesis with High-Efficiency for Electrocatalytic Oxygen Reduction

Lin Xu, Zhimin Luo, Zhanxi Fan, Xiao Zhang, Chaoliang Tan, Hai Li, Hua Zhang, and Can Xue*

We report the generation of triangular Ag-Pd alloy nanoprisms through a rationally designed synthetic strategy based on silver nanoprisms as sacrificial templates. The galvanic replacement between Ag nanoprism and H₂PdCl₄ along with co-reduction of Ag⁺/Pd²⁺ is responsible for the formation of final prismatic Ag-Pd alloy nanostructures. Significantly, these Ag-Pd alloy nanoprisms exhibited superior electrocatalytic activity for oxygen reduction reaction (ORR) as compared with the commercial Pd/C catalyst. Such a high catalytic activity is attributed to not only the alloyed Ag-Pd composition but also the dominant {111} facets of the triangular Ag-Pd nanoprisms. This work demonstrates the rational design of bimetallic alloy nanostructures with control of selective crystal facets that are critical to achieve high catalytic activity for fuel cell systems.

Introduction

Bimetallic nanostructures have received considerable attentions recently due to their unique optical, magnetic, and catalytic properties, which are not accessible by their monometallic counterparts. In particular for noble metals, such as Pd, Pt and Au, alloying a principal scarce noble metal with a secondary abundant metal can not only decrease the usage of the scarce noble metal, but also allow for tuning electronic states of the entire structure, which often exhibits superior catalytic properties, including activity, selectivity and durability, when compared with the monometallic counterpart. Considering that the catalytic performance of metal nanostructures is highly dependent on their size, shape, exposed facets and compositions, the exploration of an efficient and simple approach to synthesize bimetallic nanocatalysts with well-defined geometrical architectures is of great significance and thus becomes an intensive research topic. Hitherto, a variety of synthetic strategies, such as galvanic replacement, seed-mediated growth, and co-reduction, have been developed for the preparation of various sophisticated bimetallic nanostructures with controllable shape and composition.

One of the representative applications of noble metal catalyst is electrocatalytic oxygen reduction reaction (ORR), which is of great importance in the cathode reaction of fuel cells. Although platinum-based catalysts exhibit exceptional activity and performance toward ORR, the extremely high cost and limited resources would restrict their practical large-scale commercialization in fuel cells. As one of the alternatives, Pt-based materials have been identified as acceptable electrocatalysts comparable to platinum for ORR in alkaline solutions. In addition, Ag, as a less-expensive and relative-abundant metal, exhibits good ORR activity in alkaline media, especially for HO₂⁻ disproportionation. Thus the alloy of Ag with Pd would be promising to achieve enhanced catalytic activity with better Pd utilization efficiency and economically desirable since the Pd has much higher (>20 times) cost than Ag.

Herein, we demonstrate the successful synthesis of triangular Ag-Pd alloy nanoprisms by employing Ag nanoparticles as sacrificial templates to define both the morphology and size of the final structures. The synthetic strategy relies on the combination of galvanic replacement between Ag nanoprisms and H₂PdCl₄ and co-reduction of Ag⁺ and Pd²⁺ in the reaction system. The prepared Ag-Pd alloy nanoprisms exhibited superior electrocatalytic activity in ORR as compared with the commercial Pd/C catalyst, which might be attributed to the synergistic effects of the Ag-Pd alloyed feature along with the dominantly exposed {111} facets of triangular nanoprisms.
Results and discussion

Structural characterizations of Ag-Pd alloy nanoparticles

For the synthesis of triangular Ag-Pd alloy nanoparticles, uniform triangular Ag nanoparticles with an average edge length of 90 nm (Fig. 1a) were firstly prepared through a photoinduced approach. Then solutions of H2PdCl4 and ascorbic acid (AA) were simultaneously introduced into the Ag nanoprisms solution through two separate tubes on a programmable syringe pump. During the process, the Ag nanoparticles served as sacrificial templates to form Ag-Pd alloy nanoparticles. The representative transmission electron microscopy (TEM) image of the obtained product (Fig. 1b) indicates that the final structures retain triangular shape with slightly larger edge length as compared with the Ag prism templates. The energy-dispersive spectroscopy (EDS) (Fig. S1) confirms that the as-prepared nanoparticles are composed by Ag and Pd. The measurement by atomic force microscopy (AFM) revealed that the average thickness of the Ag-Pd alloy nanoparticles is ~18 nm (Fig. S2), which is also larger than that of the original Ag prisms (~12 nm, Fig. S3). In addition, the Ag-Pd nanoparticles exhibit darker contrast in the edges relative to the center, implying that the edges are thicker than the interior region, which is also supported by the AFM measurement result (Fig. S2). Moreover, on the surface of Ag-Pd alloy nanoparticles, there are some nanochannels that could increase the specific surface area and allow for the transport of molecules/ions, thus would be of significance to catalytic applications.

![Fig. 1 Characterizations of morphology, structure, and composition of the prepared Ag-Pd alloy nanoparticles.](image)

(a) TEM image of the original Ag nanoparticles. (b)-(c) TEM images, (d) HRTEM image, (e) HAADF-STEM image and EDX line profile, and (f) EDX mapping images of Ag-Pd alloy nanoparticles. The inset of (c) shows the SAED pattern of an individual Ag-Pd alloy nanoprisms, and the spots (triangular, square and circle) could be indexed to the {220}, {422}, and forbidden 1/3{422} reflections, respectively. This result is similar to the electron diffraction pattern of the original Ag nanoprisms (Fig. S4) and demonstrates the single crystallinity of the Ag-Pd alloy nanoprisms. The well-resolved fringes with a lattice spacing of 0.248 nm (Fig. 1d) can be ascribed to the (1/3){422} reflection that is generally forbidden for an fcc lattice.

The X-ray diffraction (XRD) pattern (Fig. S5) of the product shows diffraction peaks of typical fcc-structured metal. Notably, all peaks are locating between the standard peaks of pure Ag and Pd, and neither an Ag nor Pd single component peak was observed. This result confirms the Ag-Pd alloy phase in the prepared nanoparticles, in line with previous observations in Ag-Pd alloy nanostructures. The elemental distribution of Ag and Pd in the nanoprisms was measured through compositional line profiles and EDX mapping analyses by high-angle annular dark-field scanning TEM (HAADF-STEM). As shown in Fig. 1e, the Ag and Pd traces showed very good matching with two peaks at the edges and one valley in the center. The elemental mapping results (Fig. 1f) indicate that both Ag and Pd are homogeneously distributed throughout the nanoparticles, further confirming the alloy composition. The inductively coupled plasma atomic emission spectroscopy (ICP-AES) reveals that the weight percentage of Pd in the products is 57.7%, suggesting a molar ratio of Ag to Pd as 42:58.

Morphological evolution and formation mechanism of Ag-Pd alloy nanoparticles

In order to reveal the morphological evolution and understand the formation mechanism of the Ag-Pd alloy nanoparticles, a series of TEM images (Fig. 2) were taken from the intermediate samples collected at different reaction intervals. In the initial reaction stage (t = 45 min, Fig. 2a-2b), the prism template morphology was essentially maintained, and selective Pd deposition on the edges occurred so that the prism edges exhibited darker contrast under TEM observation. When the reaction proceeded further (t = 90 min, Fig. 2c-2d), some areas on the triangular surface displayed less contrast and small pinholes appeared in the prism structure, which is ascribed to the galvanic replacement between H2PdCl4 and Ag prisms. After 120-min reaction, Fig. 2e and 2f show that more Ag segments were excavated from the central part, leading to the formation of highly porous structures with pinholes randomly distributed throughout the surface. However, the sample collected at 180-min reaction showed relatively smaller hole regions on the nanoparticles (Fig. 2g-2h), indicating that those pinholes were backfilled. Eventually, upon such continuous backfilling and deposition, as shown in Fig. 2i and 2j, well-developed Ag-Pd alloy nanoparticles were generated after 240-min reaction.
On the basis of the above observations, we propose the formation mechanism of Ag-Pd alloy nanoprisms as illustrated by Fig. S4. Since the relative surface energies of different crystal facets are in the order of $\gamma_{\{111\}} \leq \gamma_{\{100\}} < \gamma_{\{110\}}$, and the Pd precursor (H$_2$PdCl$_4$) is introduced at a very slow rate through a programmable syringe pump to avoid self-nucleation of Pd$_0$. Therefore, at the initial reaction stage, the newly formed Pd atoms preferentially nucleate and grow on the prism edges to form Ag@Pd-framed nanoprisms due to the high-energy of {110} facets of Ag nanoprisms. Such initial site-selective Pd deposition on the prism edges is similar to the epitaxial Au growth on the edges of Ag nanoprisms. Since Ag-Pd system is a well-known binary alloy, the newly formed Pd atoms readily diffuse into the Ag matrix to form seamless and homogeneous Ag-Pd alloys. Because the reduction potential of Pd$^{2+}$/Pd (0.951 V vs. SHE, SHE = standard hydrogen electrode) is more positive than that of Ag$^{+}$/Ag (0.8 V vs. SHE), Ag nanoprisms can be oxidized by H$_2$PdCl$_4$. Therefore, owing to the selectively galvanic corrosion on {111} facets, some pinholes start to appear preferentially on the triangular face of the nanoprisms as the H$_2$PdCl$_4$ concentration increases. These pinholes can further serve as channels of ion transport for the galvanic replacement.

It should be pointed out that the Ag atoms in the newly formed Ag-Pd alloy edges could not be easily oxidized back to Ag$^+$ through galvanic replacement due to their higher stability than pure Ag atoms in the original nanoprisms. As the reaction proceeded, the Ag {111} facets were gradually removed along with the deposition of more Pd atoms at the edges, creating pinholes on the prism surfaces. At the same time, co-reduction of Pd$^{2+}$ and Ag$^+$ (from the galvanic replacement) in the solution by AA would occur, leading to backfilling of Ag and Pd atoms as alloys predominantly at the inner edges of the pinholes because these sites possess larger roughness and thus higher surface energy than the outer prism edges and flat surface sites. After the pinholes are backfilled, further deposition of Ag and Pd atoms takes place on all nanoprism surfaces, including tips, edges, and triangular faces. As a result, it finally generated well-developed Ag-Pd alloy nanoprisms. The major structural evolution during the whole reaction is schematically illustrated in Fig. 3.

![Fig. 2 TEM images of Ag-Pd alloy nanostructures collected at different reaction time: (a)-(b) 45 min, (c)-(d) 90 min, (e)-(f) 120 min, (g)-(h) 180 min, and (i)-(j) 240 min.](image1)

![Fig. 3 Schematic illustration of morphological changes in the formation of Ag-Pd alloy nanoprisms from Ag nanoprisms. It involves four major steps: (1) initial deposition of Pd on the Ag nanoprism edges; (2) pinhole generation by dissolution of Ag from {111} facets and simultaneous deposition of Pd on {110} facets; (3) backfilling of the etched pinholes with Ag-Pd alloy; and (4) further deposition on all prism facets that increases both edge length and thickness of the Ag-Pd alloy nanoprisms.](image2)
Electrocatalysis activity of Ag-Pd alloy nanoprisms toward ORR

To evaluate the catalytic performance of the prepared Ag-Pd alloy nanoprisms, we investigated the electrocatalytic activity of these structures for oxygen reduction reaction (ORR) in alkaline medium with a rotating disk electrode (RDE). The results are compared with those of commercial Pd/C catalyst (Alfa Aesar, 10 wt%). Fig. 4a shows the typical cyclic voltammograms (CVs) of the as-prepared Ag-Pd alloy nanoprisms (red curve) and commercial Pd/C catalyst (black curve) in O₂-saturated KOH solution (0.1 M).

**Fig. 4** (a) Typical CVs of the as-prepared Ag-Pd alloy nanoprisms and commercial Pd/C catalyst in O₂-saturated 0.1 M KOH solution. (b) LSV curves of commercial Pd/C catalyst and Ag-Pd alloy nanoprisms. (c) Rotation-disk voltammograms recorded for Ag-Pd alloy nanoprisms supported on a GC electrode in an O₂-saturated 0.1 M KOH solution at a scan rate of 10 mV·s⁻¹ at different rotation rates. (d) Koutecky-Levich plot of \(j^1\) versus \(ω^{1/2}\) at 0.5 V.

As shown in Fig. 4a, distinct peaks corresponding to oxygen reduction can be observed for both Ag-Pd alloy nanoprisms and the commercial Pd/C catalyst. For commercial Pd/C catalyst, the ORR peak potential located at -0.28 V, while the reduction peak for the Ag-Pd alloy nanoprisms shifts positively to -0.21 V. This result indicates that the prepared Ag-Pd alloy nanoprisms exhibit an enhanced catalytic activity toward ORR in comparison with commercial Pd/C catalyst. Fig. 4b shows the typical linear sweep voltammograms (LSV) curves of commercial Pd/C catalyst and Ag-Pd alloy nanoprisms. The mass-normalized current density generated from Ag-Pd alloy nanoprisms is about 3.2 times higher than that of commercial Pd/C catalyst, which further suggests better ORR catalytic activity of Ag-Pd alloy nanoprisms as compared to the commercial Pd/C catalyst.

The enhanced electrocatalytic performance could be rationally attributed to the alloyed feature of the Ag-Pd nanostructures. It is generally accepted that the common ORR process, as the series 4-electron pathway, should involve both the breaking of an O-O bond and the formation of O-H bonds. In the Ag-Pd alloy structure, which has different electronic structure and local reactivity from the monometallic Pd element, the Pd atoms may facilitate the initial oxygen binding, while the Ag may help to desorb the reaction products such as OH⁻, resulting in the higher activity in the overall catalytic processes.

Furthermore, the dominance of [111] facets in Ag-Pd alloy nanoprisms also contributes to the enhanced ORR activity. Since all ORR involves both the dissociation of O-O bond and the removal of surface OH groups, the strong adsorption of intermediate OH_ads on the catalyst surface would have a negative impact on the catalyst’s performance. Owing to the weak interaction between OH_ads and [111] facets, the major surfaces of Ag-Pd alloy nanoprisms as [111] facets would have low coverage of OH_ads, which promotes the activity for ORR. In addition, due to the porosity feature of the alloy prisms, as shown in Fig. 1b, this structure possesses a relatively high accessible surface area and more catalytically active sites, which are beneficial to enhance the ORR activity. Therefore, on the basis of the aforementioned explanations, the enhanced ORR activity can be reasonably attributed to the synergistic effects of the alloyed composition and dominant [111] facets of triangular Ag-Pd nanoprisms along with the porous features.

The kinetics of ORR catalyzed by Ag-Pd alloy nanoprisms was further studied through measurements with RDE at various rotating speeds. Fig. 4c displays the rotation rate-dependent current density–potential curves. High rotational speeds result in increased oxygen diffusion to electrode surface and large currents. Fig. 4d represents the inverse current density \((j^1)\) at 0.5 V as a function of the inverse of the square root of the rotation rate \((ω^{1/2})\), the so-called Koutecky-Levich plot. The linearity of this plot is an indication of first-order reaction kinetics with respect to the concentration of dissolved O₂. From the Koutecky-Levich Equation, \(1/j_k = 1/j_k + 1/B ω^{1/2}\), in which \(B = 0.62nFCD_sD_O^0/v^{1/6}\) where \(j_k\) is the measured current density, \(j_k\) is the kinetic current density, \(ω\) is the rotation rate, \(n\) is the overall number of electrons transferred in oxygen reduction, \(F\) is the Faraday constant \((F = 96485 \text{C mol}^{-1})\), \(C_O\) is the bulk concentration of \(O_2 (1.26 \times 10^{-3} \text{mol L}^{-1})\), \(D_O\) is the diffusion coefficient of \(O_2\) in the electrolyte \((1.93\times10^{-5} \text{cm}^2 \text{s}^{-1})\), \(v\) is the viscosity of the electrolyte \((0.1 \text{m}^2 \text{s}^{-1} \text{in 0.1 M KOH})\). The number of electrons, \(n\), involved in the ORR was calculated to be 3.84 at -0.5 V, which indicates a nearly four-electron-transfer reaction occurring in the ORR reaction. As such, the as-obtained triangular Ag-Pd alloy nanoprisms with high ORR activity are expected for great applications in fuel cell systems.

**Conclusions**

In summary, we have demonstrated successful synthesis of triangular Ag-Pd alloy nanoprisms by using Ag nanoparticles as sacrificial templates. The Ag nanoparticles were initially coated with a layer of Pd, followed by the dissolution of Ag from
[111] facets due to the galvanic replacement, and then the Ag and Pd atoms backfilled to the [111] facets in a form of alloy via co-reduction of the Ag⁺ and Pd²⁺ by AA. Electrocatalytic tests for ORR indicated that the synthesized Ag-Pd alloy nanoprism displayed superior activity comparing with the commercial Pd/C catalyst. The enhanced ORR activity is attributed to the synergistic effects of the alloyed composition and dominant [111] facets of the triangular Ag-Pd nanoprism along with the porous features. This work would be highly impactful to the rational design of future bimetalllic alloy nanostructures with high catalytic activity for fuel cell systems.

**Experimental**

Synthesis of triangular Ag-Pd alloy nanoprism: Silver nanoprism was prepared through a photochemical method according to literature protocols (see Supporting Information for details). The as-prepared Ag nanoprisms solution (20 mL) was added into 20 mL Millipore water in a glass vial, followed by infusion of 4 mL of 0.3 mM H₂PdCl₄ solution and 4 mL of 8 mM ascorbic acid solution through two separate tubes on a programmable syringe pump. The infusion rate was set as 1.0 mL·h⁻¹, and the solution was kept vigorous stirring during infusion. The H₂PdCl₄ aqueous solution was prepared by completely dissolving PdCl₂ in HCl (0.2 M) at 60 °C.

Electrochemical measurements: The electrochemical measurements were carried out in a standard three-electrode cell on an Autolab potentiostat/galvanostat electrochemical workstation. The Ag-Pd alloy nanoprism modified glassy carbon (GC) electrode with a geometric area of 0.071 cm² served as a working electrode for immobilization of nanoparticles, 5 μL solution of Ag-Pd alloy nanoprism (C_Pd = 24.6 μg/mL) was dropped onto the GC electrode and dried under ambient condition. Then 5 μL of Nafion solution (1%) was dropped on the Ag-Pd alloy modified GC electrode to prevent erosion of nanocatalysts during electrochemical measurements. The modified substrate was washed with DI water and dried. Pt wire and Ag/AgCl electrode were used as the counter and reference electrodes, respectively. The electrochemical measurements for oxygen reduction reaction (ORR) were performed in 0.1 M KOH solution (O₂-saturated). The current density is normalized by the electrode area.

**Acknowledgements**

This work was financially supported by NTU seed funding for Solar Fuels Laboratory, MOE AcRF-Tier1 (RG 44/11), MOE AcRF-Tier2 (MOE2012-T2-2-041, ARC 5/13), and CRP (NRF-CRP5-2009-04) from NRF Singapore. H.Z. thanks the support from MOE under AcRF Tier 2 (ARC 26/13, No. MOE2013-T2-1-034), AcRF Tier 1 (RG 61/12), and Start-Up Grant (M4080865.070.706022) in Singapore. This research is also funded by the Singapore National Research Foundation and the publication is supported under the Campus for Research Excellence And Technological Enterprise (CREATE) programme (Nanomaterials for Energy and Water Management).

**References**

School of Materials Science and Engineering
Nanyang Technological University
50 Nanyang Avenue, 639798, Singapore
E-mail: cxue@ntu.edu.sg
Electronic Supplementary Information (ESI) available. See DOI: 10.1039/b000000x/