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# One-pot Synthesis of Pd-Pt@Pd Core-shell Nanocrystals with Enhanced Electrocatalytic Activity for Formic Acid Oxidation

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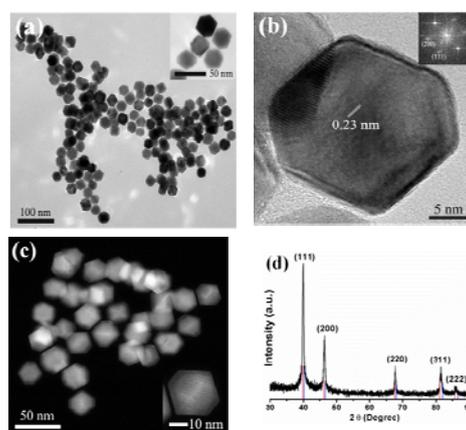
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**Well-defined Pd-Pt@Pd core-shell nanocrystals with Pd-Pt alloy core and conformal Pd shell of ~2-3 nm were directly synthesized through an one-pot, aqueous solution approach without any preformed Pd or Pt seeds. These Pd-Pt@Pd core-shell nanocrystals show an enhanced electrocatalytic activity for formic acid oxidation compared with commercial Pd black.**

Today, synthesis of bimetallic nanomaterials with controlled structure, e.g. size, shape, core/shell, alloy and composition has attracted increasing interest because of the superior catalytic, optic and magnetic properties of bimetallic nanomaterials relative to their monometallic counterparts.<sup>1-5</sup> Pd-Pt bimetallic nanocrystals are significant systems owing to the potential applications in electrocatalytic oxidation of small organic molecules (formic acid and methanol),<sup>6,7</sup> oxygen reduction reaction (ORR),<sup>8</sup> hydrogen oxidation reaction,<sup>9</sup> hydrogenation of aromatic hydrocarbons<sup>10</sup> and hydrogen storage.<sup>11</sup> Owing to the intrinsic physical properties of both palladium and platinum, namely, (1) both metals adopt a face-centered cubic (*fcc*) packing structure, (2) the atomic radii are within 15% (1.4% for Pd and Pt, 1.37 and 1.39 Å, respectively), (3) they have similar electronegativities (2.20 and 2.28 for Pd and Pt on the Pauling electronegativity scale, respectively),<sup>12-15</sup> and (4) they have a very small lattice mismatch of only 0.77%, all these factors make them easily form continuous solid solutions (alloy) for all compositions through a bottom-up, solution route.<sup>16</sup> To date, hydrothermal/solvothermal methods are mainly adopted to produce Pd and Pt bimetallic nanocrystals through coreduction of Pd and Pt precursors or a seed-mediated growth method. However, for coreduction of Pd and Pt species produced from Pd and Pt salt precursors, it is difficult to acquire core-shell nanocrystals and often results in alloy nanocrystals.<sup>7, 17-22</sup> Pd@Pt or Pt@Pd core-shell nanocrystals<sup>15, 23-28</sup> and Pt-on-Pd nanodendrites<sup>8, 29</sup> were synthesized by using two-step, seed-mediated epitaxial growth method with faceted Pd or Pt

nanoparticles as seeds. For example, Yang and co-workers reported the synthesis of Pt@Pd core-shell nanocrystals using seed-mediated epitaxial growth method with 13.4 nm Pt nanocubes as seeds.<sup>23</sup> Similar approaches were also used to synthesize Pd@Pt or Pt@Pd core-shell nanocrystals by Xia and co-workers.<sup>25</sup> Beer and co-workers have demonstrated a supramolecular route for the synthesis of Pd@Pt or Pt@Pd core-shell nanoparticles, based on an anion coordinating on the surface of preformed Pd or Pt core nanoparticles.<sup>26</sup> Moreover, Xia's and Yang's group displayed the synthesis of Pt-on-Pd bimetallic nanodendrites using sub-10 nm Pd nanoparticles as seeds.<sup>8, 28</sup> As mentioned above, Pd and Pt bimetallic heterostructures are mainly produced through two-step, seed-mediated epitaxial growth method, it seems as though it could be impossible to synthesize well-defined Pd and Pt bimetallic heterostructures (e.g., core-shell structures) by coreduction of Pd and Pt species without any Pd or Pt seeds because of the intrinsic physical properties of both Pd and Pt elements. In recently, however, Yamauchi and co-workers made a significant breakthrough in the synthesis of Pt-on-Pd nanodendrite heterostructures through one-step, direct synthesis route in aqueous solution in the presence of K<sub>2</sub>PtCl<sub>4</sub>, Na<sub>2</sub>PdCl<sub>4</sub> and Pluronic P123 with ascorbic acid (AA) as reducing agent.<sup>30</sup> It greatly inspired us to explore a reasonable route for synthesis of well-defined Pd and Pt bimetallic core-shell nanocrystals without any pre-synthesized Pd or Pt seeds.



<sup>85</sup> **Fig. 1** TEM (a), HRTEM (b) and HADDF-STEM (c) images and XRD patterns (d) of as-synthesized Pd-Pt@Pd core-shell nanocrystals (The inset in (b) is FT patterns of single nanocrystal.) The red and blue line in (d) are the standard peaks for Pd (red) and Pt (blue).

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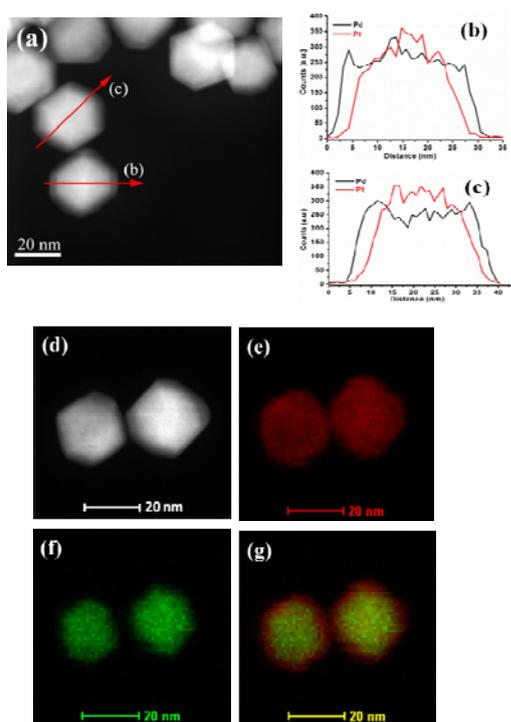
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† Electronic Supplementary Information (ESI) available: Experimental details, EDS profiles, STEM images, EDS line-scanning profiles, elemental maps, TEM, and CV. See <http://dx.doi.org/10.1039/b000000x/>

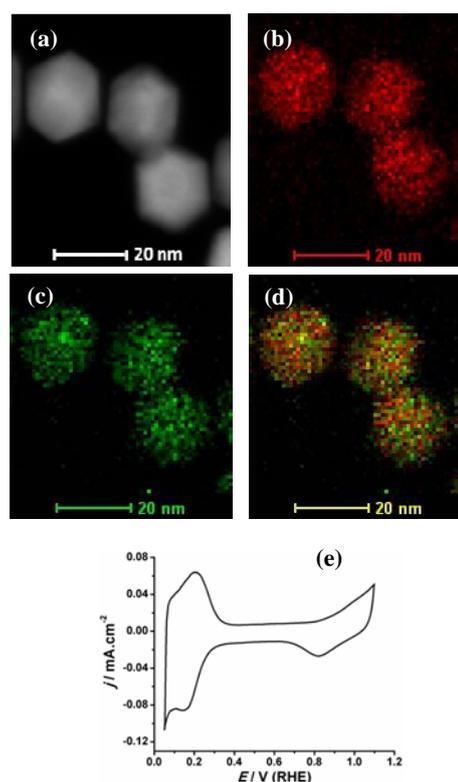
In this communication, for the first time, we reported a simple, one-pot, aqueous phase approach to directly produce Pd-Pt@Pd core-shell nanocrystals with Pd-Pt alloy core and Pd shell of ~2-3 nm without any of preformed Pd or Pt seeds. Fig. 1(a) shows the representative transmission electron microscopic (TEM) image of the product prepared with this "one-pot" approach. As can be seen, the product consists of a large quantity of uniform particles with well-defined octahedral shape and the average diameter of particles is 29 nm. The high-resolution TEM (HRTEM) image of one single particle is shown in Fig. 1(b). The well-resolved, continuous fringes in the same orientation crossing through the whole particle and the corresponding Fourier transform (FT) pattern indicate the particle is a single crystal. The interval between two lattice fringes was measured to be 0.23 nm, closed to the (111) lattice spacing of the face-centered cubic (*fcc*) Pd/Pt. The boundary between core and shell is not clear because of the weak contrast of Pd and Pt under the bright-field TEM. Nevertheless, the high-angle annular dark-field scanning transmission electron microscope (HAADF-STEM) images of the product (Fig. 1c and Fig. S1 (ESI<sup>†</sup>)) show the nanocrystals are core-shell octahedral structures with some truncation at corners and the thickness of shell is 2-3 nm. The X-ray diffraction (XRD) pattern (Fig. 1d) of the as-synthesized product shows five peaks corresponding to (111), (200), (220), (311) and (222) of *fcc* Pd/Pt. Moreover, the energy dispersive X-ray (EDX) spectra reveal the product is made of Pt and Pd (Fig. S2 (ESI<sup>†</sup>)).



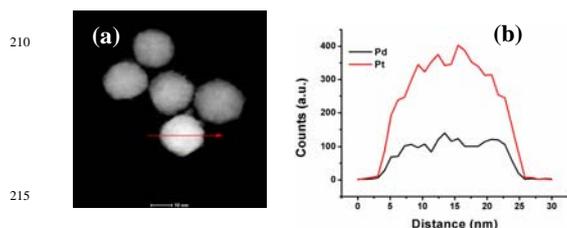
**Fig. 2** HADDF-STEM image (a) and the corresponding EDX line scanning profiles (b), (c) of two individual nanocrystals as indicated in (a). (d) HADDF-STEM image and the corresponding elemental maps (e) Pd (Red) and (f) Pt (Green). (g) Reconstructed overlay image of the maps shown in panels (e) and (f).

To clarify the structure and composition of the bimetallic core-shell nanocrystals, we further combined EDX line scanning with

element analysis mapping to analyze the core-shell nanocrystals. The positional distribution of Pd and Pt in the core-shell octahedron was revealed by EDX line-scan analysis and element analysis mapping. Fig. 2a-c display two line-scan EDX spectra of elemental Pd and Pt that were recorded through the center of two nanocrystals (marked by the red arrow in Fig. 2a). As can be seen, there is no Pt trace at the edge of nanocrystal in the range of 2-3 nm, while the Pd trace is very clear. It indicates that the nanocrystal is core-shell nanostructure with Pd shell. In the region of Pt, moreover, the signals of Pd don't decrease obviously, which is obviously different from the signals of Pt@Pd or Pd@Pt core-shell nanocrystal.<sup>24, 25</sup> It hints that the core of the as-synthesized core-shell nanocrystal is Pd-Pt alloy. The suggestion was further confirmed by the element analysis mapping. Fig. 2d-g show the HADDF STEM images and element analysis mappings of two as-synthesized nanocrystals. The Pd mappings (Fig. 2e) have the same size and shape as the as-synthesized nanocrystals shown in the Fig. 2d and the signals of Pd in the center regions of the nanocrystal are stronger than those at the edge regions. It indicates that the Pd distributes in the whole particle. The size of the Pt mappings (Fig. 2f) is obvious smaller than that of the as-synthesized nanocrystals (Fig. 2d). It indicates that there is no Pt element in the shell regions of the core-shell nanocrystals. Moreover, the overlaps of Pd and Pt mappings (Fig. 2g) further confirm the above-mentioned suggestion, namely, the core-shell nanocrystals consist of Pd shell and Pd-Pt alloy core and the thickness of the shell is 2-3 nm.



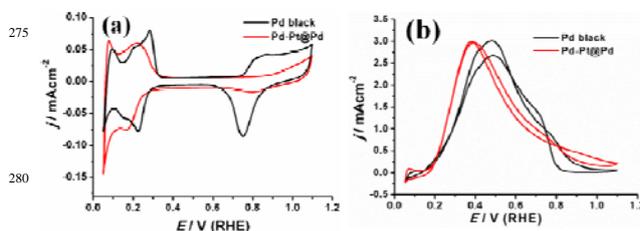
**Fig. 3** HADDF-STEM image (a), the corresponding elemental maps (b) Pd (Red); (c) Pt (Green) and (d) reconstructed overlay image shown in panels (b) and (c) and (e) CV in 0.1 M of H<sub>2</sub>SO<sub>4</sub> solution of nanocrystals synthesized with molar ratio of Na<sub>2</sub>PdCl<sub>4</sub> to H<sub>2</sub>PtCl<sub>6</sub> of 1:1.



**Fig. 4** HADDF-STEM image (a) and EDX line scanning profiles (b) of nanocrystals synthesized with molar ratio of  $\text{Na}_2\text{PdCl}_4$  to  $\text{H}_2\text{PtCl}_6$  of 1:4.

In order to get the desired nanostructures, the appropriate capping and reducing agents are often chosen to control the morphologies of nanocrystals through tuning the reduction kinetics of precursors in the solution approach. For instance, in our previous report, high quality solid Pd-Pt nanocube alloys were synthesized by coreduction of  $\text{PdCl}_2$  and  $\text{K}_2\text{PtCl}_6$  in the presence of KBr, sodium lauryl sulfate (SLS), and polyvinylpyrrolidone (PVP).<sup>19</sup> Lee and co-workers<sup>31</sup> described the formation of Au@Pd core-shell nanooctahedron in the presence of cetyltrimethylammonium chloride (CTAC) in an aqueous solution of  $\text{HAuCl}_4/\text{K}_2\text{PtCl}_6$  mixtures. Yamauchi and co-workers<sup>32</sup> demonstrated the synthesis of Au@Pd@Pt core-shell nanoparticles in an aqueous solution of  $\text{HAuCl}_4/\text{Na}_2\text{PdCl}_4/\text{K}_2\text{PtCl}_6$  mixtures with ascorbic acid (AA) and Pluronic F127 as reducing and capping agents, respectively. In our present work, to prepare Pd-Pt@Pd core-shell nanocrystals, cetyltrimethylammonium bromide (CTAB) plays an important role. To prove this suggestion, a series of control experiments were carried out. In the absence of CTAB, the product was made of non-dispersed bulky aggregates. When equal mole of CTAC replaced CTAB, no core-shell nanocrystals can be produced, and the product consists of Pd-Pt alloy nanocrystals (Fig. S3 (ESI<sup>†</sup>)). Furthermore, when equal mole of NaBr replaced CTAB, the product was made of non-dispersed irregular aggregates (Fig. S4 (ESI<sup>†</sup>)). However, in the absence of citric acid (CA), no product can't be acquired. These results indicate that CTAB and CA mainly act as capping and reducing agents, respectively. In addition, to get Pd-Pt@Pd core-shell nanocrystals, the molar ratio of  $\text{Na}_2\text{PdCl}_4$  to  $\text{H}_2\text{PtCl}_6$  is also a key factor. When decreasing the molar ratio of  $\text{Na}_2\text{PdCl}_4$  to  $\text{H}_2\text{PtCl}_6$  from 2:1 to 1:1, the results of element analysis mapping and CV in  $\text{H}_2\text{SO}_4$  (Fig. 3) show that the product consists of not Pd-Pt@Pd core-shell nanocrystals but Pd-Pt alloy nanocrystals. While decreasing the molar ratio of  $\text{Na}_2\text{PdCl}_4$  to  $\text{H}_2\text{PtCl}_6$  to 1:4, the product consists of sphere-like alloy particles (Fig. 4), no core-shell Pd-Pt@Pd nanocrystals were acquired. Furthermore, the thickness of Pd shell can be increased simply by increasing the amount of Pd precursors. When the molar ratio of  $\text{Na}_2\text{PdCl}_4$  to  $\text{H}_2\text{PtCl}_6$  is 4:1, the thickness of Pd shell is about 7.8 nm (Fig. S5 (ESI<sup>†</sup>)). Based on above-mentioned experimental results, we deduce that the formation mechanism of Pd-Pt@Pd core-shell nanocrystals is mainly through following steps: firstly, when CTAB was added in the aqueous solution, Cl<sup>-</sup> anions in the precursors were easily replaced by Br<sup>-</sup> anions to form harder to reduction of  $\text{PdBr}_4^{2-}$  and  $\text{PtBr}_6^{2-}$  species ( $\text{PdCl}_4^{2-}/\text{Pd}$  (0.62 V versus RHE);  $\text{PtCl}_6^{2-}/\text{Pt}$  (0.74 V versus RHE);

$\text{PdBr}_4^{2-}/\text{Pd}$  (0.49 V versus RHE) and  $\text{PtBr}_6^{2-}/\text{Pt}$  (0.61 V versus RHE) because of  $\text{Pd}^{2+}/\text{Pt}^{4+}$  binding Br<sup>-</sup> more strongly than Cl<sup>-</sup>,<sup>31-36</sup> then  $\text{PdBr}_4^{2-}$  and  $\text{PtBr}_6^{2-}$  species were reduced to form Pd-Pt alloy by CA, and finally, the Pd shell formed on the Pd-Pt alloy cores through epitaxial and conformal growth by reduction of the rest  $\text{Pd}^{2+}$  species.



**Fig. 5** The cyclic voltammetric curves (CVs) of as-synthesized Pd-Pt@Pd core-shell nanocrystals (the red) and commercial Pd black (the black). (a) in a 0.1 M  $\text{H}_2\text{SO}_4$  solution. (b) in a 0.1 M formic acid + 0.1 M  $\text{H}_2\text{SO}_4$  solution.

The electrocatalytic activity for formic acid oxidation of as-synthesized Pd-Pt@Pd core-shell nanocrystals was tested compared with commercial Pd black. As shown in Fig. 5a, the CV of Pd-Pt@Pd shows a similar shape with that of the commercial Pd black, and is different from the CVs of Pd-Pt alloys.<sup>7, 19</sup> It also confirms that the as-synthesized nanocrystals have a core/shell structure with Pd shell, not an alloy structure. In the positive scanning direction, the two peaks stand for hydrogen under potential desorption ( $\text{H}_{\text{upd}}$ ) and the formation of a hydroxide layer ( $\text{OH}_{\text{ad}}$ ) on the surface of nanocrystals, respectively. In the negative scanning direction, the two peaks ascribe the surface reduction of nanocrystals and hydrogen adsorption, respectively. The  $\text{H}_{\text{upd}}$  and  $\text{OH}_{\text{ad}}$  peaks of the Pd-Pt@Pd nanocrystals are shift negatively and positively, respectively, as compared with those of the commercial Pd black, which is maybe attributed to thin Pd shell of the Pd-Pt@Pd nanocrystals. The hydrogen adsorption/desorption electric charges calculated from CVs were used to estimate the electrochemically active surface area (ECSA) of catalysts. Fig. 5b shows the comparison of electrocatalytic properties of the Pd-Pt@Pd nanocrystals and commercial Pd black for formic acid oxidation. The current has been normalized by ECSA to yield current density ( $j$ ). The two samples have a similar current density, about  $3.1 \text{ mAcm}^{-2}$ . However, the peak potential of Pd-Pt@Pd nanocrystals is much lower than that of commercial Pd black, they are 0.37 and 0.48 V, respectively. It indicates that formic acid is more easily oxidized on the surface of Pd-Pt@Pd nanocrystals than that of commercial Pd black. Moreover, when the potential is at 0.3 V, the current density of Pd-Pt@Pd nanocrystals is  $2.0 \text{ mAcm}^{-2}$  and that of commercial Pd black is  $1.2 \text{ mAcm}^{-2}$ . It shows the current density of Pd-Pt@Pd nanocrystals has an obvious enhancement. However, the stability of Pd-Pt@Pd nanocrystals is weaker than that of commercial Pd black (Fig. S6 (ESI<sup>†</sup>)).

## Conclusions

In conclusion, for the first time, we have developed a simple, one-pot, aqueous phase approach to directly produce well-defined Pd-Pt@Pd core-shell nanocrystals with Pd-Pt alloy core and Pd shell of ~2-3 nm without any of preformed Pd or Pt seeds.

Characterizations of HRTEM, FT and XRD show that the as-synthesized Pd-Pt@Pd core-shell nanocrystals are single crystals. The electrocatalytic activity of Pd-Pt@Pd core-shell nanocrystals towards formic acid oxidation was investigated compared with the activity of commercial Pd black. The peak potential of Pd-Pt@Pd core-shell nanocrystals decreased 0.11 V in comparison with that of commercial Pd black.

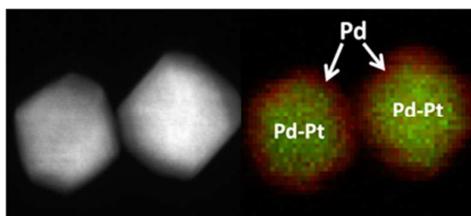
### Acknowledgements

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## Table of Content

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Pd-Pt@Pd core-shell nanocrystals were directly synthesized through an aqueous solution approach without any preformed Pd or Pt seeds.