

CrossMark  
click for updatesCite this: *RSC Adv.*, 2017, 7, 1947

# Synthesis of well-dispersive 2.0 nm Pd–Pt bimetallic nanoclusters supported on $\beta$ -cyclodextrin functionalized graphene with excellent electrocatalytic activity†

Xin Ran,<sup>‡a</sup> Long Yang,<sup>‡a</sup> Qing Qu,<sup>\*a</sup> Shunling Li,<sup>a</sup> Ying Chen,<sup>b</sup> Limei Zuo<sup>a</sup> and Lei Li<sup>\*b</sup>

In this work, ultrasmall Pd–Pt bimetallic nanoclusters with a uniform size of 2.0 nm monodispersed on  $\beta$ -cyclodextrin functionalized reduced graphene oxide ( $\beta$ -CD-RGO) were successfully prepared in aqueous solution at room temperature within 30 min without the need for any organic solvent or high temperature. The integration of  $\beta$ -CD and RGO was responsible for the formation of the monodispersed 2.0 nm Pd–Pt bimetallic nanoclusters. Inspired by the monodisperse, ultrasmall, and pristine properties of the Pd–Pt clusters, the Pd–Pt@ $\beta$ -CD-RGO nanohybrid displayed enhanced activity for methanol and ethanol oxidation in alkaline media in comparison with the commercial Pd/C catalysts. This facile and simple method is of significance for the preparation of bimetallic nanocatalysts with high catalytic activity on suitable supporting materials.

Received 8th October 2016  
Accepted 23rd November 2016

DOI: 10.1039/c6ra24893d

www.rsc.org/advances

## 1. Introduction

Over the past decades, extensive research has been devoted to electrocatalysis studies to improve the performance of noble metal catalysts.<sup>1</sup> One of the most promising strategies is to design and synthesize noble metals with ultrafine sizes because of their increased surface area and the number of edge and corner atoms, which could greatly improve their catalytic activities.<sup>2,3</sup> In particular, ultrasmall noble metal nanoclusters ( $\leq 2$  nm) with several to hundreds of atoms, have received significant attention.<sup>4</sup> However, the surface energy increases with decreasing particle size, which usually leads to serious aggregation of the small particles in order to minimize the total surface energy.<sup>2,5</sup> To overcome this aggregation, the metal particles must be anchored to suitable supports.<sup>6,7</sup> For this reason, extensive interests have been drawn to the uses of  $sp^2$ -hybridized carbon materials as the supporting materials.<sup>2,3,7</sup> Among all the available supporting materials, graphene is considered as the most promising candidate for growing or anchoring metal nanoparticles because of its high surface area ( $2630 \text{ m}^2 \text{ g}^{-1}$ ), high conductivity, and superior chemical inertness.<sup>7,8</sup> Moreover, theoretical research based on first-principles calculations indicates that metal atoms can interact with and

thereby bind with the  $sp^2$ -hybridized carbons of graphene because more interaction states and transmission channels are generated between both.<sup>3,9–11</sup> This implies that graphene may be an ideal substrate for growing and anchoring metal nanoparticles for high-performance electrocatalytic or electrochemical devices.

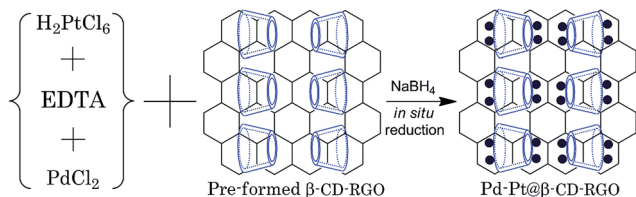
The Pt-based bimetallic nanostructures have attracted considerable attention in recent years because these materials are highly promising catalysts.<sup>12–14</sup> In comparison with monometallic Pt nanostructures, Pt-based bimetallic nanostructures show enhanced catalytic activities.<sup>1,13,15</sup> The catalytic properties of the bimetallic nanostructures are dependent on their sizes, shapes, and compositions. Fine control of the structural features, sizes, and compositions is highly favorable for improving catalytic performance Pt-based catalysts.<sup>13,14</sup> Among various Pt-based bimetallic nanostructures, Pt–Pd bimetallic nanostructures are considered as the most promising metallic nanostructures due to their superior catalytic activity.<sup>16</sup> Up to now, there are numerous studies have been reported<sup>13,14,17–27</sup> by controlling the morphology and composition to improve the catalytic activity of Pt–Pd bimetallic nanocatalyst. It is well known that the catalytic activities of noble metal particles are highly size dependent.<sup>4</sup> For instance, ultrafine Pd nanoparticles with a size of 3.5 nm supported on graphene oxide (GO) reported by Xie's group have displayed enhanced electrocatalytic activity for formic acid and ethanol electrooxidation.<sup>2</sup> Tang's group demonstrated that 1.8 nm Au clusters could be supported on reduced GO (RGO), which displayed high electrocatalytic performance towards oxygen reduction reactions.<sup>7</sup> Mao's group reported electroless deposition of 4.0 nm Pd clusters on

<sup>a</sup>School of Chemical Science and Technology, Yunnan University, Kunming 650091, China. E-mail: quqing@ynu.edu.cn; Fax: +86 871 65036538; Tel: +86 871 65035798

<sup>b</sup>Laboratory for Conservation and Utilization of Bio-Resources, Yunnan University, Kunming, 650091, China. E-mail: leelei@ynu.edu.cn

† Electronic supplementary information (ESI) available. See DOI: 10.1039/c6ra24893d

‡ These authors contributed equally to this work.



Scheme 1 Illustration for the preparation of the Pd-Pt@β-CD-RGO nanohybrid using the *in situ* reduction method.

graphdiyne oxides with high catalytic activity for the reduction of 4-nitrophenol.<sup>3</sup> Recently, a “soft” nitriding method to grow *in situ* 1.6 nm Au clusters on carbon reported by He's group exhibit superior activity for methanol oxidation.<sup>4</sup> However, it is still highly desirable to develop a facile and general approach for the fabrication of ultrafine bimetallic nanocatalysts such as Pd-Pt clusters with ultrasizes ( $\leq 2$  nm) though some monometallic clusters supported on carbon materials have been synthesized by different approaches if considering the potential industrial applications.

Herein, we report a facile aqueous approach (Scheme 1) to prepare 2.0 nm Pd-Pt bimetallic nanoclusters supported on β-cyclodextrin functionalized reduced graphene oxide (β-CD-RGO) at room temperature within 30 min without the need for any organic solvent or high temperature. The introduction of β-CD is used to immobilize Pd-Pt clusters on RGO and to avoid aggregation. Most importantly, it was found that the as-prepared Pd-Pt@β-CD-RGO nanohybrid exhibited higher electrocatalytic activities towards methanol and ethanol oxidation.

## 2. Experimental

### 2.1. Chemicals and materials

Graphite oxide was purchased from Nanjing XFNANO Materials Tech Co., Ltd (Nanjing, China). The  $\text{PdCl}_2$  (99.99%),  $\text{H}_2\text{PtCl}_6$  (99.99%), Nafion (5.0 wt%), and commercial Pd/C (10 wt%) were purchased from Sigma Chemical Co. (St. Louis, MO, USA). The β-CD and ethylene diamine tetraacetic acid disodium salt (EDTA) were obtained from Aladdin Chemical Reagent Co., Ltd (Shanghai, China). All other reagents were of analytical grade and used without further purification. Milli-Q ultrapure water (Millipore,  $\geq 18.25$  MΩ cm) was used throughout the experiments.

### 2.2. Preparation of the β-CD-RGO nanocomposite

Graphene oxide (GO) sheets were exfoliated from graphite oxide by ultrasonication in ultrapure water at room temperature for 1 h. A water-based wet-chemical route was adopted to prepare the β-CD-RGO composite. In a typical experiment, 20 mL of  $1.0 \text{ mg mL}^{-1}$  β-CD aqueous solution was mixed with 20 mL of  $0.5 \text{ mg mL}^{-1}$  GO aqueous suspension and stirred at room temperature for 30 min. After that, a small amount of NaOH aqueous solution (1.0 M) was used to adjust the pH of the mixture to 12.0. Finally, the suspension was transferred to a flask and stirred at  $90^\circ\text{C}$  in an oil bath for 4.0 h. After the reaction, the stable black suspension of the β-CD-RGO mixture

was centrifuged at a relative centrifugal force (RCF) of 30 600g. After washing with ultrapure water for three times, the resulting β-CD-RGO composite was collected by freeze-drying. Additionally, the RGO was prepared with the similar procedure in the absence of β-CD.

### 2.3. Synthesis of Pd-Pt@β-CD-RGO nanohybrid

Initially, a certain amount of β-CD-RGO freeze-dried powder was dispersed in ultrapure water with the aid of ultrasonication to obtain a concentration of  $1.0 \text{ mg mL}^{-1}$  β-CD-RGO homogeneous suspension. In a typical synthesis of Pd-Pt@β-CD-RGO, 3.0 mL of 10.0 mM  $\text{PdCl}_2$ , 3.0 mL of 10.0 mM  $\text{H}_2\text{PtCl}_6$ , and 2.0 mL of 0.1 M EDTA were mixed together and stirred to form a homogeneous solution. After adjusting the solution pH to 9.0, twenty milliliter of  $1.0 \text{ mg mL}^{-1}$  β-CD-RGO aqueous suspension was added into the above mixture and stirred for 5 min. Then, three milliliter of 200 mM sodium borohydride solution was added to the mixture and stirred for 30 min. After centrifuging and washing with ultrapure water for three times, the obtained products were collected by freeze-drying. In addition, the Pd-Pt@RGO, Pd@β-CD-RGO, and Pt@β-CD-RGO were prepared with the similar procedure.

### 2.4. Material characterization

The morphologies of the prepared samples were characterized by a JEM 2100 transmission electron microscopy (TEM, JEOL, Japan) with an accelerating voltage of 200 kV. High-resolution TEM (HRTEM) measurements, high-angle annular dark-field scanning TEM (HAADF-STEM), and elemental mapping were made on a JEM-2100F HRTEM (JEOL, Japan) operating at 200 kV. The samples for TEM and HRTEM characterization were prepared by depositing a drop of the diluted suspension on a copper grid coated by amorphous carbon. X-ray diffraction (XRD) patterns were recorded on a D8 ADVANCE (BRUKER, Germany) diffractometer using Cu-Kα radiation with a Ni filter ( $\lambda = 0.154059 \text{ nm}$  at 30 kV and 15 mA). X-ray photoelectron spectroscopy (XPS) measurements were performed with Al Kα X-ray radiation as the X-ray source for excitation, which were carried out on an ESCALAB-MKII spectrometer (VG Co., United Kingdom). Fourier transform infrared (FTIR) study was performed over the wavenumber, range of  $4000\text{--}400 \text{ cm}^{-1}$  by a Thermo Fisher SCIENTIFIC Nicolet IS10 (Massachusetts, USA) FTIR impact 410 spectrophotometer using KBr pellets. Thermogravimetric analysis (TGA) was carried out on a Q50 TGA (TA Instruments, New Castle, USA), at a heating rate of  $5^\circ\text{C min}^{-1}$  from 35 to  $800^\circ\text{C}$  in nitrogen.

### 2.5. Electrochemical measurements

All the electrochemical measurements were carried out with a CHI 660E Electrochemical Workstation from Chenhua Instrument (Shanghai, China) and conducted using a three-electrode system, with the modified glassy carbon electrode (GCE, 3 mm in diameter) as working electrode, a platinum wire as the counter electrode, a saturated calomel electrode as the reference electrode. Before the modifications, GCE was polished carefully with 0.3 and  $0.05 \mu\text{m}$   $\text{Al}_2\text{O}_3$  powder to obtain a mirror-



like surface, and then followed by sonication in ethanol and ultrapure water respectively. Then, the GCE was coated with the as-produced samples (5.0  $\mu\text{g}$ ) and dried at room temperature. Finally, 5.0  $\mu\text{l}$  of Nafion (0.1 wt%) was coated on the surface of the modified GCE and dried before electrochemical experiments. Methanol and ethanol electro-oxidation measurements were performed in a solution of 1.0 M KOH containing 1.0 M methanol or 1.0 M ethanol at a scan rate of 50  $\text{mV s}^{-1}$ . The methanol or ethanol solution was deoxygenated with  $\text{N}_2$  bubbling for 30 min before electrochemical experiments. The electrochemically active surface area (ECSA) of each sample was estimated by cyclic voltammetry (CV) measurement carried out in 0.5 M  $\text{H}_2\text{SO}_4$  solution under  $\text{N}_2$  protection with a sweep rate of 50  $\text{mV s}^{-1}$ . Mass current densities were normalized by the loaded Pt + Pd amount.

### 3. Results and discussion

The  $\beta$ -CD-RGO composite was successfully prepared as proved by FTIR and TGA characterizations. As shown in Fig. S1,<sup>†</sup> the  $\beta$ -CD displayed the typical CD absorption features. It is found that the FTIR spectrum of RGO is featureless except the stretching vibrations of  $-\text{OH}$  ( $3400\text{ cm}^{-1}$ ),  $\text{C}=\text{C}$  conjugation ( $1630\text{ cm}^{-1}$ ), and  $\text{C}-\text{C}$  band ( $1190\text{ cm}^{-1}$ ), etc. Whereas the FTIR spectra of  $\beta$ -CD-RGO exhibit the typical CD absorption features of the coupled  $\text{C}-\text{O}/\text{C}-\text{C}$  stretching/ $\text{O}-\text{H}$  bending vibrations at  $1032\text{ cm}^{-1}$ , the coupled  $\text{C}-\text{O}-\text{C}$  stretching/ $\text{O}-\text{H}$  bending vibrations at  $1152\text{ cm}^{-1}$ , the  $\text{C}-\text{H}/\text{O}-\text{H}$  bending vibrations at  $1410\text{ cm}^{-1}$ , the  $\text{CH}_2$  stretching vibrations at  $2925\text{ cm}^{-1}$ , and  $\text{O}-\text{H}$  stretching vibrations at  $3400\text{ cm}^{-1}$ . The presence of these peaks confirmed that the  $\beta$ -CD molecules were successfully attached to the surface of the RGO.<sup>28</sup> Additionally, the TGA was used to determine the amount of  $\beta$ -CD molecules on the surface of RGO. As shown in Fig. S2,<sup>†</sup> the  $\beta$ -CD showed an abrupt mass loss when the temperature was approximately  $260^\circ\text{C}$ . For the pristine RGO, the loss in mass (17.0 wt%) at a temperature of approximately  $600^\circ\text{C}$  was ascribed to the pyrolysis of the remaining oxygen-containing functional groups in RGO.<sup>28</sup> The  $\beta$ -CD@RGO composite revealed an abrupt mass loss when the temperature was approximately  $260^\circ\text{C}$  because of the decomposition of  $\beta$ -CD;<sup>29</sup> the mass loss reached about 45.0 wt% when the temperature was  $600^\circ\text{C}$ . The amount of  $\beta$ -CD grafted to RGO was estimated to be approximately 28.0 wt%.

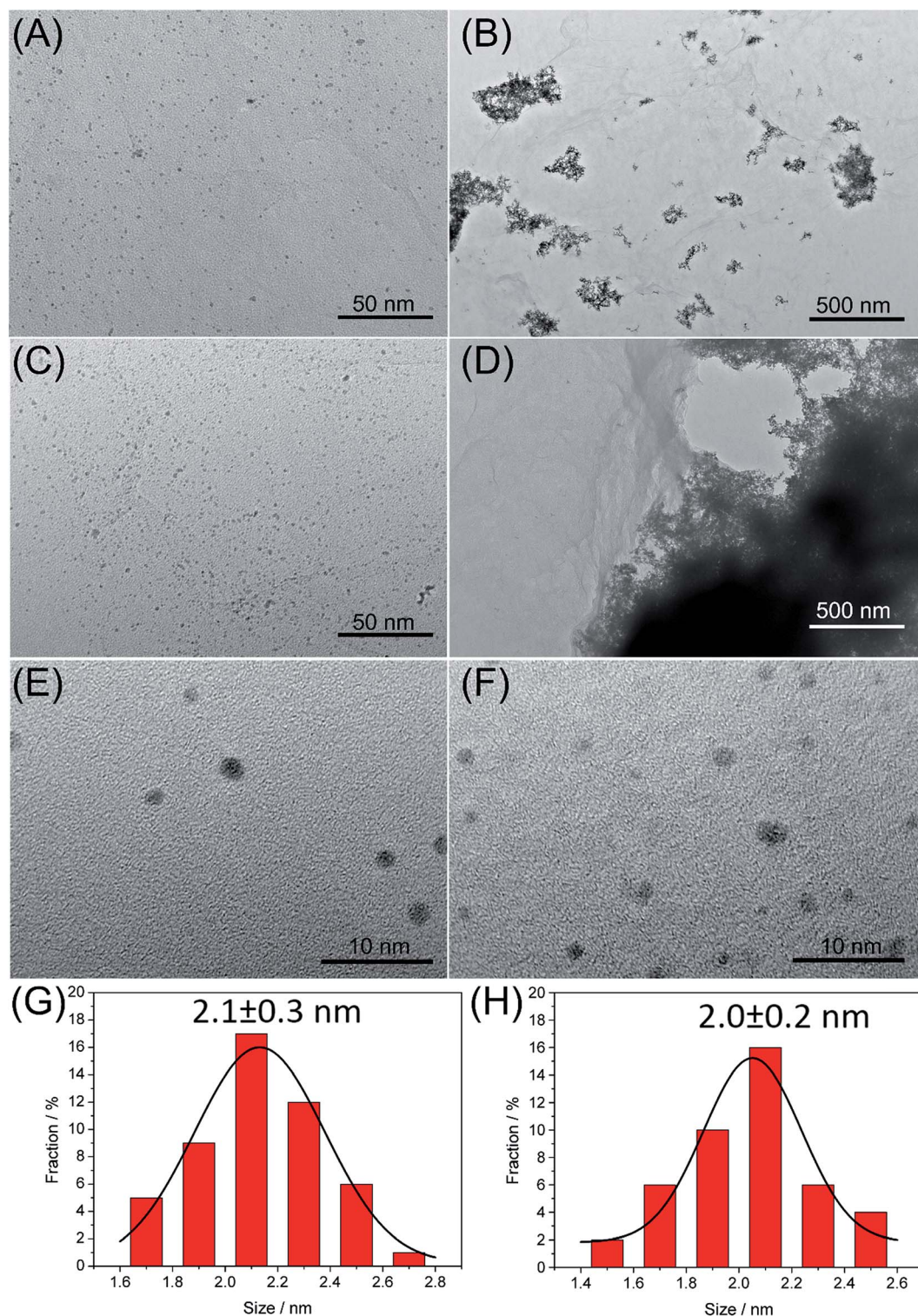
The Pd-Pt@ $\beta$ -CD-RGO and the related materials were characterized by TEM. As shown in Fig. 1A, the TEM image of Pd@ $\beta$ -CD-RGO indicates that the Pd clusters with uniform size could be obtained by using  $\beta$ -CD-RGO as the support. The high-resolution TEM (HRTEM) image (Fig. 1E) shows that the size of the Pd cluster is approximately  $2.1 \pm 0.3\text{ nm}$  with narrow size distribution, as demonstrated by the size distribution histogram (Fig. 1G). The well-dispersive Pd clusters might be attributed to abundant  $-\text{OH}$  groups on  $\beta$ -CD that are favorable for the coordination of cationic  $\text{Pd}^{2+}$  precursor *via* electrostatic interactions, followed by the *in situ* formation of the corresponding clusters on the surface of RGO. A similar function of  $-\text{OH}$  groups was also reported by Zhang *et al.* for the immobilization of Pd nanoparticles on hydroxypropyl- $\beta$ -CD modified

$\text{C}_{60}$ .<sup>30</sup> The TEM image of Pt@ $\beta$ -CD-RGO was illustrated in Fig. 1B. However, the Pt clusters showed an increased size with serious aggregation when using anionic  $\text{PtCl}_6^{2-}$  precursor. Thus, the nucleation of nanocatalysts on the  $\beta$ -CD-RGO is somewhat dependent on electrostatic interactions. These results are similar to that reported by Liu *et al.* using nitrated carbon as support to load metal clusters.<sup>4</sup> In their work, the positive charged N-doped carbon could load and stabilize Au clusters to avoid overgrowth due to its strong affinity to anionic  $\text{AuCl}_4^-$ . Fig. 1C shows the representative TEM image of the Pd-Pt@ $\beta$ -CD-RGO nanohybrid. It can be seen that the Pd-Pt bimetallic nanoclusters with a uniform size are well monodispersed on the surface of  $\beta$ -CD-RGO. The HRTEM image (Fig. 1F) reveals that the size of the Pd cluster is approximately  $2.0 \pm 0.2\text{ nm}$  with narrow size distribution, as proved by the size distribution histogram (Fig. 1H). The size of Pd-Pt in the bimetallic Pd-Pt@ $\beta$ -CD-RGO material is also very small. It is well known that Pd and Pt are very easy to form an alloy through the co-reduction of Pd and Pt salts in the solution phase because they have a very small lattice mismatch of only 0.77% and similar redox potential  $\text{Pd}^{2+}/\text{Pd}$  (0.62 V *versus* RHE);  $\text{PtCl}_6^{2-}/\text{Pt}$  (0.74 V *versus* RHE).<sup>20</sup> Thus, the very small size and uniform dispersibility of Pd-Pt clusters in the Pd-Pt@ $\beta$ -CD-RGO material may be ascribed to the fact that the Pd and Pt formed an alloy or heterostructure. The HRTEM image shows that the Pd-Pt clusters contain defects (several steps, corners, and edges), which can act as highly active sites in catalytic reactions.<sup>13,20</sup> In order to further investigate the effect of surface properties of the support on the formation of Pd-Pt clusters, a control experiment was carried out by using RGO as support under the same conditions except for the absence of  $\beta$ -CD. As shown in Fig. 1D, the TEM image of Pd-Pt@RGO reveals that the Pd-Pt bimetallic clusters exhibit an obviously increased size and serious aggregations. This confirms that  $\beta$ -CD plays a key role in controlling the size and the monodispersity of the Pd-Pt clusters. Therefore, well-dispersive Pd-Pt clusters with a uniform size and a narrow distribution could be obtained using  $\beta$ -CD-RGO as the composite support.

The crystal structures of the Pd@ $\beta$ -CD-RGO, Pt@ $\beta$ -CD-RGO, and Pd-Pt@ $\beta$ -CD-RGO composites were investigated by wide-angle X-ray diffraction XRD. The XRD patterns (Fig. 2A) of the synthesized materials show four peaks corresponding to  $\{111\}$ ,  $\{200\}$ ,  $\{220\}$ , and  $\{311\}$  of face-centered cubic (fcc) Pd/Pt. The ratio (3.3) between the intensities of the  $\{111\}$  and  $\{200\}$  peaks is higher than the value reported for some conventional samples (approximately 2.2),<sup>31,32</sup> indicating that the Pd-Pt clusters are mainly covered with  $\{111\}$  facets. Moreover, a broad diffraction peak (002) centered at  $2\theta = 23.9^\circ$  was observed, which was ascribed to the reduction of GO.<sup>33</sup> The corresponding energy dispersive X-ray (EDX) spectra revealed the peaks of C, O, Pd, and Pt elements, indicating the existence of bimetallic Pd-Pt clusters on the surface of  $\beta$ -CD-RGO (Fig. 2B). The Pd-Pt@ $\beta$ -CD-RGO composite was also characterized by high-angle annular dark-field scanning TEM (HAADF-STEM) (Fig. 2C). It is difficult to see the Pd-Pt clusters in the Pd-Pt@ $\beta$ -CD-RGO composite due to small size. The positional distribution of Pd, Pt, C, and O in the Pd-Pt@ $\beta$ -CD-RGO was revealed by element mapping







**Fig. 1** TEM images of Pd@β-CD-RGO (A), Pt@β-CD-RGO (B), Pd-Pt@β-CD-RGO (C), and Pd-Pt@RGO (D); the HRTEM images of Pd@β-CD-RGO (E) and Pd-Pt@β-CD-RGO (F); the corresponding size distribution histograms of Pd@β-CD-RGO (G) and Pd-Pt@β-CD-RGO (H).

analysis. Fig. 3A–D show that the Pd, Pt, C, and O elements are uniformly distributed in the Pd-Pt@β-CD-RGO composite. The Pd-Pt nanostructures are usually alloy structure<sup>14</sup> or hetero-structure<sup>13</sup> in published works. However, it is difficult to distinguish whether the present Pd-Pt cluster is alloy structure

due to the too small size. This can be ascribed to the fact that resolution of the instrument is not enough to distinguish the too small Pd-Pt clusters.

XPS analysis was used to study the electronic structure and compositions of the Pd-Pt@β-CD-RGO composite. As shown in



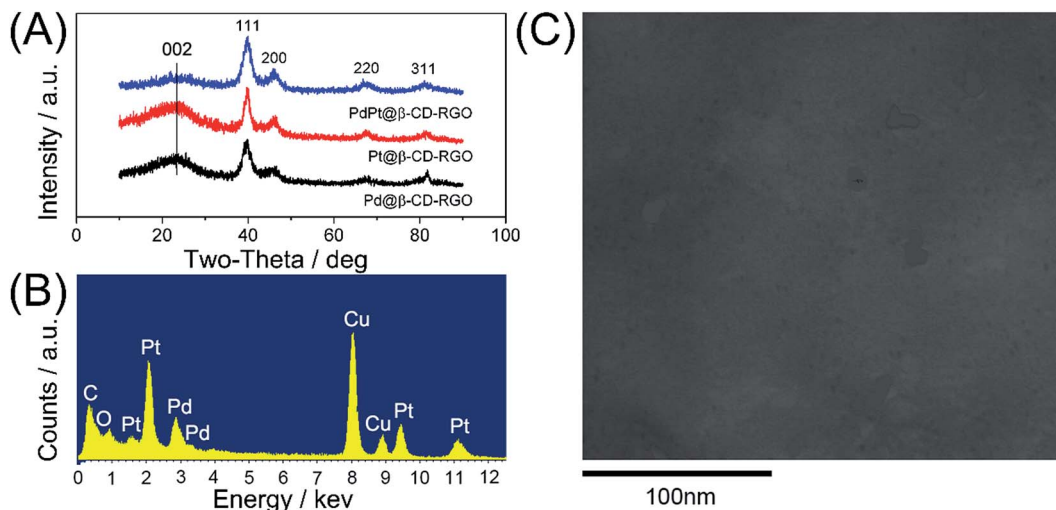


Fig. 2 (A) XRD patterns of Pd@β-CD-RGO, Pt@β-CD-RGO, and Pd-Pt@β-CD-RGO. (B) EDX of spectrum of Pd-Pt@β-CD-RGO. (C) HAADF-STEM image of Pd-Pt@β-CD-RGO.

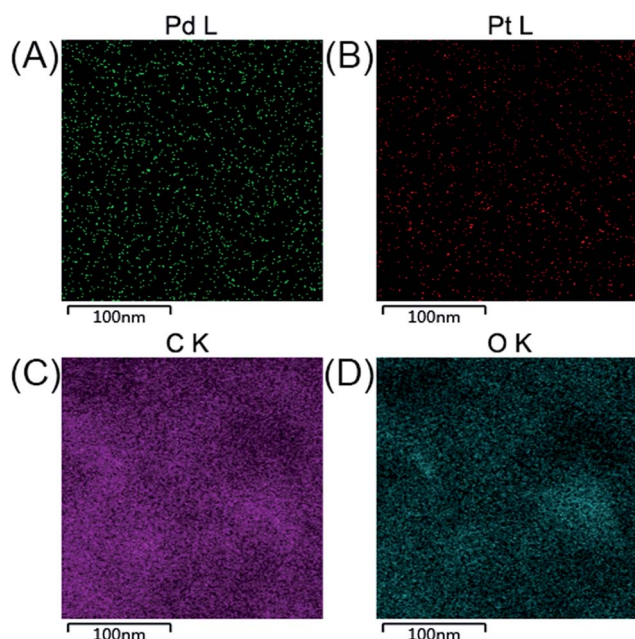


Fig. 3 The elemental maps of C (A), O (B), Pd (C), and Pt (D) in Pd-Pt@β-CD-RGO.

Fig. 4A, the survey spectrum confirms the presence of Pd, Pt, C, and O in the Pd-Pt@β-CD-RGO composite. The Na signal is attributed to residual NaOH. Fig. 4B is the C 1s spectrum, which reveals that there are three components of carbon bond, namely  $sp^2$  C (C=C, 284.6 eV),  $sp^3$  C (C-C, 285.3 eV), and C-O (286.8 eV) are presented in the composite.<sup>34–36</sup> The high-resolution spectrum of Pd 3d is illustrated in Fig. 4C. The peaks centered at 335.2 and 340.5 eV can be assigned to  $3d_{5/2}$  and  $3d_{3/2}$  peaks, respectively, which are attributed to metallic Pd.<sup>2,20,37</sup> In addition, the spectrum of Pt 4f is depicted in Fig. 4D. The  $4f_{7/2}$  and  $4f_{5/2}$  peaks at approximately 71.2 and 74.5 eV are ascribed to metallic Pt.<sup>20</sup>

The electrochemical active surface areas (ECSAs) of the Pd-Pt@β-CD-RGO, Pd-Pt@RGO, Pd@β-CD-RGO, Pt@β-CD-RGO, and commercial Pd/C catalysts were determined by the CV measurements in the  $N_2$  saturated electrolyte of 0.5 M  $H_2SO_4$  at a scanning rate of  $50\text{ mV s}^{-1}$ . As shown in Fig. S3,<sup>†</sup> all the CV curves exhibit obvious multiple peaks in the lower potential range of  $-0.3$  to  $0$  V associated with hydrogen adsorption/desorption and Pt/Pd oxide formation/reduction. The anodic peaks at the higher potential approximately  $0.6$  V and its cathodic counterparts in the potential range of  $0.45$  to  $0.33$  V are due to the Pt/Pd oxide formation/reduction. The ECSAs were estimated by the integrated charge ( $Q_H$ ) in the hydrogen desorption region after deduction of the double-layer region. On the basis of the assumption that the charge density for the formation of a fully covered  $Pd(OH)_2$  monolayer was  $430\text{ }\mu\text{C cm}^{-2}$ , according to the equation  $ECSA = Q_H / (430\text{ mC cm}^{-2} \times \text{Pd loading})$ , the ECSA for Pd/C and Pd@β-CD-RGO were  $27.9$  and  $38.0\text{ m}^2\text{ g}^{-1}$ , respectively. According to the equation  $ECSA = Q_H / (210\text{ mC cm}^{-2} \times \text{Pt loading})$ , the ECSA of Pt@β-CD-RGO was  $21.4\text{ m}^2\text{ g}^{-1}$ . According to the equation  $ECSA = Q_H / (210\text{ mC cm}^{-2} \times \text{Pt} + \text{Pd loading})$ , the ECSA of Pd-Pt@β-CD-RGO and Pd-Pt@RGO were  $51.0$  and  $41.3\text{ m}^2\text{ g}^{-1}$ , respectively. The higher ECSA of Pd-Pt@β-CD-RGO is favorable to the electrocatalytic activity for methanol or ethanol oxidation.

The catalytic performance of the Pd-Pt@β-CD-RGO toward methanol oxidation reaction (MOR) was investigated with three-electrode system in the  $N_2$  saturated electrolytes of  $1.0\text{ M KOH}$  and  $1.0\text{ M CH}_3\text{OH}$  at a scan rate of  $50\text{ mV s}^{-1}$  at room temperature. For comparison, the catalytic performance of the Pd-Pt@RGO, Pd@β-CD-RGO, Pt@β-CD-RGO, and commercial Pd/C was also tested. The currents were normalized to the loading amount of Pt + Pd in order to compare the mass activity of different catalyst. As shown in Fig. 5A, the Pd-Pt@β-CD-RGO exhibits the highest catalytic activity with a mass-normalized current density of  $1609\text{ mA mg}^{-1}$ , which is approximately 1.5, 2.7, 3.8, and 5.3 times greater than that of Pd-Pt@RGO ( $1077$





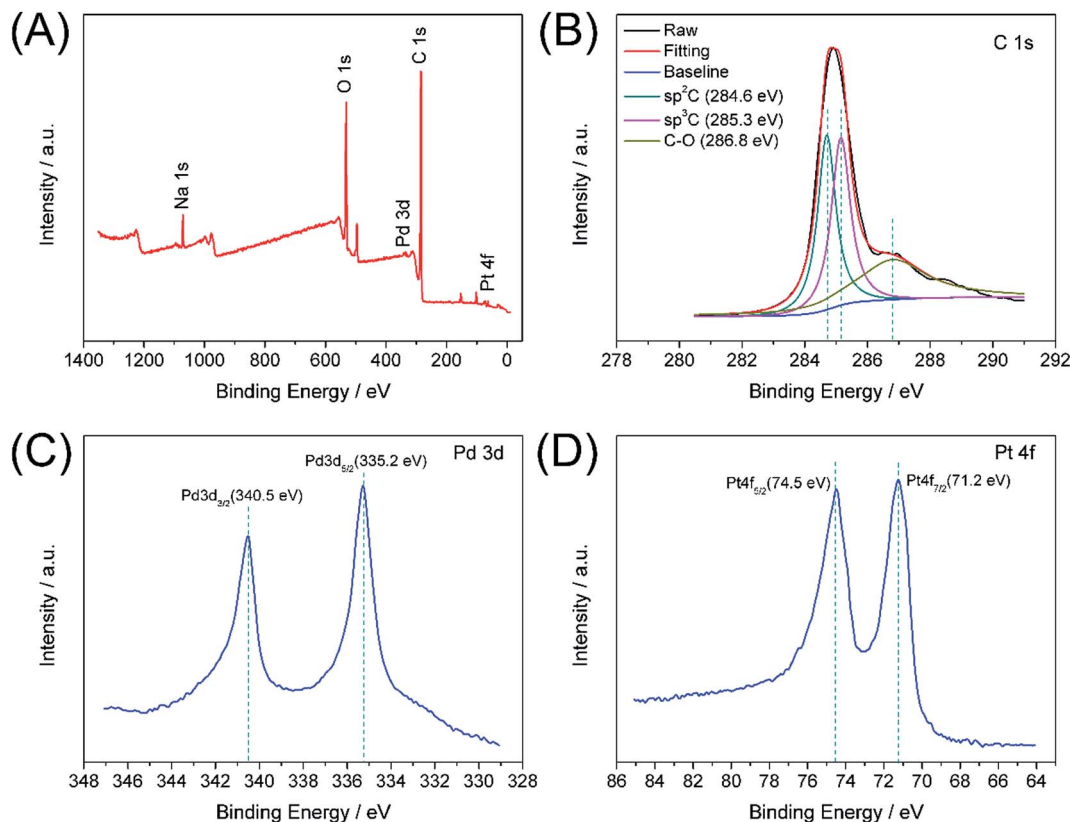


Fig. 4 XPS survey spectrum of Pd-Pt@β-CD-RGO (A); the narrow spectra of C 1s (B), Pd 3d (C), and Pt 4f (D).

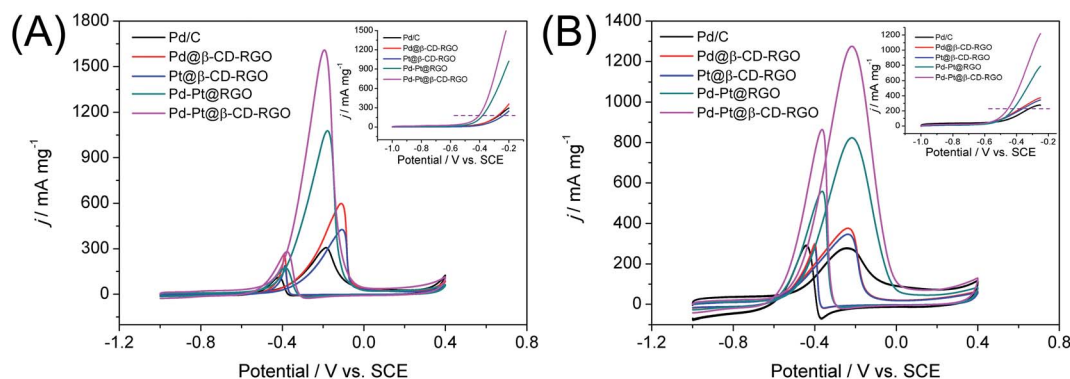


Fig. 5 Stable CV curves for methanol (A) and ethanol (B) oxidation reactions catalyzed by Pd/C, Pd@β-CD-RGO, Pt@β-CD-RGO, Pd-Pt@RGO, and Pd-Pt@β-CD-RGO in a solution of 1.0 M KOH containing 1.0 M methanol or 1.0 M ethanol at a scan rate of 50 mV s<sup>-1</sup>.

mA mg<sup>-1</sup>), Pd@β-CD-RGO (595 mA mg<sup>-1</sup>), Pt@β-CD-RGO (426 mA mg<sup>-1</sup>), and commercial Pd/C (305 mA mg<sup>-1</sup>) catalyst, respectively. The detailed electrochemical parameters of the as-prepared catalysts obtained from the CV curves were listed in Table 1. It should be noted that the specific activity of Pd-Pt@β-CD-RGO (3.15 mA cm<sup>-2</sup>) was also higher than that of Pd-Pt@RGO (2.60 mA cm<sup>-2</sup>), Pd@β-CD-RGO (1.57 mA cm<sup>-2</sup>), Pt@β-CD-RGO (1.99 mA cm<sup>-2</sup>), and commercial Pd/C (1.09 mA cm<sup>-2</sup>). Moreover, at the same current density in the positive scan (as indicated by the dashed line in inset of Fig. 5A), the corresponding oxidation potential on the Pd-Pt@β-CD-RGO

was the lowest, suggesting the enhancement of the methanol oxidation kinetics on the Pd-Pt@β-CD-RGO.<sup>13</sup> The improved electrooxidation kinetics of the Pd-Pt@β-CD-RGO may originate from the bimetallic Pd-Pt composition and the too small size (2.0 nm). In addition, the ratio of the mass current density values in two sequential forward and backward sweeps ( $J_f/J_b$ ), (as considered to be an important indicator of the catalyst tolerance to poisoning species<sup>14</sup>) showed significant differences for these catalysts. A higher ratio indicates more effective removal of the poisoning species on the catalyst surface.<sup>38</sup> The  $J_f/J_b$  values were 5.8, 5.6, 2.4, 2.3, and 2.7 for Pd-Pt@β-CD-RGO,



Table 1 The electrochemical parameters of the as-prepared different catalysts

Catalyst	Potential (V)		ECSA ( $\text{m}^2 \text{g}^{-1}$ )	Mass activity ( $\text{mA mg}^{-1}$ )		Specific activity ( $\text{mA cm}^{-2}$ )	
	Methanol	Ethanol		Methanol	Ethanol	Methanol	Ethanol
Pd/C	−0.20	−0.23	27.9	305	272	1.09	0.97
Pd@ $\beta$ -CD-RGO	−0.08	−0.23	38.0	595	376	1.57	0.99
Pt@ $\beta$ -CD-RGO	−0.10	−0.23	21.4	426	346	1.99	1.62
Pd–Pt@RGO	−0.20	−0.24	41.3	1077	820	2.60	1.98
Pd–Pt@ $\beta$ -CD-RGO	−0.24	−0.24	51.0	1609	1274	3.15	2.50

Pd–Pt@RGO, Pd@ $\beta$ -CD-RGO, Pt@ $\beta$ -CD-RGO, and commercial Pd/C, respectively. Thus, the Pd–Pt@ $\beta$ -CD-RGO exhibited best tolerance to the poisoning species than other and the Pd/C catalysts.

The Pd–Pt@ $\beta$ -CD-RGO, Pd–Pt@RGO, Pd@ $\beta$ -CD-RGO, Pt@ $\beta$ -CD-RGO, and commercial Pd/C were further studied for ethanol oxidation reaction (EOR) in 1.0 M KOH containing 1.0 M  $\text{C}_2\text{H}_5\text{OH}$  under  $\text{N}_2$  protection. As illustrated in Fig. 5B, the Pd–Pt@ $\beta$ -CD-RGO also displayed the highest catalytic activity with a mass current density of  $1274 \text{ mA mg}^{-1}$ , which is 1.6, 3.4, 3.7, and 4.6 times higher than that of Pd–Pt@RGO ( $820 \text{ mA mg}^{-1}$ ), Pd@ $\beta$ -CD-RGO ( $376 \text{ mA mg}^{-1}$ ), Pt@ $\beta$ -CD-RGO ( $346 \text{ mA mg}^{-1}$ ), and Pd/C ( $272 \text{ mA mg}^{-1}$ ) catalyst, respectively. The catalytic activity follows the order: Pd–Pt@ $\beta$ -CD-RGO > Pd–Pt@RGO > Pd@ $\beta$ -CD-RGO > Pt@ $\beta$ -CD-RGO > Pd/C, which is similar to the order of MOR. As shown in Table 1, the specific activity for EOR of Pd–Pt@ $\beta$ -CD-RGO ( $2.50 \text{ mA cm}^{-2}$ ) was also higher than that of Pd–Pt@RGO ( $1.98 \text{ mA cm}^{-2}$ ), Pd@ $\beta$ -CD-RGO ( $0.99 \text{ mA cm}^{-2}$ ), Pt@ $\beta$ -CD-RGO ( $1.62 \text{ mA cm}^{-2}$ ), and commercial Pd/C ( $0.97 \text{ mA cm}^{-2}$ ). The onset potential (inset of Fig. 5B) of Pd–Pt@ $\beta$ -CD-RGO was more negative than those of other catalysts, also indicating the higher electrocatalytic performance of the Pd–Pt@ $\beta$ -CD-RGO toward ethanol oxidation. For the EOR, the  $J_{\text{L}}/J_{\text{b}}$  value of Pd–Pt@ $\beta$ -CD-RGO (1.5) was also higher than that of Pd–Pt@RGO (1.4), Pd@ $\beta$ -CD-RGO (1.2), Pt@ $\beta$ -CD-RGO (1.3), and Pd/C (0.9), suggesting the better tolerance of Pd–Pt@ $\beta$ -CD-RGO to the poisoning species than other catalysts. In addition, the activity of the Pd–Pt@ $\beta$ -CD-RGO catalyst is also higher than those of recently reported Pd–Pt bimetallic catalysts as summarized in Table 2.

The enhanced electrocatalytic activity of the Pd–Pt@ $\beta$ -CD-RGO could be attributed to three main factors: (a) the Pd–Pt clusters with very small size (2.0 nm) would offer more

accessibility methanol or ethanol oxidation, thus remarkably improving the atomic utilization efficiency; (b) the excellent monodispersity and uniformity of Pd–Pt loaded on  $\beta$ -CD-RGO with high surface area might be favorable for improving the catalytic performance; (c) the formation of bimetallic surface atom arrangement (coexistence of surface Pd and Pt sites) and/or the synergistic effect originating from the coexistence of surface Pd and Pt atoms should also be emphasized. The TEM images of Pd/C were investigated and displayed in Fig. S4.† The size of Pd in Pd/C is approximately  $7.5 \pm 2.0 \text{ nm}$ . From these results, we can conclude that decreasing the size of noble metal nanoparticles can significantly improve the catalytic activity.

The durability of Pd–Pt@ $\beta$ -CD-RGO and commercial Pd/C was compared in 1.0 M KOH solution containing 1.0 M methanol using chronoamperometry and CV. At a fixed potential, the intermediate products such as CO from the methanol oxidation would adsorb and accumulate on the electrode surface, which might lead to the catalysts inactivation and the rapid current attenuation.<sup>39</sup> As shown in Fig. 6A, the current–time curves recorded at  $-0.2 \text{ V}$  for 4000 s are obtained. The oxidation current on commercial Pd/C modified electrode decreases much more rapidly than that on Pd@ $\beta$ -CD-RGO, Pt@ $\beta$ -CD-RGO, Pd–Pt@RGO, and Pd–Pt@ $\beta$ -CD-RGO modified electrodes at the initial stage. The currents on these modified electrodes gradually decay and achieve a pseudo-steady state. Obviously, the Pd–Pt@ $\beta$ -CD-RGO catalyst retains the highest current than other catalysts for the whole time. This result confirms that Pd–Pt@ $\beta$ -CD-RGO catalyst has the higher poison tolerance and better electrochemical stability than commercial Pd/C. The electrocatalytic cycling stabilities of these catalysts and commercial Pd/C in 1.0 M KOH solution containing 1.0 M methanol have also been investigated using CV, as shown in Fig. 6B. It can be observed that the forward peak current density

Table 2 Comparison of the Pt–Pd@ $\beta$ -CD-RGO with similar catalysts for methanol oxidation

Catalysts	Shapes of Pd–Pt	Potential (V)	ECSA ( $\text{m}^2 \text{g}^{-1}$ )	Specific activity ( $\text{mA cm}^{-2}$ )	Ref.
PdPt nanoalloys	Islands	0.60	21.7	1.08	20
Pt–Pd	Cubes	0.85	—	1.49	14
Pt-on-Pd	Nanodendrites	0.70	48.0	1.04	13
Pd@Pt	Porous particles	0.70	38.2	1.31	24
Pt–Pd@ $\beta$ -CD-RGO	Clusters	−0.24	51.0	3.15	This work



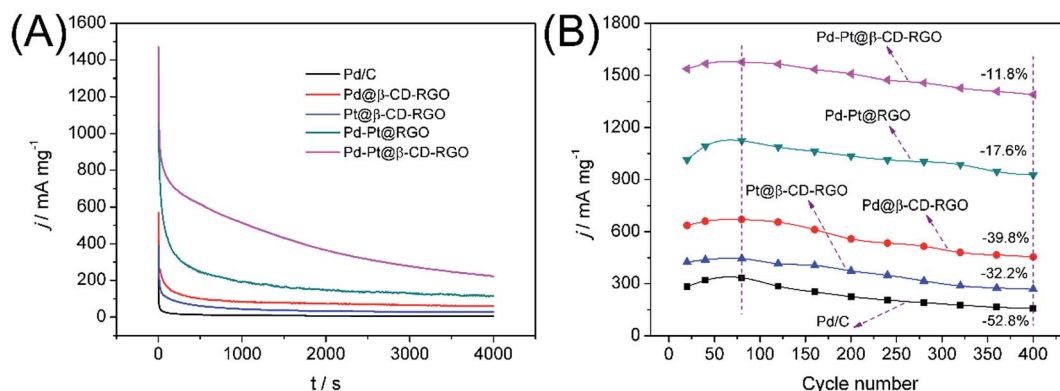


Fig. 6 (A) Chronoamperometric curves for methanol oxidation reactions catalyzed by Pd/C, Pd@β-CD-RGO, Pt@β-CD-RGO, Pd-Pt@RGO, and Pd-Pt@β-CD-RGO recorded at  $-0.2$  V (vs. SCE) for 4000 s in 1.0 M KOH containing 1.0 M methanol. (B) The durability performance of the Pd/C, Pd@β-CD-RGO, Pt@β-CD-RGO, Pd-Pt@RGO, and Pd-Pt@β-CD-RGO reflected by the forward peak current density based on 400 cycles in 1.0 M KOH containing 1.0 M methanol.

for methanol oxidation on Pd-Pt@β-CD-RGO modified electrode decreases by about 11.8% within the 400 scans. However, the peak current density on Pd-Pt@RGO, Pd@β-CD-RGO, Pt@β-CD-RGO, and commercial Pd/C modified electrodes declines about 17.6%, 39.8%, 32.2%, and 52.8% in the same scans. The good cycling stabilities may be attributed to the more durable nature of the {111} facet of Pd-Pt bimetallic nanoalloys and the synergetic effect of the Pd and Pt atoms,<sup>14,20</sup> which has been reported in the MOR with Pd-Pt bimetallic electrocatalysts. The TEM image of Pd-Pt@β-CD-RGO after durability was obtained as shown in Fig. S5.† From the TEM image, we can see that part of the Pd-Pt nanoclusters have begun to aggregate. This aggregation may result in the decrease of the activity of the Pd-Pt@β-CD-RGO catalyst. This implies that the small size and uniform dispersibility are indeed very important for the catalytic activity of the catalyst. Thus, we will further study on this problem in our lab for improving the durability of such catalyst.

## 4. Conclusions

In conclusion, we have developed a facile aqueous method to load Pd-Pt bimetallic nanoclusters with a uniform size of 2.0 nm on β-CD-RGO. The integration of β-CD and RGO was responsible for the formation of the monodispersed 2.0 nm Pd-Pt bimetallic clusters. The β-CD functionalization enhanced the affinity between supports and metal precursors, thus preventing aggregation and overgrowth of nanocrystals. Moreover, compared with commercial Pd/C catalyst, the β-CD-RGO supported Pd-Pt nanoclusters exhibited better electrocatalytic activities for methanol and ethanol oxidation. The Pd-Pt@β-CD-RGO nanocatalyst may also be used in other potential applications such as CO oxidation or hydrogenation reaction.

## Acknowledgements

This work was financially supported by the National Natural Science Foundation of China under the Grant No. 51361028, 51161025, 51661033, and 31660538.

## Notes and references

- 1 H.-H. Li, S.-Y. Ma, Q.-Q. Fu, X.-J. Liu, L. Wu and S.-H. Yu, *J. Am. Chem. Soc.*, 2015, **137**, 7862–7868.
- 2 X. M. Chen, G. H. Wu, J. M. Chen, X. Chen, Z. X. Xie and X. R. Wang, *J. Am. Chem. Soc.*, 2011, **133**, 3693–3695.
- 3 H. T. Qi, P. Yu, Y. X. Wang, G. C. Han, H. B. Liu, Y. P. Yi, Y. L. Li and L. Q. Mao, *J. Am. Chem. Soc.*, 2015, **137**, 5260–5263.
- 4 B. Liu, H. Q. Yao, W. Q. Song, L. Jin, I. M. Mosa, J. F. Rusling, S. L. Suib and J. He, *J. Am. Chem. Soc.*, 2016, **138**, 4718–4721.
- 5 R. J. White, R. Luque, V. L. Budarin, J. H. Clark and D. J. MacQuarrie, *Chem. Soc. Rev.*, 2009, **38**, 481–494.
- 6 Y. J. Wang, D. P. Wilkinson and J. Zhang, *Chem. Rev.*, 2011, **111**, 7625–7651.
- 7 H. J. Yin, H. J. Tang, D. Wang, Y. Gao and Z. Y. Tang, *ACS Nano*, 2012, **6**, 8288–8297.
- 8 H. P. Cong, X. C. Ren, P. Wang and S. H. Yu, *ACS Nano*, 2012, **6**, 2693–2703.
- 9 P. A. Khomyakov, G. Giovannetti, P. C. Rusu, G. Brocks, J. Van den Brink and P. Kelly, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2009, **79**, 195425.
- 10 Q. J. Wang and J. G. Che, *Phys. Rev. Lett.*, 2009, **103**, 066802.
- 11 I. Cabria, M. J. López and J. A. Alonso, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2010, **81**, 035403.
- 12 A. C. Chen and P. Holt-Hindle, *Chem. Rev.*, 2010, **110**, 3767–3804.
- 13 L. Wang, Y. Nemoto and Y. Yamauchi, *J. Am. Chem. Soc.*, 2011, **133**, 9674–9677.
- 14 A.-X. Yin, X.-Q. Min, Y.-W. Zhang and C.-H. Yan, *J. Am. Chem. Soc.*, 2011, **133**, 3816–3819.
- 15 S. J. Guo, S. J. Dong and E. K. Wang, *ACS Nano*, 2010, **4**, 547–555.
- 16 B. Lim and Y. N. Xia, *Angew. Chem., Int. Ed.*, 2011, **50**, 76–85.
- 17 Q. Yuan, Z. Y. Zhou, J. Zhuang and X. Wang, *Chem. Commun.*, 2010, **46**, 1491–1493.
- 18 X. Q. Huang, H. H. Zhang, C. Y. Guo, Z. Y. Zhou and N. F. Zheng, *Angew. Chem., Int. Ed.*, 2009, **48**, 4808–4812.





- 19 A.-X. Yin, X.-Q. Min, W. Zhu, H.-S. Wu, Y.-W. Zhang and C.-H. Yan, *Chem. Commun.*, 2012, **48**, 543–545.
- 20 D.-B. Huang, Q. Yuan, H.-H. Wang and Z.-Y. Zhou, *Chem. Commun.*, 2014, **50**, 13551–13554.
- 21 B. Lim, J. G. Wang, P. H. C. Camargo, M. J. Jiang, M. J. Kim and Y. N. Xia, *Nano Lett.*, 2008, **8**, 2535–2540.
- 22 Y. Chen, J. Yang, Y. Yang, Z. Y. Peng, J. H. Li, T. Mei, J. Y. Wang, M. Hao, Y. L. Chen, W. L. Xiong, L. Zhang and X. B. Wang, *Chem. Commun.*, 2015, **51**, 10490–10493.
- 23 W. Hong, C. S. Shang, J. Wang and E. K. Wang, *Energy Environ. Sci.*, 2015, **8**, 2910–2915.
- 24 B. Jiang, C. L. Li, J. Henzie, T. Takei, Y. Bando and Y. Yamauchi, *J. Mater. Chem. A*, 2016, **4**, 6465–6471.
- 25 Y. Q. Li, C. L. Li, B. P. Bastakoti, J. Tang, B. Jiang, J. Kim, M. Shahabuddin, Y. Bando, J. H. Kim and Y. Yamauchi, *J. Mater. Chem. A*, 2016, **4**, 9169–9176.
- 26 B. Lim, M. J. Jiang, P. H. C. Camargo, E. C. Cho, J. Tao, X. M. Lu, Y. M. Zhu and Y. N. Xia, *Science*, 2009, **324**, 1302–1305.
- 27 Z. M. Peng and H. Yang, *J. Am. Chem. Soc.*, 2009, **131**, 7542–7543.
- 28 Y. J. Guo, S. J. Guo, J. T. Ren, Y. M. Zhai, S. J. Dong and E. K. Wang, *ACS Nano*, 2010, **4**, 4001–4010.
- 29 G. B. Zhu, X. Zhang, P. B. Gai, X. H. Zhang and J. H. Chen, *Nanoscale*, 2012, **4**, 5703–5709.
- 30 Q. Zhang, Z. Y. Bai, M. Shi, L. Yang, J. L. Qiao and K. Jiang, *Electrochim. Acta*, 2015, **177**, 113–117.
- 31 Q. B. Zhang, J. P. Xie, J. H. Yang and J. Y. Lee, *ACS Nano*, 2009, **3**, 139–148.
- 32 Y. J. Xiong, J. M. McLellan, Y. D. Yin and Y. N. Xia, *Angew. Chem., Int. Ed.*, 2007, **46**, 790–794.
- 33 Z. Gao, W. L. Yang, J. Wang, H. J. Yan, Y. Yao, J. Ma, B. Wang, M. L. Zhang and L. H. Liu, *Electrochim. Acta*, 2013, **91**, 185–194.
- 34 Z. Fan, W. Kai, J. Yan, T. Wei, L. Zhi, J. Feng, Y. Ren, L. Song and F. Wei, *ACS Nano*, 2011, **5**, 191–198.
- 35 X. W. Mao, D. M. Tian and H. B. Li, *Chem. Commun.*, 2012, **48**, 4851–4853.
- 36 D. H. Du, P. C. Li and J. Y. Ouyang, *ACS Appl. Mater. Interfaces*, 2015, **7**, 26952–26958.
- 37 G. T. Fu, L. Tao, M. Zhang, Y. Chen, Y. W. Tang, J. Lin and T. H. Lu, *Nanoscale*, 2013, **5**, 8007–8014.
- 38 M. Chen, Y. Meng, J. Zhou and G. W. Diao, *J. Power Sources*, 2014, **265**, 110–117.
- 39 H. Huang, D. Sun and X. Wang, *J. Phys. Chem. C*, 2011, **115**, 19405–19412.

