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Branched PdAu nanowires with superior electrocatalytic formic acid oxidation activities†

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In this work, novel branched palladium–gold (PdAu) bimetallic nanowires supported on graphene (PdAu/graphene) were fabricated at relatively low temperatures for the first time by a one-pot co-reduction method with the aid of poly(diallyldimethyl-ammonium chloride). The structural characteristics of PdAu/graphene were confirmed by X-ray diffraction, X-ray photoelectron spectroscopy, transmission electron microscopy, energy-dispersive X-ray spectroscopy, high-angle annular dark-field scanning transmission electron microscopy and element mapping observations. The electrochemical properties of the PdAu/graphene were characterised by cyclic voltammetry and chronoamperometry (CA) in the formic acid oxidation reaction. Compared with a commercial Pd/C catalyst, the PdAu/graphene catalyst prepared in this work exhibited enhanced electrocatalytic activity including a higher oxidation current density, enhanced CO tolerance, and a more negative onset potential. Moreover, CA analyses demonstrated that PdAu/graphene shows higher stability than the Pd/C catalyst. The PdAu/graphene catalyst exhibited higher activity and strong poisoning resistance, which could be attributed to its special branched structure from PdAu nanowires.

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1. Introduction

Given increasing concerns over the environment and accelerated depletion of fossil fuels, over the past few decades the development of new energy sources as alternatives to fossil fuel has resurged. Fuel cells have attracted intense research activity due to ever-increasing environmental issues and the impending depletion of fossil fuels.^{1–7} Low temperature fuel cell types, including direct methanol fuel cells (DMFCs) and direct formic acid fuel cells (DFAFCs), present unique properties, such as low operating temperatures and relatively high energy densities. As such, these fuel cell types hold great promise as potential alternative power sources for portable electronic devices.^{8–10} DFAFCs operating near room temperature are potential power sources for stationary or portable electronic devices because of their relatively high electromotive force, relatively low fuel crossover flux through a Nafion membrane, competitive energy density and facile power-system integration.^{11–14} In general, the performance and efficiency of DFAFCs greatly depend on the intrinsic properties of the applied electrode catalysts, as well as the chemical and electrochemical stability of these materials in an electrolytic medium.

Most fuel cells at low temperatures use Pt as a catalyst.^{15–17} Pt-based alloys with other transition metals, especially nanostructured materials, exhibit higher performance because of the so-called bi-functional mechanism.^{18–20} However, the surface of Pt as an anode catalyst is usually heavily poisoned by the strong adsorption of CO intermediates, which are produced during the oxidation of organic fuels, thereby resulting in lower catalytic performance.^{21–23} Among the metal-based catalysts that have been studied to date, palladium (Pd) is considered a promising alternative because of its similar properties but much lower cost compared with platinum.²⁴ Recent studies revealed that nanoscale Pd can catalyse formic acid oxidation at the anode of a polymer electrolyte membrane fuel cell.²⁵ Oxidation of HCOOH on Pd mainly proceeds *via* the desired direct dehydrogenation pathway without forming poisonous CO species as a reaction intermediate. However, the catalytic performance of pure Pd is unstable over time, particularly with high concentrations of formic acid, which may be attributed to the progressive dissolution of Pd in the oxidized state. The addition of other metals to Pd/C, such as Au has beneficial effect on the oxidation of HCOOH.^{26–28} Suo and Hsing found that PdAu alloys exhibited higher Pd-specific activities towards the oxidation of HCOOH compared with individual Pd catalyst and easier removal of adsorbed oxygen species (*e.g.* O_{ads} or OH_{ads}) was observed from the surface of the PdAu alloy with higher Au content.²⁹ Yuan and Liu found the enhanced catalytic activity of PdAu nanoparticles resulted from the Pd site containing Au and the atomic distribution of the alloy.³⁰

For nanostructured electrocatalysts, catalytic performance is largely dependent on their surface structures, such as the

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surface facets, particle size, particle shape, *etc.* Xia and coworkers have shown that one-dimensional bimetallic nanowires exhibit enhanced electrocatalytic activity for anodic oxidation fuels because of the large specific areas.³¹ However, in the published studies, the seed-based epitaxial synthetic route is the most used method to prepare bimetallic nanowires. First the metal seeds were synthesized, and then the nanowires grew subsequently onto the metal core nanoparticles.^{32–34} Therefore, synthesis of PdAu bimetallic nanowires using a facile method is promising for designing novel nanostructured fuel cell electrocatalysts.

The catalytic performance of catalysts is also tightly associated with their support. Reduced graphene oxide (GO) with an interesting 2-dimensional plane structure has attracted considerable attention as a carbon support because of its convenient large-scale preparation. GO has been extensively studied in recent years because of its high conductivity (10^3 to 10^4 S m⁻¹), large theoretical surface area (calculated value, 2600 m² g⁻¹), unique graphitized basal plane structure, and potentially low manufacturing cost.^{35–37} Functionalized GO has also attracted special attention because it enables customization of the properties of the hybrids of nanoparticles and graphene while preserving nearly all of the intrinsic properties of graphene.^{38,39} Functionalized GO has been demonstrated to be a promising candidate material for supporting bimetallic nanoparticles (*e.g.* PtRu,⁴⁰ PtPd,⁴¹ and PtCu⁴²) in fuel cells.

In this study, we report a facile, efficient, and controllable route to disperse branched palladium–gold (PdAu) nanowires on reduced GO with the aid of poly(diallyldimethyl-ammonium chloride) (PAC). Adsorption of PAC on GO can lead to a highly dense and homogeneous distribution of positive charges on the GO surface and preserve the intrinsic properties of GO without any chemical oxidation treatment. The structure of PdAu/graphene was characterised in detail by X-ray diffraction (XRD), X-ray photoelectron spectroscopy (XPS), transmission electron microscopy (TEM), high-resolution transmission electron microscopy (HR-TEM), energy-dispersive X-ray spectroscopy (EDS), high-angle annular dark-field scanning transmission electron microscopy (HAADF-STEM), and element mapping. The electrocatalytic activity of PdAu/graphene during formic acid oxidation was examined by electrochemical cyclic voltammetry (CV) and chronoamperometry (CA), and results showed that the synthesised PdAu/graphene exhibits better activity including a higher current density and durability than commercial Pd/C.

2. Experiments

2.1. Reagents

Graphite flake (99.8%, 325 mesh) was provided by Alfa. Poly(diallyldimethyl-ammonium chloride) (PAC, 20 wt% solution in water, average molecular weight 200–350 K) was purchased from Aladdin Chemical Company, China. Analytical grade HAuCl₄ and Na₂PdCl₄ were obtained from Tianjin Chemical Reagents, China. NaNO₃, H₂SO₄, KMnO₄ and other chemicals were of analytical grade and used without further purification. A 0.02 mol L⁻¹ HAuCl₄ solution was prepared by

dissolving HAuCl₄·4H₂O powder in twice-distilled water. High-purity nitrogen and twice-distilled water were used in all experiments. Sodium borohydride (NaBH₄) was purchased from commercial sources in China and used as supplied. All solutions were prepared with deionized water.

2.2. Preparation of PAC-functionalized graphene

The GO was firstly synthesized according to a modified Hummers method.^{43,44} Then GO (40 mg) was sonicated in 50 mL distilled water for 1 h, and then 0.1 g of PAC was added. The solution was sonicated for another 1.5 h. Fastly, the suspension was centrifuged to isolate the solid product which was then washed with distilled water. This washing process was repeated three times to remove excess PAC. The PAC-functionalized GO powders were dried in a vacuum oven at 60 °C for 24 h and the final product is denoted as PAC-functionalized GO.

2.3. Preparation of PdAu/graphene

20 mg PAC-functionalized GO was dissolved in 50 mL water by ultrasonic treatment 2 h. Then 3.8 mg Na₂PdCl₄ and 650 μL 0.02 M HAuCl₄ were added into the mixture, and the solution was sonicated for 0.5 h. Then an aqueous solution of NaBH₄ (40 mL, 0.05 M) in a water-ice bath was then slowly added dropwise into the above solution under magnetic stirring, and the color of mixture changed from yellow to black. After continuous reaction for 2 h under stirring, the black product was collected by centrifugation through washing with ethanol and deionized water (*V* : *V* = 1 : 1) to remove the ions possibly remaining in the final products, and then vacuum-dried 24 h at 60 °C. For Comparison, Pd/graphene was prepared with a similar method for PdAu/graphene.

2.4. Characterization

X-ray diffraction (XRD) analysis was carried out on Bruker D8-ADVANCE diffractometer with Cu K α radiation (λ = 1.5406 Å, 40 kV and 40 mA). The composition of the product was measured by the inductively coupled plasma-atomic emission spectrometer (ICP-AES; USA Thermo Jarrell-Ash Corp. ICP-9000 (N + M)). Field-emission scanning electron microscopy (FESEM; JEOL, JSM-7500F, 8 kV) was used to analyze the morphology of the samples. Transmission electron microscopy (TEM), energy-dispersive X-ray (EDS) and EELS elemental mapping observations were performed on a Philips Tecnai F20 microscope, working at 200 kV. All samples subjected to TEM measurements were dispersed in ethanol ultrasonically and were dropped on copper grids. X-ray photoelectron spectroscopy (XPS) measurement was conducted with Kratos Axis Ultra DLD spectrometer with an Al K α radiation as the X-ray source ($h\nu$ = 1486.7 eV). The peak positions were corrected for sample charging by setting the C 1s binding energy at 284.8 eV.

2.5. Electrochemical measurements

All electrochemical measurements were carried out with a CHI 660D electrochemical workstation system (CH Instruments, Chenhua Co., Shanghai, China) and a conventional three-



electrode system. The working electrode was a Glassy Carbon Electrode (GCE) with a diameter of 3 mm. Ag/AgCl (in saturated KCl) electrode was employed as reference *via* a salt bridge and a Pt wire was used as counter electrode. Prior to use, the glassy carbon electrode was first polished subsequently with a 1.0, 0.3, and 0.05 μm alumina slurry and was ultrasonically cleaned with 1 : 1 nitric acid, ethanol and distilled water, and then dried in a stream of nitrogen gas. For electrochemical measurements, 4 mg of catalyst were dispersed in 1 mL ethanol under ultrasonication for 1 h and 10 μL of the catalyst suspension was dropped onto the GCE. After drying in air for 2 hours, a 5 μL Nafion (Sigma Aldrich) solution (0.2%) was added, followed by drying at ambient temperature. The dried GCE were cleaned again by sequentially washing with water. Prior to electrochemical measurements, the electrolytes were deoxygenated by bubbling with N_2 for 30 min. The formic acid oxidation reaction (FAOR) experiments were carried out by sweeping the potential from -0.20 V to $+0.90$ V at a scan rate of 50 mV s^{-1} in a solution containing 0.5 M H_2SO_4 and 0.5 M HCOOH . CA curves were recorded at $+0.2$ V in a solution containing 0.5 M H_2SO_4 and 0.5 M HCOOH for 7 h to investigate the durability of the catalysts. Prior to CO adsorption, the electrolytes were deoxygenated by bubbling with N_2 for 30.0 min. In the CO stripping experiments, CO was adsorbed on the pre-cleaned electrode by holding the potential at 0.10 V for 5.0 min while bubbling CO gas through the electrolyte solution. CO dissolved in the electrolyte solution and was removed by purging nitrogen through the solution for 15.0 min. All electrochemical experiments were carried out at room temperature.

For the electrochemical surface area (ECSA) study, results were calculated by integrating the charge passing the electrode during palladium oxide reduction and the CO stripping. For example, ECSA values were calculated by integrating the area under curves for the oxygen adsorption charge (Q_{O}) region between -0.2 V and $+0.4$ V of the reverse sweep in the CV and the Q_{O} can be determined using $Q_{\text{O}} = 0.5 \times Q$, where Q is the charge in the oxygen adsorption/desorption area obtained after double-layer correction. The specific ECSA was calculated based on the following relation:

$$\text{Specific ECSA} = Q_{\text{O}}/m \times q_{\text{O}} \quad (1)$$

where m is the loading amount of metal, q_{O} is a conversion factor of 424 $\mu\text{C cm}^{-1}$ for the adsorption of a monolayer of oxygen on a Pd surface, and Q_{O} is the charge for hydrogen adsorption, which can be calculated from the area under the CV curve by integrating the charge passing the electrode during palladium oxide reduction after the correcting for the double layer formation.

3. Results and discussion

3.1. Physicochemical properties of the samples

The morphology and structure of the as-synthesized materials were characterised by TEM (Fig. 1). Fig. 1A and B show TEM micrographs of the final products obtained from one-pot preparation methods at different magnifications. In Fig. 1A,

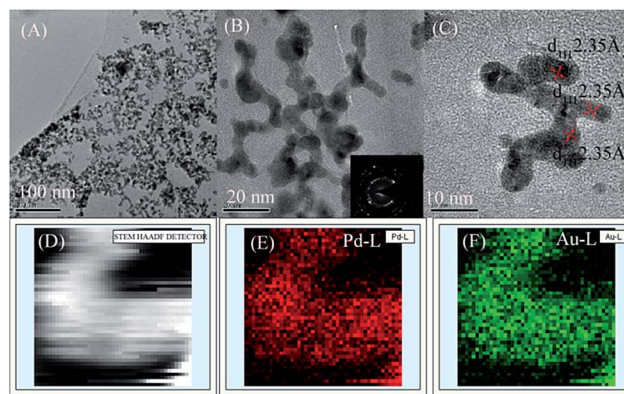


Fig. 1 (A, B) TEM micrographs of PdAu/graphene at different magnifications. The scale bars are (A) 100 nm and (B) 20 nm, inset of (B) is SAED of PdAu/graphene. (C) High-resolution TEM micrographs (D) the high angle annular dark field (HAADF) STEM image of PdAu and the corresponding elemental mapping of Pd (E) and Au (F) in the PdAu alloy.

several branched PdAu nanowires formed, and PdAu intermetallics were well-dispersed on graphene. Upon closer examination at higher magnification (Fig. 1B), a high yield of distinct nanobranched structures were observed from the proposed one-pot preparation method. The selected-area electron-diffraction (SAED) pattern of the product (inset of Fig. 1B) illustrates that the PdAu nanowires were crystalline in nature. Fig. 1C depicts an HRTEM micrograph of the branched PdAu nanowires. Here, the worm-like nanowires were well dispersed and a well-resolved lattice fringe can be clearly observed. Careful measurements revealed that primarily one type of lattice fringe with an interplanar spacing of 2.35 Å, which can be ascribed to the Au (111) plane, was present in the sample. To investigate the crystal structure and the element distribution of the branched PdAu nanowires, HAADF-STEM analysis was performed. Fig. 1D–F show the HAADF-STEM micrographs and selected-area element analysis maps of Pd and Au. The element maps revealed that Pd and Au were homogeneously dispersed in the PdAu nanobranched structures, thereby indicating the formation of an alloy structure. The HRTEM and elements analysis results strongly suggested that the PdAu alloy nanoparticles were first formed, after which the branched structure was developed. The STEM-EDS line-scan profiles (Fig. S1A–D†) of two regions of the sample showed that the distribution of Pd and Au in the branched nanowires coincided well with their alloy structures, and agreed with the results of HAADF-STEM. Fig. S2A–C† show the Pd/graphene TEM, HRTEM, and EDS results, where Pd was observed to form branched nanowires.

The EDS of PdAu/graphene shown in Fig. 2A, revealed that both Pd and Au, in addition to the Cu substrate, are present in the branched nanowires. EDS revealed that the average atomic ratio of Pd : Au was $0.45 : 0.55$, which agrees well with the value (Pd : Au = $0.48 : 0.52$) obtained by inductively coupled plasma-atomic emission spectrometry (ICP-AES). The weight percentage of metal species in the PdAu/graphene, Pd/graphene and Pd/C were measured to be approximately 26, 22.3 and 17.3 wt%, respectively with thermogravimetric analysis (TGA, Fig. S3†).



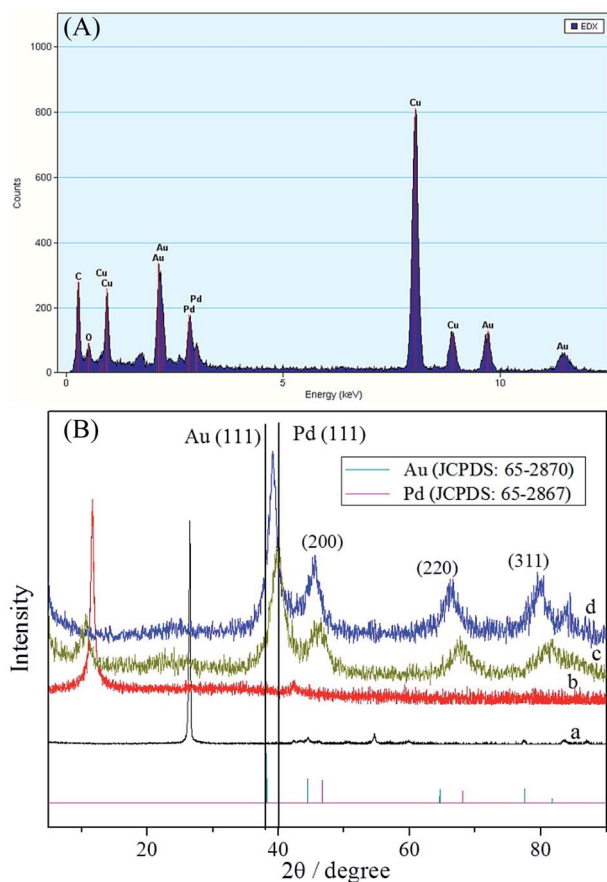


Fig. 2 (A) EDS of PdAu/graphene. (B) XRD patterns of (a) graphite, (b) GO, (c) Pd/graphene, and (d) PdAu/graphene.

The crystalline phase of the PdAu nanowires was studied by powder XRD measurements. The XRD patterns of GO and PdAu/graphene are shown in curves *b* and *d* of Fig. 2B. The X-ray (002) peak of graphite is observed at 2θ value of 26.5° (curve *a*, Fig. 2B). This peak shifted to 10.0° (curve *b*, Fig. 2B) after oxidation treatment. Once PdAu/graphene was formed, the 2θ value of (002) shifted to approximately 23.6° (curve *d*, Fig. 2B). Therefore, GO was reduced to graphene after the reduction process *via* sodium borohydride. XRD analysis revealed the better crystallinity of PdAu/graphene (Fig. 2B, curve *d*), the four representative diffraction peaks corresponding to the (111), (200), (220), and (311) planes of the PdAu alloy. Pd/graphene showed clear diffraction peaks in curve *c*. All of the diffraction peaks from PdAu/graphene were located between the expected positions of pure Pd (no. 65-2867) and Au (no. 65-2870). This XRD phenomenon has also been observed in other bimetallic alloy nanostructures.^{45–47}

XPS measurements were performed to investigate C 1s, Pd 3d, and Au 4f. The high-resolution spectra of the C 1s peak are shown in Fig. 3A: four components namely, C–C (284.5 eV), C–O (286.3 eV), C=O (288.7 eV), and O=C–O (289.7 eV) could be observed. The intensities of carbon atoms became predominant compared to the C 1s for GO (inset pattern of Fig. 3A), whereas the intensities of carbon atoms bonded to oxygen were obviously reduced. The XPS spectra of the Au 4f and Pd 3d regions of

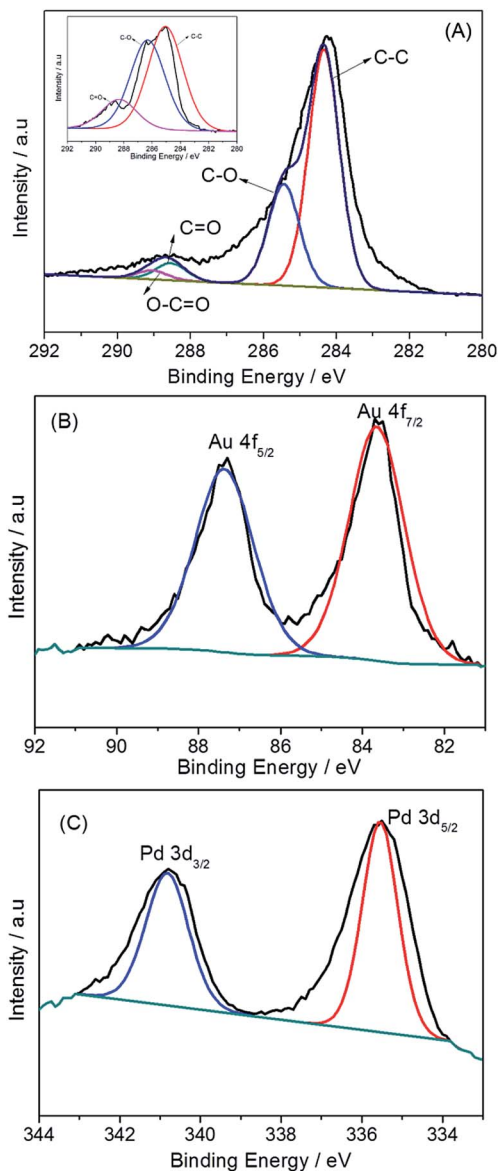
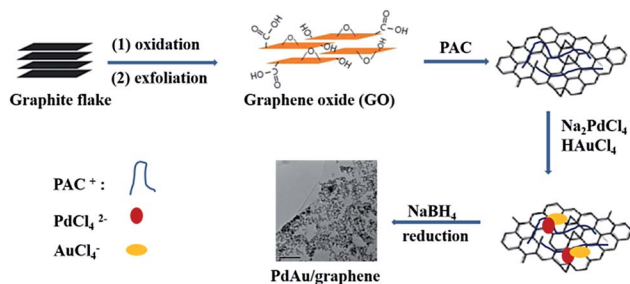


Fig. 3 XPS spectrum of PdAu/graphene in (A) C 1s regions (B) Au 4f regions and (C) Pd 3d regions. Inset of (A) shows the C 1s XPS pattern of GO.

PdAu/graphene were shown in Fig. 3B and C, respectively. The results revealed the binding states of Au^{III} and Pd^{II} could not be observed, thereby demonstrating the Au^{III} and Pd^{II} precursors were successfully reduced in our synthesis. The two 4f zero valence Au⁰ were identified to be 4f_{7/2} and 4f_{5/2} and were located at 83.7 eV and 87.4 eV, respectively. By comparison, the Pd⁰ 3d_{5/2} and 3d_{3/2} states were located at 335.6 and 340.8 eV, respectively. This trend indicates a small positive shift relative to pure polycrystalline Au (Au 4f_{7/2} at 82.2 eV and Au 4f_{5/2} at 86.1 eV), which confirms the expected electronic interaction between Au and Pd upon the formation of an intermetallic compound.

Based on these experiments, we suggest a formation mechanism of branched PdAu nanowires on GO (Scheme 1). The steps of the synthetic route of the PdAu/graphene are as follows: first,





Scheme 1 Synthetic scheme of nanobranched palladium-gold nanowires supported on graphene.

the GO nanosheets are functionalised with amine moieties carrying PAC, thereby producing a large amount of positively charged sites on the GO sheets. The amine groups function as preferential reactive sites in electrostatic conjunction with PdCl_4^{2-} and AuCl_4^- . Finally, the reduction of the Pd and Au precursors by NaBH_4 causes the formation of nanoparticles, and GO is simultaneously reduced to graphene. The PAC layer is a polyamine with fine biocompatibility, and abundant amine that help stabilize the metal nanoparticles. Moreover, the sites of the longer PAC chains cause PdAu to form branched nanowires.

3.2. Electrocatalytic behavior of the PdAu/graphene catalyst

The ECSA of an electrocatalyst can provide important information on the availability of its active sites. The CV curves of the PdAu/graphene and Pd/C electrocatalysts in a 0.5 M H_2SO_4 solution were in the range of -0.2 V to 1.2 V (vs. Ag/AgCl), with a sweep rate of 0.1 V s^{-1} . The CV curves show potential regions in the ranges -0.2 – 0.15 V, 0.15 – 0.4 V and 0.4 – 1.0 V which are characteristics of hydrogen adsorption-desorption, a double layer charge, and the formation and reduction of PdO, respectively. The ECSA was calculated by integrating the charge passing the electrode during palladium oxide reduction and CO stripping (by assuming a q_0 value of 424 $\mu\text{C cm}^{-2}$).⁴⁸ Fig. 4A also demonstrates that the area of the reduced peak of the palladium oxide for PdAu/graphene, Pd/graphene, and Pd/C, respectively. The ECSA per unit weight of Pd was estimated from the palladium oxide reduction charge to be 99.6 $\text{m}^2 \text{g}^{-1}$ Pd for PdAu/graphene and 93.6 $\text{m}^2 \text{g}^{-1}$ Pd for Pd/graphene, both of which are higher than that of the Pd/C catalyst (79.3 $\text{m}^2 \text{g}^{-1}$ Pd, Fig. 4B). By calculating the stripping charges of pre-adsorbed CO, the ECSA of PdAu/graphene, Pd/graphene and Pd/C were 40.6 $\text{m}^2 \text{g}^{-1}$, 43.5 $\text{m}^2 \text{g}^{-1}$ and 51.8 $\text{m}^2 \text{g}^{-1}$, respectively.

The large ECSA of the PdAu/graphene catalyst may be attributed to its branched nanostructure. The loss of ECSA with cycling was plotted in Fig. S5.† After 1000 cycles, the PdAu/graphene catalyst lost only 25% of its initial ECSA, while the commercial Pd/C catalyst lost 50% of its initial ECSA.

The catalytic activity of PdAu/graphene, Pd/graphene, and Pd/C in formic acid oxidation was investigated. To compare the mass activity of different catalysts, the kinetic current was calculated from the formic acid oxidation curve by mass-transport correction. Fig. 5A shows the steady CV curves of formic acid oxidation with PdAu/graphene and commercial Pd/C in

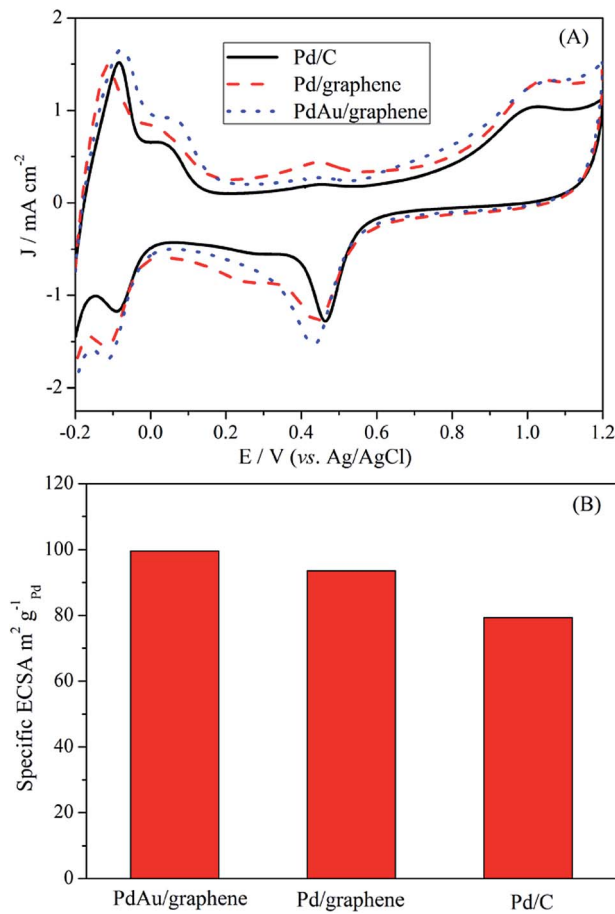


Fig. 4 (A) CV profiles of commercial Pd/C, Pd/graphene and PdAu/graphene catalysts in a 0.5 M H_2SO_4 solution. (B) Specific ECSA for the PdAu/graphene, Pd/graphene and commercial Pd/C catalysts.

0.5 M H_2SO_4 /0.5 M HCOOH at a potential scan of 50 mV s^{-1} . The voltammetric currents were normalized to the actual active surface area (*i.e.* the ECSA by calculating the stripping charges of pre-adsorbed CO) of the respective electrodes. The current density of the anode peak (J_a) was larger than that of the cathodic peak (J_c) during HCOOH electro-oxidation by PdAu/graphene, thereby qualitatively indicating the improved CO tolerance of the Pd nanomaterials compared with that of Pd-based catalysts.^{49,50} At least two aspects of the CV formic acid oxidation curve at the PdAu/graphene electrode warrant special attentions. First, the onset potential of formic acid oxidation on PdAu/graphene was more negative than that on the Pd/C catalyst. Second, the current density of formic acid oxidation in the forward and reverse potential sweeps obtained from the Pd catalysts was much higher than that obtained from the commercial Pd/C catalysts. The current density and onset potential results indicated that PdAu/graphene has much better catalytic activity than the commercial Pd/C catalyst. The specific activity of the kinetic current was calculated from the ECSA and mass activity. The peak current densities of formic acid oxidation were: PdAu/graphene > Pd/graphene > Pd/C.

To further explore the observed enhancements in the electrocatalytic activity and CO-tolerance of the proposed catalyst,



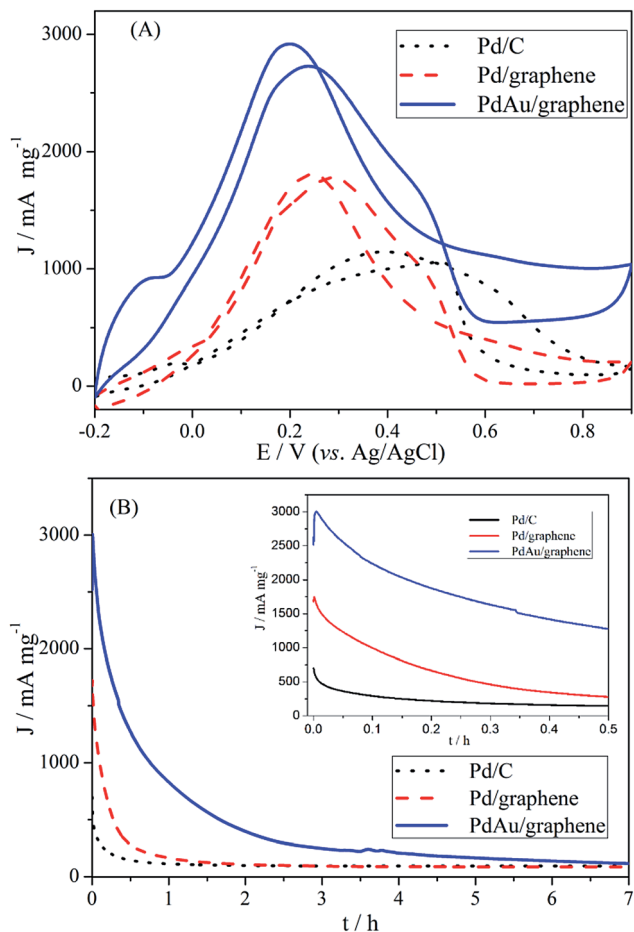


Fig. 5 (A, B) CV and CA results of HCOOH on PdAu/graphene, Pd/graphene and Pd/C catalysts (50 mV s^{-1}) for formic acid electro-oxidation in $0.5 \text{ M H}_2\text{SO}_4/0.5 \text{ M HCOOH}$ solutions. CA curves were recorded at 0.2 V .

the activity and stability of PdAu/graphene, Pd/graphene and Pd/C catalysts in formic acid oxidation were evaluated by CA in $0.5 \text{ M H}_2\text{SO}_4/0.5 \text{ M HCOOH}$. The CA curves at a potential of $+0.20 \text{ V}$ for the PdAu/graphene, Pd/graphene, and Pd/C electrodes are shown in Fig. 5B. The maximum initial and steady-state current densities obtained with PdAu/graphene were much larger than that on Pd/C, which agreed well with the observed CV curves (Fig. 5A). According to the literature, Pd/C catalysts suffer from poor durability in formic acid oxidation, although they show higher catalytic activity. Pd electrocatalysts could easily aggregate, dissolve from the carbon support into the electrolyte, or become poisoned by the intermediates, thereby resulting in poor durability.⁵¹ We also performed TEM on PdAu/graphene after 300 electrochemical cycles. The structure of the PdAu nanowires did not change and aggregated partly (Fig. S4†). The comparison of the catalytic activities to the previously reported results are shown in Table S1.† Compared to some earlier catalysts, which were Pd-based catalysts, the PdAu/graphene catalysts investigated in this work demonstrated enhanced activity.

Accordingly, the long-term durability of the electrocatalyst was evaluated by continuously scanning the CV curves to follow

the changes in the anode peak current density with cyclic number, as shown in Fig. 6. Similar to the literature,^{52,53} the peak current density of commercial Pd/C sharply decreased and only 20% of the initial current remained after 60 cycles, thereby confirming the poor durability of Pd/C (Fig. 6). By contrast, the peak current density of PdAu/graphene initially increased (*via* electrochemically activation) and decreased to 40% after 300 cycles, thereby showing that the PdAu/graphene is relatively more stable.

CO stripping experiments were also performed and the results are shown in Fig. 7. For the Pd/C catalyst, a sharp peak was observed at approximately 0.69 V . Compared with that of the Pd/C catalyst, the CO stripping voltammograms of the PdAu/graphene and Pd/graphene catalysts presented sharp peaks with potentials of 0.60 and 0.67 V , respectively, thereby suggesting higher CO oxidation activity on the PdAu/graphene and Pd/graphene catalysts because of their slightly more negative

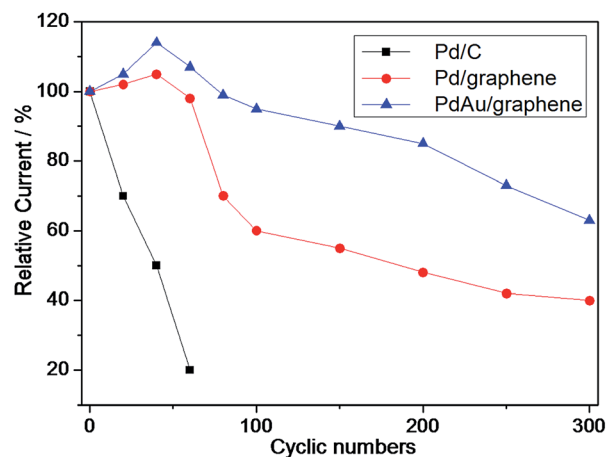


Fig. 6 Durability of PdAu/graphene and Pd/C electrocatalysts for formic acid electro-oxidation in $0.5 \text{ M H}_2\text{SO}_4 + 0.5 \text{ M HCOOH}$ solutions.

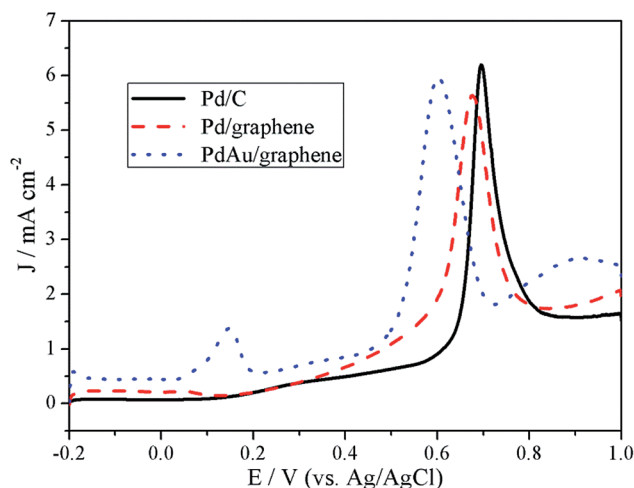


Fig. 7 CO stripping curves of the Pd/C, PdAu/graphene and Pd/graphene catalysts in $0.5 \text{ M H}_2\text{SO}_4$.



potential. Furthermore, the onset potentials of PdAu/graphene and Pd/graphene catalysts were lower than that of Pd/C, thus indicating the higher CO tolerance of the former compared with that of the latter.

The enhanced electrocatalytic performance of branched PdAu nanowires for formic acid oxidation should be attributed to the following reasons. (1) Special nanostructures: the branched nanowires networks can improve electron transport characteristics during the catalysis. What's more, graphene can offer high electron conductivity and a large specific surface. (2) The Au co-metals on Pd can provide an opportunity to improve the adsorption of OH_{ads} onto the surface of the catalyst. Additionally, the Au can result in higher CO tolerance on the surface of the catalyst and thus activate the catalyst's surface and help to enhance the formic acid process.

4. Conclusions

Branched PdAu bimetallic nanowires enclosed by a stepped surface supported on reduced GO (PdAu/graphene) catalysts with the aid of PAC were prepared using a one-pot method. The nanostructures and electronic properties of the catalysts were investigated with XRD, XPS, TEM, HR-TEM, and HAADF-TEM. Based on its electrocatalytic performance for formic acid electro-oxidation, PdAu/graphene is a highly durable and more efficient electrocatalyst compared with commercial Pd/C in formic acid electro-oxidation. The improved performance of PdAu/graphene may be attributed to its PdAu alloy structure and the increased electrical conductivity of its graphene support. The superior overall electrocatalytic performance, especially the higher CO tolerance, suggests that a PdAu/graphene catalyst could be a promising anode catalyst for fuel cell applications. The enhanced performance of the proposed catalyst for the formic acid oxidation could be attributed to its special structure, which includes branched nanowires enclosed by a stepped surface and possible synergetic effects among its Pd and Au components. Further product analyses by *in situ* FTIR and HPLC are required to better understand the enhancement effects of the branched PdAu/graphene nanowires.

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Notes and references

- 1 R. Elosato, G. Cordaro, D. Stucchi, C. Cristiani and G. Dotelli, *J. Power Sources*, 2015, **298**, 46–67.
- 2 J. K. Wang, J. Zhou, T. Wang, G. Chen, K. Wu and Y. H. Cheng, *J. Alloys Compd.*, 2016, **689**, 581–586.
- 3 Z. J. He, H. Li and E. Birgersson, *Renewable Energy*, 2016, **85**, 1301–1305.

- 4 G. R. Molaeimanesh, H. S. Googarchin and A. Q. Moqaddam, *Int. J. Hydrogen Energy*, 2016, **41**, 22221–22245.
- 5 A. Baldinelli, L. Barelli and G. Bidni, *Energy*, 2015, **90**, 2070–2084.
- 6 L. Duclos, L. Svecova, V. Laforest, G. Mandil and P. X. Thivel, *Hydrometallurgy*, 2016, **160**, 79–89.
- 7 S. Dresp, F. Luo, R. Schmack, S. Kuhl, M. Gliech and P. Strasser, *Energy Environ. Sci.*, 2016, **9**, 2020–2024.
- 8 M. M. Liu, S. J. He and W. Chen, *Electrochim. Acta*, 2016, **199**, 218–226.
- 9 S. Zhang, Y. Shao, G. Yin and Y. Lin, *Angew. Chem., Int. Ed.*, 2010, **49**, 2211–2214.
- 10 Z. Q. Niu, D. S. Wang, R. Yu, Q. Peng and Y. D. Li, *Chem. Sci.*, 2012, **3**, 1925–1929.
- 11 Y. Z. Lu, Y. Y. Jiang, H. B. Wu and W. Chen, *J. Phys. Chem. C*, 2013, **117**, 2926–2938.
- 12 Z. L. Xin, S. H. Wang, J. Wang, X. Huang, X. B. Ji, Y. Y. Yao and L. D. Shao, *Electrochem. Commun.*, 2016, **67**, 26–30.
- 13 Z. L. Xin, J. Wang, X. Huang, Y. Y. Yao and L. D. Shao, *J. Electrochem. Soc.*, 2015, **162**, H898–H902.
- 14 M. D. Obradovic and S. L. Gojkovic, *Electrochim. Acta*, 2013, **88**, 384–389.
- 15 L. Zhang, L. Wang, C. M. B. Holt, B. Zahiri, Z. Li, K. Malek, T. Navessin, M. H. Eikerling and D. Mitlin, *Energy Environ. Sci.*, 2012, **5**, 6156–6172.
- 16 C. Alegre, D. Sebastian, M. E. Galvez, R. Moliner and M. J. Lazaro, *Appl. Catal., B*, 2016, **192**, 260–267.
- 17 N. Doan, M. C. Figueiredo, C. Johans and T. Kallio, *Int. J. Electrochem. Sci.*, 2016, **11**, 7631–7643.
- 18 Y. X. Wang, H. J. Zhou, P. C. Sun and T. H. Chen, *J. Power Sources*, 2014, **245**, 663–670.
- 19 X. L. Sui, D. M. Gu, Z. B. Wang, J. Liu, L. Zhao and L. M. Zhang, *RSC Adv.*, 2016, **6**, 79254–79262.
- 20 C. H. Wan, M. T. Lin and B. F. Su, *Int. J. Hydrogen Energy*, 2016, **39**, 7447.
- 21 S. J. Yoo, T. Y. Jeon, K. S. Kim, T. H. Lim and Y. E. Sung, *Phys. Chem. Chem. Phys.*, 2010, **12**, 15240–15246.
- 22 A. X. Yin, X. Q. Min, W. Zhu, W. C. Liu, Y. W. Zhang and C. H. Yan, *Chem.–Eur. J.*, 2012, **18**, 777–782.
- 23 J. Kugai, T. Moriya, S. Seino, T. Nakagawa, Y. Ohkubo, H. Nitani, H. Daimon and T. A. Yamamoto, *Int. J. Hydrogen Energy*, 2012, **37**, 4787–4797.
- 24 C. Bianchini and P. K. Shen, *Chem. Rev.*, 2009, **109**, 4183–4206.
- 25 D. D. Sun, L. Si, G. T. Fu, C. Liu, D. M. Sun, Y. Chen, Y. W. Tang and T. H. Lu, *J. Power Sources*, 2015, **280**, 141–146.
- 26 S. G. da Silver, J. C. M. Silva, G. S. Buzzo, E. V. Spinace, A. O. Neto and M. H. M. T. Assumpcao, *Electrocatalysis*, 2015, **6**, 442–446.
- 27 J. Nandenha, R. F. B. de Souza, M. H. M. T. Assumpcao, E. V. Spinace and A. O. Neto, *Ionics*, 2013, **19**, 1207–1213.
- 28 S. X. Zhang, M. Qing, H. Zhang and Y. N. Tian, *Electrochem. Commun.*, 2009, **11**, 2249–2252.
- 29 Y. Suo and I.-M. Hsing, *Electrochim. Acta*, 2011, **56**, 2174–2183.
- 30 D. W. Yuan and Z. R. Liu, *J. Power Sources*, 2013, **224**, 241–249.



- 31 Y. Wang, S. I. Choi, X. Zhao, S. F. Xie, H. C. Peng, M. F. Chi, C. Z. Huang and Y. N. Xia, *Adv. Funct. Mater.*, 2014, **24**, 131–139.
- 32 Q. Yuan, Z. Y. Zhou, J. Zhuang and X. Wang, *Chem. Mater.*, 2010, **22**, 2395–2402.
- 33 S. J. Guo, S. J. Dong and E. K. Wang, *Energy Environ. Sci.*, 2010, **3**, 1307–1310.
- 34 X. W. Teng, M. Feygenson, Q. Wang, J. Q. He, W. X. Du, A. I. Frenkel, W. Q. Han and M. Aronson, *Nano Lett.*, 2009, **9**, 3177–3184.
- 35 Y. P. Yang, P. Cheng and S. P. Huang, *J. Alloys Compd.*, 2016, **688**, 1172–1180.
- 36 W. L. Wang, Q. H. Zhang, J. L. Li, X. Liu, L. J. Wang, J. J. Zhu, W. Luo and W. Jiang, *RSC Adv.*, 2015, **5**, 8988–8995.
- 37 L. Li, Y. Y. Dou, L. F. Wang, M. Luo and J. Liang, *RSC Adv.*, 2014, **4**, 25658–25665.
- 38 J. D. Qiu, G. C. Wang, R. P. Liang, X. H. Xia and H. W. Yu, *J. Phys. Chem. C*, 2011, **115**, 15639–15645.
- 39 S. D. Yang, C. M. Shen, X. J. Lu, H. Tong, J. J. Zhu, X. G. Zhang and H. J. Gao, *Electrochim. Acta*, 2012, **62**, 242–249.
- 40 H. Wang, J. Du, Z. Yao, R. Yue, C. Zhai, F. Jiang, Y. Du, C. Wang and P. Yang, *Colloids Surf., A*, 2013, **436**, 57–61.
- 41 S. S. Li, J. J. Lv, Y. Y. Hu, J. N. Zheng, J. R. Chen, A. J. Wang and J. J. Feng, *J. Power Sources*, 2014, **247**, 213–218.
- 42 F. Li, Y. Guo, Y. Liu, H. Qiu, X. Sun, W. Wang, Y. Liu and J. Gao, *Carbon*, 2013, **64**, 11–19.
- 43 N. I. Kovtyukhova, P. J. Ollivier, B. R. Martin, T. E. Mallouk, S. A. Chizhik, E. V. Buzaneva and A. D. Gorchinskiy, *Chem. Mater.*, 1999, **11**, 771–778.
- 44 W. S. Hummers and R. E. Offeman, *J. Am. Chem. Soc.*, 1958, **80**, 1339.
- 45 B. N. Wanjala, J. Luo, R. Loukrakpam, B. Fang, D. Mott, P. N. Njoki, M. Engelhard, H. R. Naslund, J. K. Wu, L. C. Wang, O. Malis and C. J. Zhong, *Chem. Mater.*, 2010, **22**, 4282–4294.
- 46 J. C. Huang, Z. L. Liu, C. B. He and L. M. Gan, *J. Phys. Chem. B*, 2005, **109**, 16644–16649.
- 47 Y. Z. Lu and W. Chen, *Chem. Commun.*, 2011, **47**, 2541–2543.
- 48 M. Hara, U. Linke and T. Wandlowski, *Electrochim. Acta*, 2007, **52**, 5733–5748.
- 49 W. Chen, J. Kim, S. H. Sun and S. W. Chen, *Phys. Chem. Chem. Phys.*, 2006, **8**, 2779–2786.
- 50 W. Chen, J. M. Kim, S. H. Sun and S. W. Chen, *Langmuir*, 2007, **23**, 11303–11310.
- 51 S. Y. Wang and A. M. Manthiram, *Electrochim. Acta*, 2013, **88**, 565–570.
- 52 X. Yu and P. G. Pickup, *Electrochem. Commun.*, 2009, **11**, 2012–2014.
- 53 X. Yu and P. G. Pickup, *J. Power Sources*, 2009, **187**, 493–499.

