Encapsulating Pd Nanoparticles in Double-Shelled Graphene@Carbon Hollow Spheres for Excellent Chemical Catalytic Property

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Double-shelled hollow carbon spheres with reduced graphene oxide (RGO) as inner shell and carbon (C) layer as outer shell have been successfully designed and prepared. This tailor-making structure acts as an excellent capsule for encapsulating with ultrafine Pd nanoparticles (Pd NPs), which could effectively prevent Pd NPs from aggregation and leaching. As a result, the as-obtained RGO@Pd@C nanohybrid exhibits superior and stable catalytic performance. With the aid of RGO@Pd@C, the reduction reaction of 4-nitrophenol (4-NP) to 4-aminophenol with NaBH₄ as reducing agent can be finished within only 30 s, even the content of Pd is as low as 0.28 wt%. As far as we know, RGO@Pd@C is one of the most effective catalyst for 4-NP reducing reaction up to now.

Noble metals have attracted considerable attention because of their potential applications in catalysis, gas sensor, energy conversion, and fuel cells¹–³. However, the high surface energy of noble metal nanoparticles (NPs) makes them easily agglomerated or shape-changed during catalytic reactions, which results in the dramatic decrease of their activity and selectivity⁴–⁵. To avoid aggregation, various isolated nanoreactors are employed to restrict these NPs⁶–⁸. Among them, the design of yolk-shell structure has been proven to be an effective strategy to prevent the aggregation of NPs. Recently, yolk/shell structured nanohybrid composed of metal particles and silica/carbon (C) shells have become the research hotspot in chemistry due to its widely applications in confined nanoreactors, catalysts and drug delivery systems⁹–¹¹. For example, Lee and coworkers fabricated Au@SiO₂ yolk-shell nanospheres through a two-step etching treatment with potassium cyanid¹². Liu and coworkers used dopamine as the C source and fabricated yolk-shell structured Au@C nanocomposites. Taking the advantages of C shell, the catalyst exhibited high catalytic activity, and the reduction of 4-nitrophenol (4-NP) could be finished in 5 min¹³. However, the size of metal NPs in these single-shelled structures are usually larger than 10 nm, which greatly attenuates their catalytic activity and limits their extensive application. Therefore, it is still a challenge to develop new types of yolk-shell nanostructures that can effectively prevent aggregation of NPs, retain their ultrafine particle size, and keep their high catalytic activity.

Herein, we report for the first time the design and synthesis of double-shelled reduced graphene oxide@palladium@carbon (RGO@Pd@C) hollow spheres with RGO as inner shell and amorphous C as outer shell. Graphene has been successfully utilized as a support to disperse and stabilize metal nanoparticles, because of its large surface area, extraordinary electronic transport property and strong mechanical strength¹⁴,¹⁵. Ultrafine Pd NPs grow well on the surface of inner shell by spontaneous redox reaction between graphene oxide (GO) and PdCl₄²⁻. Moreover, the outer shell combined with inner shell can confine Pd NPs in it, which effectively prevent the aggregation and leaching of Pd NPs and increase their catalytic active area, therefore, enhance their stability and catalytic performance. Representatively, the reduction of 4-NP catalyzed by RGO@Pd@C nanocomposites can be finished within only 30 s even the content of Pd is as low as 0.28 wt%, which outperforms the catalytic activity of other reported nanocomposites¹²,¹³.
step 4, the SiO2 cores were etched by HF solution and the double-ordered structure of graphene by thermally reduction. In the final precursor shell. During this process, GO layer was partially the 2D and heated at 500°C and the thickness of the RGO@Pd@C hollow spheres are quite uniform (Fig. 3B). The selected-area electron diffraction (SAED) pattern corresponds to the (111), (200), (220) and (311) planes of the expected fcc Pd (Fig. 3C). The X-ray photoelectron spectroscopy (XPS) of RGO@Pd@C exhibits two peaks at 335.5 eV and 340.5 eV, respectively (Fig. 3D), which are in good agreement with the reported XPS data of Pd 2p3/2 and Pd 2p1/2 in metallic Pd20. In addition, energy-dispersive X-ray (EDX) spectrum of composite materials also identifies the peak of Pd (Fig. S3). In comparison, RGO/Pd nanocomposites are fabricated under the same condition. But no hollow structures are formed due to the thin nanosheets of RGO can not preserve the structural integrity and spherical morphology after etching of the SiO2 template (Fig. S4), which indicates that the outer C shell avoid the collapse of the hollow spheres.

Inspired by the high catalytic activity of Pd NPs as well as the unique structure of the double-shelled hollow spheres, we explore the applications of RGO@Pd@C as a catalyst in the reduction reaction of 4-NP to 4-AP with the aid of NaBH4. It is well-known that this reaction is simple and fast in the presence of metallic surfaces21–24. While upon the addition of RGO@Pd@C catalysts, the reduction reaction can be finished within only 30 s. Fig. 4A shows the UV-vis spectrum changes of the reaction mixture in the presence of RGO@Pd@C catalysts. The absorption of 4-NP at 400 nm disappears quickly and absorption of 4-AP at about 300 nm peak increased accordingly. Meanwhile, the complete reduction of 4-NP to 4-AP can also be detected by color change of the solution, which changes from originally bright yellow to colorless (Fig. 4A insert). The reduction with RGO@Pd@C composite as catalyst can be done in only 30 s, even when the content of Pd in it is as low as 0.28 wt%, whereas reactions in the presence of Au@SiO213, Au@C15, Ag/
Carbon nanofiber\textsuperscript{25} and Au/Graphene\textsuperscript{26} catalysts are finished more than 5 min (Table 1). Furthermore, the turnover frequency (TOF), defined as moles of the reactant (4-NP) converted by per mole of active metal in catalyst per minute, also shows much higher for RGO@Pd@C than that of Ag/Carbon nanofiber and Au/Graphene catalysts.

**Discussion**

The excellent catalytic performance of RGO@Pd@C composite can be ascribed to synergistic effects between Pd catalyst and double-shelled RGO@C layers. Firstly, the double-shelled RGO@C layers can effectively inhibit the aggregation and leaching of Pd NPs, and render RGO@Pd@C with high catalytic stability due to the chemical inertness and excellent mechanical stability of RGO and C shells. On the other hand, RGO can absorb 4-NP via $\pi-\pi$ stacking interactions, and facilitate transportation of electron from RGO to Pd NPs, which to a great extent improve the catalytic performance of RGO@Pd@C nanocomposite. The stability of catalyst was investigated by repeated measurements in 4-NP under the same condition. After each measurement, the catalyst was recycled by simple

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**Figure 3** | (A) and (B) SEM images of RGO@Pd@C hollow spheres with different magnifications. (C) The selected-area electron diffraction (SAED) pattern and (D) XPS spectrum of Pd 3d of RGO@Pd@C.

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**Figure 4** | (A) UV/Vis spectra of 4-NP reduction reaction in the absence and presence of RGO@Pd@C catalyst. Insert: color changes of the conversion of 4-NP. (B) The stability of catalyst by performing the same reduction reaction with 10 cycles.
centrifugation, followed by washing with distilled water and drying in an oven overnight for the next cycle of catalysis. The catalysts exhibited well stability after 10 cycles of reactions, with 100% conversion within 40 s reaction periods. However, the reaction time increased from 25 s to 420 s under the same condition when RGO/Pd nanocomposite was used as the catalyst. (Fig. 4B). This proves that the double-shelled RGO@Pd@C hollow spheres are highly effective and stable catalysts with great potential applications.

In summary, we have developed tailor-made double-shelled hollow carbon spheres with RGO as inner shell and C layer as outer shell, which acts as an excellent capsule for encapsulating with ultrafine Pd NPs. This unique structure can effectively prevent the aggregation and leaching of Pd NPs as well as render them large active area, therefore, facilitate the high stability and catalytic performance of the nanocomposites. Using RGO@Pd@C as catalyst, the reduction of 4-NP to 4-AP could be finished in 30 s, which demonstrates that it is one of the most effective catalyst as far as we know. In addition, RGO@Pd@C also shows excellent tolerance to the chemical environment. We envision that the unique double-shelled hollow spheres will have broad applications in many fields including catalyst and drug delivery.

**Methods**

**Synthesis of GO wrapped SiO2(3SiO2-GO) nanospheres.** GO was prepared according to a modified Hummer’s method.

**Synthesis of Pd NPs loaded SiO2@GO (SiO2@GO@Pd) nanospheres.** 150 mg of as-synthesized SiO2@GO@Pd nanospheres were dispersed in 60 mL of DI water, then 3 mL of 20 mM H2PdCl4 aqueous solution was added and the mixture was stirred for 8 h at 60 °C under inert atmosphere.

**Synthesis of Pd NPs encapsulated double-shelled hollow carbon (RGO@Pd@C) nanospheres.** 150 mg of as-prepared SiO2@GO@Pd nanospheres were dispersed in 16 mL of water/ethanol (volume ratio = 3/1) mixture by ultrasonication, then 4 mL of 0.5 M aqueous glucose solution was added with vigorous stirring for 30 min. After that, the suspension was transferred to a 25 mL Teflon-lined autoclave, and heated in an oven at 180 °C for 16 h. The dark gray products were collected by centrifugation and washed with ethanol and DI water for six times, respectively. After drying at 60 °C overnight, the resulting dark gray powder was characterized by TG-DTA, and the weight loss was less than 5%.

**Materials characterization.** The morphology and structure of products were characterized with a field-emission scanning electron microscope (SEM, FEI, Nova NanoSEM 450) and a transmission electron microscope (TEM, FEI, Tecnai G2 20), X-ray photoelectron spectroscopy (XPS) measurements were performed on a VG ESCALAB 250 spectrometer with monochromatic Al Kα (1486.71 eV) X-ray radiation (15 kV and 10 mA) and hemispherical electron energy analyzer. The UV-vis measurements were performed on a UV-2550 spectrophotometer (Shimadzu, Japan). The Pd contents in the catalysts were determined using Microwave Plasma-Atom Emission Spectrometer (MP-AES, Agilent 4100, USA).

**Catalytic study.** 5 mg of RGO@Pd@C nanocomposites was added to 3 mL of 1 × 10−4 M 4-NP solution. 0.1 mL of 3 × 10−4 M NaBH4 solution was then added with constant magnetic stirring. The changes of the reduction reaction were recorded in the UV-vis spectrophotometer.

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| Catalyst | Mass of catalyst (mg) | Amount of 4-NP (mmol) | Amount of NaBH4 (mmol) | Metal size (nm) | Metal content (wt%) | Conversion time (min) | TOF (mmol 4-NP/(mmol metal min)) | Reference |
|----------|----------------------|-----------------------|------------------------|----------------|---------------------|----------------------|-------------------------------|----------|
| RGO@Pd@C | 5                    | 3 × 10−4              | 3 × 10−2               | 4              | 0.28                | 0.5                  | 4.56                          | Our work |
| Au@C     | 5                    | 3 × 10−4              | 3 × 10−2               | 15             | NA                  | NA                   | NA                           | 13       |
| Ag/Carbon nanofiber | 1 | 3.6 × 10−3          | 15 × 10−2            | 28.1          | 8.4                   | 8                    | 0.58                          | 25       |
| Au@Graphene | 0.1      | 2.8 × 10−4          | 2 × 10−2              | 14.6          | 24                   | 12                   | 0.19                          | 26       |
| Au@SiO2  | NA                   | 3.4 × 10−3           | 1.2                   | 103            | NA                   | 60                   | NA                           | 12       |

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Author contributions
Z.Z. and F.X. contributed equally. Z.Z., F.X., H.W., S.W. and Y.L. proposed, planned, and designed the project. Z.Z., J.X., T.S. and S.X. performed the material preparation, characterizations, and catalytic tests. All authors contributed to writing the manuscript.

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