Surfactant-Mediated One-Pot Method To Prepare Pd–CeO$_2$ Colloidal Assembled Spheres and Their Enhanced Catalytic Performance for CO Oxidation

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Supporting Information

ABSTRACT: A simple, one-pot method to fabricate ordered, monodispersed Pd–CeO$_2$ colloidal assembled spheres (CASs) was developed using the surfactant-mediated solvothermal approach, which involves a tunable self-assembled process by carefully controlling different chemical reactions. The evolution process and formation mechanism of the CASs were thoroughly investigated by time-controlled and component-controlled experiments. For CO oxidation, this CAS nanocatalyst exhibited much higher catalytic activity and thermal stability than Pd/CeO$_2$ prepared by an impregnation method, and its complete CO conversion temperature is $\sim$120 °C. The enhanced catalytic performance for CO oxidation could be attributed to the synergistic effect of highly dispersed PdO species and Pd$^{2+}$ ions incorporated into the CeO$_2$ lattice. For this CAS catalyst, each sphere can be viewed as a single reactor, and its catalytic performance can be further improved after being supported on alumina, which is obviously higher than results previously reported. Furthermore, this method was used to successfully prepare M–CeO$_2$ CASs (M = Pt, Cu, Mn, Co), showing further that this is a new and ideal approach for fabricating active and stable ceria-based materials.

1. INTRODUCTION

Pd-based catalysts have been intensively researched in recent years because of their wide applications in various catalytic processes.$^{3-4}$ For example, Pd-catalyzed carbon–carbon cross-coupling reactions are the most important and practical reactions in organic chemistry.$^{5,6}$ Supported PdO$_x$ species have been recognized as the active sites for CH$_4$ combustion, and an application of Pd-only three-way catalysts represents a meaningful progress for the removal of pollutants from exhaust gas.$^{7,8}$

It is known that the catalytic activity of Pd nanoparticles (NPs) is closely related to their size, shape, and valence state;$^{9-12}$ thus it is of great importance to maintain the morphology and physicochemical states of Pd-active species under various reaction conditions. For this purpose, Pd NPs are often loaded on supports, such as metal oxides, zeolites, carbon materials, and so forth. Among these kinds of supports, cerium oxide is considered as a unique one because of its oxygen storage/release capacity and its redox capability to influence the chemical and electronic states of Pd species.$^{13}$

Pd/CeO$_2$ has been widely researched and used in various catalytic reactions, such as three-way catalytic exhaust gas purification, volatile organic compound (VOC) elimination, selective hydrogenations, and photocatalytic reactions.$^{14-19}$ In recent years, with the help of advanced nanomaterial synthesis technology, it is now possible to purposefully design Pd–CeO$_2$ hybrid nanocatalysts with unique features and further improve their functionality.$^{20-24}$ For instance, Cargnello et al. designed and prepared a Pd@CeO$_2$-type catalyst by means of the concepts of supramolecular chemistry and showed an exceptional activity and stability in methane combustion.$^{25-27}$ Zhang et al. explored a clean synthesis of high-quality Pd@CeO$_2$ nanospheres by salting-out-assisted growth and self-assembly, which improved their catalytic activity for CO oxidation.$^{28}$ They recently also fabricated a Pd@CeO$_2$ hybrid structure with tunable Pd core sizes with a simple two-step biomolecular-assisted process without any expensive or toxic ligand.$^{29}$ We have synthesized a hollow-structured core–shell-like Pd–CeO$_2$ nanocomposite by the hard-template method, which exhibited enhanced catalytic activities for CO oxidation and selective reduction of 4-nitrophenol. The Pd particles can be well protected by the CeO$_2$ shell, and the catalytic stability could be well maintained even after calcination at 700 °C.$^{30}$

Although many gratifying achievements have been made, there are still some challenges remaining. In Cargnello’s protocol, multiple procedures are needed to obtain the hybrid structures, which is time consuming, inevitably. Besides, large amounts of toxic organic solvents and surfactants are required during the assembly processes. In Zhang’s process, the size of Pd NPs is

Supporting Information

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relatively large (>8 nm); thus, the utilization of the Pd element is not efficient enough. Therefore, it is still necessary to develop simple and green methods for preparing highly active Pd−CeO₂ hybrid nanocatalysts with more efficient Pd utilization.

With the help of the hands-on experience of researchers, the key factors for successfully synthesizing novel Pd−CeO₂ hybrid structures include the manipulation of individual chemical reactions rates, especially tuning interfacial chemical processes among different components. In this paper, we developed a one-pot, surfactant-assisted solvothermal method to prepare Pd−CeO₂ colloidal assembled spheres (CASₜ). The Pd clusters were dispersively distributed in CeO₂ CASₜ by a one-pot method, which is efficient enough than some of the two-step processes to prepare noble-metal/oxide nanospheres.31,32 The results showed that the Pd−CeO₂ CAS superstructure exhibited an enhanced catalytic activity for CO oxidation compared with a conventional Pd/CeO₂-supported catalyst. Moreover, this method can be easily extended to the synthesis of Pt−CeO₂ CASₜ and transition metal-incorporated CeO₂ CASₜs.

2. EXPERIMENTAL SECTION

2.1. Sample Synthesis. Pd−CeO₂ CASₜ. CeCl₃·7H₂O was dissolved in ethylene glycol (6 g CeCl₃·7H₂O per 100 mL of solvent) under stirring for 2 h. CeCl₃·7H₂O ethylene glycol solution (10 mL) and 30 mL ethylene glycol were added into a 250 mL flask under stirring at room temperature. Then a certain amount of K₂PdCl₄ was added under intensive stirring for 30 min, followed by adding 0.1 g of polyvinylpyrrolidone (PVP), 2 mL of acetic acid, 0.5 mL of tetramethyammonium hydroxide (N(CH₃)₄OH, 25% in H₂O), forming a transparent light-brown solution. This solution was immersed in an oil bath, and the temperature was increased up to 130 °C; a grayish suspension solution was obtained after 15 h of the reaction.

After the reaction was completed and cooled down to room temperature, this mixture was diluted with 40 mL of acetone and then centrifuged at 4500 rpm for 15 min. The obtained solid was dispersed in the mixed solution of acetone and ethanol (v/v = 1:1), followed by centrifugation twice at 4000 rpm for 10 min. The obtained product was dried at 60 °C overnight and calcined at 600 °C for 2 h.

Pd−CeO₂ CASₜ Supported on γ-Al₂O₃. A certain amount of γ-Al₂O₃ was dispersed in 50 mL of deionized water under ultrasonication for 10 min. A calcined Pd−CeO₂ CASₜ suspension (20 mL, 2.5 mg/mL) was slowly added into the Al₂O₃ suspension suspension under stirring at room temperature for 1 h. The product was gathered by centrifugation and dried at 80 °C overnight.

Pd/CeO₂-Imp Catalyst. The conventional Pd/CeO₂ catalyst was prepared by the impregnation method with PdCl₂ as the precursor. The collected powder was dried at 80 °C and calcined at 600 °C for 2 h.

Pt−CeO₂ CASₜ. K₂PtCl₄ was used as a precursor instead of K₂PdCl₄. The preparation reaction was conducted at 140 °C for 20 h, and other processes were the same as those of Pd−CeO₂ CASₜs.

Transition Metal (M = Cu, Co, Mn)−CeO₂ CASₜ (M−CeO₂ CASₜ). CuCl₂, CoCl₂, and Mn(CH₃COO)₂ were used as precursors with a molar ratio of 1:9 (transition metal: Ce), and the synthesis processes were the same as those of Pd−CeO₂ CASₜs except the addition of PVP. For the Mn−CeO₂ CAS sample, the reaction time was prolonged to 24 h.

2.2. Catalytic Activity Testing. The catalytic performances of samples for CO oxidation were evaluated in a fixed-bed reactor. For a typical testing, the Pd−CeO₂ catalyst diluted with silica was loaded in a quartz-tube reactor. Then, the feed gas containing 1 vol % CO, 20 vol % O₂, and balance N₂ was introduced into the reactor at a flow rate of 50 mL/min. After the sample was pretreated in 10 vol % O₂/N₂ at 150 °C for 30 min, the temperature of the reactor was raised at a rate of 2 °C/min from room temperature. The gas composition was monitored online using a GC (GC 2060 system) with a flame ionization detector (FID) and methane converter. The conversion of CO was calculated on the basis of the change of CO concentrations in the inlet and outlet gases.

Kinetic experiments were performed at a gas hourly space velocity (GHSV) of 100 000 or 150 000 mL/(gcat·h). Typical conversions of CO were kept well below 10% so that the differential conditions could be assumed.

The stability testing was performed under a cycling experiment. Before the testing, the catalyst was pretreated in the feed gas at 600 °C for 30 min and then cooled down to 85 °C to test the conversion of CO to CO₂. The cycle was repeated three times.

Turnover frequency (TOF) measurements were conducted at 60 °C at a GHSV of 100 000 mL/(gcat·h). The CO conversion was obtained after 20 min of the CO reaction, and the TOF was calculated based on the equation TOF = \( \frac{V_{\text{CO}}A}{N} \), where \( V_{\text{CO}} \) is the CO conversion, \( V_{\text{CO}} \) is the gas flow of CO (mol/s), \( N \) is the Avogadro constant, and \( N \) is the number of the catalytic sites (\( N = N_tD_{\text{pd}} \), where \( N_t \) is the total amount of Pd atom in the catalyst and \( D_{\text{pd}} \) is the Pd dispersion).

2.3. Characterization of Samples. Elemental analysis of the sample was done by inductively coupled-plasma atomic emission spectroscopy (ICPAES) on a Varian 710ES instrument (Varian Co. Palo Alto, US). Powder X-ray diffraction (XRD) patterns were recorded on a PANalytical PW 3040/60 X Pert Pro powder diffractometer with Cu Kα radiation, which was operated at 40 kV and 40 mA and a scanning speed of 0.5°/min. N₂ adsorption/desorption isotherms of samples were measured at −196 °C on a micromeritics ASAP 2020 instrument, and the surface area was calculated by the Brunauer−Emmett−Teller (BET) method. The Pd dispersion in the catalysts was calculated by the CO chemisorption method on AutoChem II 2920 chemical adsorption equipment at room temperature.

H₂ temperature-programmed reduction (H₂-TPR) was carried on a PX200 apparatus (Tianjin Pengxiang Technology Co. Ltd.) with a thermal conductivity detector (TCD). The catalyst (40 mg) was used, and the reduction gas consisted of 5% H₂/N₂ with a flow rate of 40 mL/min. TPR was run from room temperature to 800 °C at 10 °C/min.

Transmission electron microscopy (TEM) images were taken on a JEM-2100 transmission electron microscope. The sample was ultrasonically suspended in the ethanol solvent, and one or two drops of this slurry were deposited on a copper grid. The liquid phase was evaporated before the grid was loaded into the microscope.

The X-ray photoelectron spectroscopy (XPS) spectra of samples were obtained on a Kratos Axis Ultra-DLD photoelectron spectrometer equipped with Al Kα (1486.6 eV) radiation as the excitation source. All binding energies (BEs) were determined with respect to the C1s line (284.8 eV) originating from adventitious carbon.

The size distribution of spheres was measured by photon correlation spectroscopy on a Zetasizer instrument (Malvern Instruments Ltd.) at room temperature. The suspension was
prepared with a concentration of 0.5 mg/mL followed by ultrasonication for 15 min.

3. RESULTS AND DISCUSSION

The Pd–CeO2 CASs were synthesized in a simple one-pot fashion by the modified solvothermal method. As shown in TEM images (Figure 1a,b) and size distribution profiles (Figure S1), the as-synthesized Pd–CeO2 CASs are well monodispersed, and their average size is 100–120 nm. The as-synthesized hybrid spheres are composed mainly of small 2–3 nm CeO2 nanocrystals, which is consistent with the XRD results (Figure 1g). However, the dense stacking of CeO2 nanocrystals makes it difficult to discern the distribution of Pd species in the spheres, so high-angle annular dark-field scanning transmission electron microscopy (HAADF-STEM) was used to illustrate the mappings of Pd, Ce, and O components in the CAS. As shown in Figure 1f, Pd is randomly scattered in the entire colloidal sphere, which is different from some results reported. After calcination at 600 °C to completely remove organic residues, the morphology of spheres remains unchanged and the size of the spheres obviously shrinks, and their most probable diameter decreases from 110 to 80 nm (Figure S1). Also, the CeO2 size

Figure 1. TEM images of (a,b) the as-prepared Pd–CeO2 CASs, (c–e) sample calcined at 600 °C, (f) STEM image and elemental mapping of Pd–CeO2 CASs, and (g) XRD profiles of the as-prepared and calcined Pd–CeO2 CASs.

Figure 2. (A) Pd 3d and (B) Ce 3d XPS spectra of the as-synthesized and calcined Pd–CeO2 CASs at 600 °C.
increases a bit (Figure 1c), and its XRD peaks strengthened to some extent (Figure 1g). In the HRTEM micrograph of calcined Pd−CeO₂ CASs (Figure 1d), the lattice fringes with an interplanar spacing of 0.31 nm is associated with the (111) facet of CeO₂ NPs, and the obvious lattice discontinuities can be observed, which characterizes further the nanosized CeO₂ and the possible lattice strain. Figure 1e is the magnified picture of a single sphere and well demonstrates that ultrasmall Pd clusters (indicated by white arrows) are embedded in the assembled CeO₂ NPs, resulting in a strong interaction between Pd clusters and CeO₂ NPs.

To further confirm the close interaction between Pd and CeO₂ components, XPS was applied to investigate the valence state of Ce and Pd. As shown in Figure 2A, the peaks at BEs of 336.8 and 337.9 eV can be attributed to the dispersed PdO clusters and the Pd−O−Ce structure, respectively, which clearly demonstrated that the Pd species in the Pd−CeO₂ CASs interacted closely with CeO₂ nanocrystals. The Ce 3d XPS spectra in Figure 2B showed a valence variation before and after calcination. The as-prepared sample possesses more Ce³⁺ species that is caused by ethylene glycol during the synthesis, and after thermal treatment, the oxidation of Ce³⁺ to Ce⁴⁺ obviously occurred, indicating a thermal-induced crystallization process.

The N₂ adsorption/desorption isotherms of the as-synthesized Pd−CeO₂ CASs before and after calcination at 600 °C are shown in Figure 3. In the low relative pressure region, the curves reflect a microporous structure, which may be caused by the stacking of nanocrystals that assembled the spheres. A hysteresis, typical of type IV, appeared in the high relative pressure region, indicating a relatively large pore structure, which is likely caused by the stacking of spheres. The surface area of the sample before and after being calcined at 600 °C is 89 and 80 m²/g, respectively, which is larger than that of the core−shell Pt/Pd@CeO₂ spheres reported.35

The formation mechanism of Pd−CeO₂ CASs can be discussed in detail based on the time-controlled experiments. Followed by the visual variation of the reaction system, the formation process undergoes three stages, that is, the nucleation of Pd clusters, formation of CeO₂ nanocrystals, and self-assembly of these two components.

First, the brown transparent solution gradually turned black after 20−30 min of the reaction when the temperature ramped up to 130 °C, indicating the possible formation of Pd clusters in the solution. To verify this assumption, we characterized the solid sample formed for 1.5 h. As shown in Figure 4, the sheetlike structure and ultrasmall Pd clusters coexisted in the sample, but no lattice fringes can be observed. Also, the elemental analysis shows that this sample is consisted of C, H, O, N, Ce, and Pd elements (Table S1). The XRD results (Figure 4f) show that it is a Ce−glycate-like inorganic−organic network structure and was caused by the Ce ions interacted with acetic acid and ethylene glycol at 130 °C. The FT-IR spectrum (Figure S2) also confirmed that the Ce ions have bonded with ethylene glycol and acetic acid on the basis of the typical IR absorption peaks at 1562 cm⁻¹ ν(OCO), 2942, and 2895 cm⁻¹ ν(C−H) C₂H₄.34

With the increase in the synthesis time, the hydrolysis of Ce species was gradually promoted, and the subsequent self-assembly of nucleated CeO₂ nanocrystals formed the primary spheres (Figure 4b−d). After 10 h of the synthesis reaction, the black transparent solution gradually became blurred, indicating the initial formation of the CeO₂ structure. As shown in Figure 4c,d, the primary CAS and CeO₂ nuclei can coexist simultaneously. Finally, the Pd clusters and CeO₂ nanocrystals coassembled into the Pd−CeO₂ hybrid structure (Figure 4e).

The XRD patterns in Figure 4f can well reflect the structure change of different intermediate compounds formed in the fabrication process of Pd−CeO₂ CASs.

To obtain a highly dispersible Pd−CeO₂ hybrid structure in a one-pot fashion, the key is to tune the reaction rates for different chemical processes; that is, the balance between the slow hydrolysis of the Ce precursor, Pd-cluster formation, and self-assembled process should be critically controlled, which can be well achieved with the help of surfactants and cosolvents. To clarify the effect of each reagent on the synthetic results, we also designed a series of component-controlled experiments.

(1) We tested the essential role of ammonia hydroxide, acetic acid, and N(CH₃)₄OH in the synthesis of Pd−CeO₂ CASs. When metal salt precursors were added in the solution including acetic acid and N(CH₃)₄OH but without ammonia hydroxide, the obtained synthesis solution was transparent black after 15 h, and only the reduction of Pd²⁺ to Pd⁰ occurred and no assembled CeO₂ nanocrystal spheres formed. This was verified by the typical Pd peaks in the XRD pattern of the obtained product (Figure S3).

(2) When the amount of NH₄H₂O was increased 2-fold up to 0.8 mL, the solution immediately became opaque at 70−80 °C, indicating that a large amount of CeO₂ crystals formed very quickly and its nucleation could not be controlled. The obtained spheres would severely conjugate one another, and the bulklike structure formation was inevitable (TEM image shown in Figure S4). These phenomena confirmed the inevitable role of an appropriate amount of NH₄H₂O, which functions as a pH regulator and affects the hydrolysis rate of Ce ions.

(3) Although NH₄H₂O is indispensable in the successful fabrication of Pd−CeO₂ CASs, its role can be played only in the presence of acetic acid. Without the addition of this organic acid, the produced Pd particles would aggregate, and CeO₂ nanocrystals could not be well self-assembled into spheres (TEM image shown in Figure S5). It was reported recently that appropriate amounts of formic acid and ammonia hydroxide could be used as a formic acid/ammonium formate buffer solution to effectively control the OH⁻ concentration.35 We consider that the acetic
acid/ammonium acetate plays the similar role in the fabrication of Pd–CeO₂ CASs by tuning the reaction process.

(4) N(CH₃)₄OH was commonly used as a pH regulator as well as a dispersing agent in the synthetic process, which was reported in the synthesis of aqueous dispersible CeO₂ NPs. PVP helps to stabilize Pd clusters against aggregation. If PVP was not added, the black precipitate would be formed and stuck at the flask bottom when the synthesis reaction occurred at 130 °C for a short time.

On the basis of the above results, each reagent is necessary and its dosage needs to be carefully controlled or balanced to obtain the dispersible Pd–CeO₂ CASs. To our knowledge, it is difficult to design and prepare noble metal- or transition metal-incorporated CeO₂ hybrid nanospheres by a universal method. However, we luckily found that the present approach of structural design could be well used to synthesize other CAS architectures. For example, platinum and some common transition metals (Cu, Mn, and Co) can be easily introduced into CeO₂ CASs by slight changes of experimental conditions. As shown in Figure 5, the diameter of the as-prepared Pt–CeO₂ CASs is nearly the same as that of Pd–CeO₂ CASs, and they are mainly composed of ultrasmall CeO₂ nanocrystallites, and no Pt NPs can be observed. The XPS analysis indicates that the Pt species existed in an ionic state (Figure S6), confirming the highly dispersed Pt species in Pt–CeO₂ CASs. For transition metal-incorporated M–CeO₂ CASs (M = Cu, Co, Mn), their sphere diameters are less than 50 nm, and their XRD profiles showed only the characteristic diffraction peaks of the fluorite phase.
CeO₂ structure (Figure S7), which indicates that the successful incorporation into the CASs. Their transition metal contents were measured by the EDS analysis (Figure S8), and Cu/Ce (at %) = 9.5, Co/Ce = 4.2, and Mn/Ce = 6.5.

To investigate the catalytic performance of the Pd−CeO₂ CAS material, the CO catalytic oxidation was used as the model reaction. For comparison, we tested the catalytic activities of the Pd/CeO₂-imp sample prepared by the impregnation method and pure CeO₂ for CO oxidation. As shown in Figure 6A, Pd−CeO₂ CASs exhibited the best catalytic activity, and the temperature (T₁₀₀) of 100% CO conversion is around 125 °C, which is much lower than that of the Pd/CeO₂-imp catalyst (190 °C). T₁₀₀, T₅₀, and T₉₀ (the reaction temperatures of 10, 50, and 90% CO conversions) were shown in Table 1. Besides, the Pd−CeO₂ CASs also showed a good thermal stability after the cycling testing 3 times at 600 and 85 °C (Figure 6B). A decline of 2−3% CO conversion can be observed, which can be explained by the minor accumulation of carbonate species on the catalyst surface. The TEM image (Figure S9) showed that the morphology of Pd−CeO₂ CASs was well maintained and no agglomeration occurred after the cycling testing 3 times. A recent study on similar the Pd@CeO₂ nanostructure showed that at medium-high temperature and an oxidizing atmosphere cerium could be quite volatile and some new structural features would appear, including highly dispersed SiO₂−CeOₓ species and rather bigger CeO₂ particles. Therefore, after the relatively long time utilization of Pd−CeO₂ CASs in the cycling experiment at 600 °C, we cannot neglect their possible structural change, although the morphology of Pd−CeO₂ CASs could be well maintained. Also, further detailed characterization and analysis are needed.

The kinetic data of CO oxidation on Pd−CeO₂ CASs and Pd/CeO₂-imp are shown in Figure 6C. The results showed that the Pd−CeO₂ CASs exhibited much higher catalytic activity (or much higher reaction rate) than the Pd/CeO₂-imp catalyst. Also, the apparent activation energy (Eₐ) of the Pd−CeO₂ CAS catalyst was 44 kJ/mol, which is much lower than that (Eₐ = 75 kJ/mol) of the Pd/CeO₂-imp catalyst. We have further calculated the TOF at 60 °C; the TOF of the Pd/CeO₂-imp catalyst is ~0.026 s⁻¹, and the TOF of Pd−CeO₂ CASs is ~0.051 s⁻¹ (Table 1), which also confirmed that the Pd−CeO₂ CAS nanocatalyst possessed higher catalytic activity than the Pd/CeO₂-imp catalyst for CO oxidation under the same reaction condition. Furthermore, we listed TOFs of reported Pd−CeO₂

**Table 1. Pd Contents, Pd Dispersions (Dₚd), and the Catalytic Activities of Pd−CeO₂ CASs and Pd/CeO₂-imp for CO Oxidation**

| sample          | Pd content (wt %) ¹   | Dₚd (%) ²   | T₁₀₀ (°C) | T₅₀ (°C) | T₉₀ (°C) | TOF (s⁻¹) |
|-----------------|----------------------|-------------|-----------|----------|----------|-----------|
| Pd−CeO₂ CASs    | 1.1                  | 28          | 24        | 74       | 113      | 0.051     |
| Pd/CeO₂-imp     | 0.97                 | 34          | 48        | 115      | 150      | 0.026     |

¹Pd content was measured by ICP. ²Pd dispersion was determined by CO chemisorption. ³TOFs were measured at 60 °C.
catalysts at certain temperatures for CO oxidation in Table 2, and the results show that the catalytic activity of Pd–CeO2 CASs is higher than that reported.

Table 2. Catalytic Activities of Pd–CeO2 Catalysts for CO Oxidation

| Catalyst               | Reaction Temperature (°C) | TOF (s⁻¹) | Ref. |
|------------------------|---------------------------|-----------|------|
| Pd/CeO2-nanorod        | 50                        | 0.04      | 38   |
| Pd/CeO2                | 97                        | 0.017     | 41   |
| Pd–mesoporous CeO2     | 20                        | 0.024     | 42   |
| PdO–CeO2/SiO2          | 150                       | 0.038     | 43   |
| PdO/Fe–CeO2 CASs       | 140                       | 0.036     | 44   |
| Pd/Al2O3 model catalyst | 80                        | ~0.04     | 45   |
| Pd–CeO2 CASs           | 60                        | 0.051     | this work |

To investigate and compare the differences of reducibility and an interaction of Pd species with CeO2 between the Pd–CeO2 CASs and Pd–CeO2-imp catalyst, H2-TPR was used to test these two catalysts, and the results are shown in Figure 7. In the two TPR profiles, there are two peaks at below 200 °C, but their intensities and positions are different. We may attribute the first peak at below 200 °C to the reduction of PdO species and the second to the reduction of the Pd2+-O–Ce4+ linkage structure.37,40 The calculated H2 consumption of the first peak in Pd/CeO2-imp at 60 °C is ~104 μmol/g and is close to the theoretical value (88 μmol/g) for PdO reduction to metallic Pd. For Pd–CeO2 CASs, the H2 consumption of the first peak is much lower than its theoretical value. This is because of the fact that in the Pd–CeO2-imp sample the Pd species existed mostly in a form of PdO by the impregnation process with few Pd2+ incorporated into the CeO2 support; in the Pd–CeO2 CAS sample, most Pd species formed a solid solution with CeO2 nanocrystallites produced during the solvothermal process. Moreover, we can observe a very small negative peak in the TPR curves of two samples (~60 °C for Pd–CeO2 CASs and ~80 °C for Pd/CoO2-imp), which could be attributed to the decomposition of PdHx species.

In the reduction peaks at <200 °C, the total H2 consumption of Pd/CoO2-imp and Pd–CeO2 CASs is ~211 and 427 μmol/g, respectively, which are higher than that when reducing PdO to Pd. Thus, it is inevitable that the reduction peak may contain simultaneous reduction of Ce4+ ions near Pd species. This is because that the presence of noble metals can promote the reduction of metal oxides by the H2 spillover from the metal to the oxide support, as well as the increase of oxygen vacancies on the ceria support. With more Pd2+ ions incorporated into the CeO2 nanocrystals in the synthesis process, the Pd–CeO2 CASs possessed more oxygen vacancy defects; thus, more Ce4+ can be easily reduced at lower temperatures.

For the reduction peak at 320–370 °C, it should be attributed to the reduction of surface lattice oxygen. In the Pd–CeO2 CAS catalyst, because the Pd2+-O–Ce4+ linkage structure (Pd2+ incorporated into CeO2) existed besides highly dispersed PdO clusters, part of surface lattice oxygen was reduced at <200 °C, which was promoted by the reduction of Pd2+ ions, resulting in the decrease in the peak area at 324 °C. The reduction peak at >700 °C is ascribed to the reduction of bulk CeO2 and affected by the CeO2 particle size. The average particle size of CeO2 is 5–6 nm for Pd–CeO2 CASs, and for the Pd/CoO2-imp catalyst, the particle size of its commercial nanosized CeO2 powder is over 20 nm. This leads to a distinguishing difference in their H2-TPR profiles. With the decrease of the CeO2 particle size, more surface oxygen existed in the Pd–CeO2 CASs, so the reduction of bulk CeO2 decreased obviously.46

As shown in the characterization results above for the Pd–CeO2 CAS catalyst, the active Pd species existed mainly in highly dispersed PdO clusters and Pd2+ incorporated into CeO2 (Pd2+-O–Ce4+ linkage structure), which act the active sites for CO oxidation. In this Pd–CeO2 CAS hybrid spheres, the enhanced catalytic activity can be attributed to the synergetic effect provided by two aspects: the surface dispersed PdO clusters are more active for CO adsorption, and the Pd2+ ions incorporated into the CeO2 lattice creates more oxygen vacancies, which are beneficial to oxygen activation in CO oxidation. For the Pd/CoO2-imp catalyst, the O2 activation might be weaker than that for the Pd–CeO2 CAS catalyst because of the lack of oxygen vacancies on the surface, resulting in its lower catalytic activity for CO oxidation.

One of the advantages of ordered hybrid nanocatalysts is that every independent unit can be viewed as an individual active reactor in the catalytic reaction, and it is helpful to load these nanocatalysts on high-surface area supports to further improve the overall catalytic performance.26,29 Hence, we prepared the Pd–CeO2 CAS/Al2O3 by a simple process (seen in the Experimental Section). The results showed that Pd–CeO2 CAS particles could be uniformly supported on the Al2O3 carrier (TEM image, Figure S10). The BET surface area (144 m²/g) of Pd–CeO2 CAS/Al2O3 is higher than that (80 m²/g) of the unsupported one, and their N2 adsorption/desorption isotherms are shown in Figure S11. After Pd–CeO2 CASs were supported, their dispersion was improved, resulting in an increase in their catalytic performance (Figure 6D). Besides, this procedure was also applied to fabricate the Pt–CeO2 CAS/Al2O3 sample; as shown in Figure S12, CO was completely converted to CO2 over this Pt catalyst at 115 °C, which has a higher catalytic activity than those reported for the Pt catalysts for CO oxidation.3,47,48

4. CONCLUSIONS

In summary, we have developed a facile one-pot method to successfully synthesize Pd–CeO2 CASs, in which ultrasmall Pd clusters are embedded in the CeO2 colloidal assembly. The synthesis process involves careful control of Pd-cluster formation, the Ce4+ hydrolysis rate, and the subsequent assembly process. Each surfactant/cosolvent is necessary for the successful
fabrication of this material. For CO oxidation, this Pd−CeO₂ CAS sample shows a much higher catalytic activity compared with the Pd/CeO₂ catalyst prepared by the impregnation method. The enhanced catalytic activity can be attributed to the maximized Pd−CeO₂ interface of this ordered hybrid spheres. Moreover, after the Pd−CeO₂ CAS sample was simply supported on alumina, its overall catalytic performance can be further improved, for instance, the temperature of the CO complete conversion was only around 90 °C. Because this Pd−CeO₂ CAS also exhibit a relatively high thermal stability, it can potentially be valuable for other hydrocarbon abatement processes, such as methane or propane combustion. The successful synthesis of M−CeO₂ CASs (M = Pt, Cu, Mn, Co) shows further that this method is a new and ideal approach for fabricating active and stable ceria-based materials.

**ASSOCIATED CONTENT**

**Supporting Information**

The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/acsomega.6b00050.

Size distributions of Pd−CeO₂ CASs, FT-IR spectrum and composition of the sample prepared for 1.5 h, XRD profile and TEM images of the sample prepared at 130 °C for 15 h with different additives, XPS spectrum of Pt−CeO₂ CASs, XRD profiles and EDS spectra of M−CeO₂ CASs, TEM images of cycled Pd−CeO₂ CASs and supported Pd−CeO₂ CAS/Al₂O₃, N₂ adsorption/desorption isotherms of Al₂O₃ and Pd−CeO₂ CAS/Al₂O₃, and activity profiles of Pt−CeO₂ CASs and Pt−CeO₂ CAS/Al₂O₃ (PDF)

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**Notes**

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**REFERENCES**

(1) Buchwald, S. L. Cross Coupling. *Acc. Chem. Res.* 2008, 41, 1439.
(2) García-Melchor, M.; Braga, A. A. C.; Lledós, A.; Ujaque, G.; Maseras, F. Computational Perspective on Pd-catalyzed C−C Cross-Coupling Reaction Mechanisms. *Acc. Chem. Res.* 2013, 46, 2626−2634.
(3) Peng, H.-C.; Xie, S.; Park, J.; Xia, X.; Xia, Y. Quantitative Analysis of the Coverage Density of Br⁻ Ions on Pd[100] Facets and Its Role in Controlling the Shape of Pd Nanocrystals. *J. Am. Chem. Soc.* 2013, 135, 3780−3783.
(4) Choi, S.; Shao, M.; Lu, N.; Ruditskij, A.; Peng, H.-C.; Park, J.; Guerrero, S.; Wang, J.; Kim, M. J.; Xia, Y. Synthesis and Characterization of Pd@Pt−Ni Core−Shell Octahedra with High Activity Towards Oxygen Reduction. *ACS Nano* 2014, 8, 10363−10371.
(5) Miyaura, N.; Suzuki, A. Palladium-Catalyzed Cross-Coupling Reactions of Organoboron Compounds. *Chem. Rev.* 1995, 95, 2457−2483.
(6) Xue, L.; Lin, Z. Theoretical Aspects of Palladium-Catalysed Carbon−Carbon Cross-Coupling Reactions. *Chem. Soc. Rev.* 2010, 39, 1692−1705.
(7) Cargnello, M.; Jaén, J. D.; Garrido, J. C. H.; Baklmutsy, K.; Montini, T.; Gaméz, J. J. C.; Gorte, R. J.; Fornasiero, P. Exceptional Activity for Methane Combustion over Moduler Pd@CeO₂ Subunits on Functionalized Al₂O₃. *Science* 2012, 337, 713−717.
(8) Wang, J.; Chen, H.; Hu, Z.; Yao, M.; Li, Y. A Review on the Pd-Based Three-Way Catalysts. *Catal. Rev. 2015*, 57, 79−144.
(9) Chen, X.; Wu, G.; Chen, J.; Chen, X.; Xia, Z.; Wang, X. Synthesis of “Clean” and Well-Dispersive Pd Nanoparticles with Excellent Electrocatalytic Property on Graphene Oxide. *J. Am. Chem. Soc.* 2011, 133, 3693−3695.
(10) Cheong, S.; Watt, J. D.; Tilley, R. D. Shape Control of Platinum and Palladium Nanoparticles for Catalysis. *Nanoscale* 2010, 2, 2045−2053.
(11) Martins, J.; Batali, N.; Silva, S.; Rafik-Clement, S.; Karelovic, A.; Debecker, D. P.; Chaumonnot, A.; Uzio, D. CO Hydrogenation with Shape-Controlled Pd Nanoparticles Embedded in Mesoporous Silica: Elucidating Stability and Selectivity Issues. *Catal. Commun.* 2015, 58, 11−15.
(12) Wang, Y.; Choi, S.; Zhao, X.; Xie, S.; Peng, H.-C.; Chi, M.; Huang, C. Z.; Xia, Y. Polyol Synthesis of Ultrathin Pd Nanowires via Attachment-Based Growth and Their Enhanced Activity towards Formic Acid Oxidation. *Adv. Funct. Mater.* 2014, 24, 131.
(13) Montini, T.; Melchionna, M.; Monai, M.; Fornasiero, P. Fundamental and Catalytic Applications of CeO₂-Based Materials. *Chem. Rev.* 2016, 116, 5987−6041.
(14) Zhang, Z. X.; Jiang, Z.; Shangguan, W. Low-temperature Catalysts for VOCs Removal in Technology and Application: A State-of-the-Art Review. *Catal. Today* 2016, 264, 270−278.
(15) Vlè, G.; Bridier, B.; Wichert, J.; Pérez-Ramírez, J. Ceria in Hydrogenation Catalysis: High Selectivity in the Conversion of Alkynes to Olefins. *Angew. Chem., Int. Ed.* 2012, 51, 8620−8623.
(16) Zhang, S.; Chang, C.-R.; Huang, Z.-Q.; Li, J.; Wu, Z.; Ma, Y.; Zhang, Z.; Wang, Y.; Qu, Y. High Catalytic Activity and Chemo-selectivity of Sub-Nanometric Pd Clusters on Porous Nanorods of CeO₂ for Hydrogenation of Nitroaromatics. *J. Am. Chem. Soc.* 2016, 138, 2629−2637.
(17) Zhang, S.; Chang, C. R.; Huang, Z. Q.; Ma, Y. Y.; Gao, W.; Li, J.; Qin, Y. Q. Visible-Light-Activated Suzuki-Miyaura Coupling Reactions of Aryl Chlorides over the Multifunctional Pd/Au/porous Nanorods of CeO₂ Catalyst. *ACS Catal.* 2015, 5, 6481−6488.
(18) Tan, H.; Wang, J.; Yu, S.; Zhou, K. Support-Morphology-Dependent Catalytic Activity of Pd/CeO₂ for Formaldehyde Oxidation. *Environ. Sci. Technol.* 2015, 49, 8675−8682.
(19) Li, G.; Wang, Q.; Zhao, B.; Zhou, R. Promoting Effect of Synthesis Method on the Property of Nickel Oxide Doped CeO₂−ZrO₂ and the Catalytic Behaviour of Pd-only Three-Way Catalysts. *Appl. Catal., B* 2011, 105, 151−162.
(20) Zhang, N.; Xu, Y.-J. Aggregation- and Leaching-Resistant, Reusable, and Multifunctional Pd@CeO₂ as a Robust Nanocatalyst Achieved by a Hollow Core−Shell Strategy. *Chem. Mater.* 2013, 25, 1979−1988.
(21) Chen, C.; Fang, X.; Wu, B.; Huang, L.; Zheng, N. A Multi-Yolk−Shell Structured Nanocatalyst Containing Sub-10 nm Pd Nanoparticles in Porous CeO₂. *ChemCatChem* 2012, 4, 1578−1586.
(22) Zhang, N.; Liu, S.; Fu, X.; Xu, Y.-J. A Simple Strategy for Fabrication of “Plum-Pudding” Type Pd@CeO₂ Semiconductor Nanocomposite as a Visible-Light-Driven Photocatalyst for Selective Oxidation. *J. Phys. Chem. C* 2011, 115, 22901−22909.
(23) Cargnello, M.; Montini, T.; Polizzi, S.; Wieder, N. L.; Gorte, R. J.; Graziani, M.; Fornasiero, P. Novel Embedded Pd@CeO₂ Catalysts: A Way to Active and Stable Catalysts. *Dalton Trans.* 2010, 39, 2122−2127.
(24) Song, S.; Wang, X.; Zhang, H. CeO₂-Encapsulated Noble Metal Nanocatalysts: Enhanced Activity and Stability for Catalytic Application. *NPG Asia Mater.* 2015, 7, No. e179, DOI: 10.1038/asiam.2015.27.
(25) Cargnello, M.; Wieder, N. L.; Montini, T.; Gorte, R. J.; Fornasiero, P. Synthesis of Dispersible Pd@CeO$_2$ Core–Shell Nanostructures by Self-Assembly. *J. Am. Chem. Soc.* 2010, 132, 1402–1409.

(26) Adjani, L.; Bennett, D. A.; Chen, C.; Yu, A. S.; Cargnello, M.; Fornasiero, P.; Gorte, R. J.; Vois, J. M. Exceptional Thermal Stability of Pd@CeO$_2$ Core–Shell Catalyst Nanostructures Grafted onto an Oxide Surface. *Nano Lett.* 2013, 13, 2252–2257.

(27) Monai, M.; Montini, T.; Chen, C.; Fonda, E.; Gorte, R. J.; Fornasiero, P. Methane Catalytic Combustion over Hierarchical Pd@CeO$_2$/Si-Al$_2$O$_3$: Effect of the Presence of Water. *ChemCatChem* 2015, 7, 2038–2046.

(28) Wang, X.; Liu, D.; Li, J.; Zhen, J.; Wang, F.; Zhang, H. γ-Al$_2$O$_3$-Supported Pd@CeO$_2$ Core@Shell Nanospheres: Salting-out Assisted Growth and Self-assembly, and Their Catalytic Performance in CO Oxidation. *Chem. Sci.* 2015, 6, 2877–2884.

(29) Wang, X.; Zhang, Y.; Song, S.; Yang, X.; Wang, Z.; Jin, R.; Zhang, H. L-Arginine-Triggered Self-Assembly of CeO$_2$ Nanosheets on Palladium Nanoparticles in Water. *Angew. Chem., Int. Ed.* 2016, 55, 4543–4546.

(30) Du, C.; Guo, Y.; Guo, Y.; Gong, X.-Q.; Lu, G. Polymer-templated synthesis of hollow Pd–CeO$_2$ nanocomposite spheres and their catalytic activity and thermal stability. *J. Mater. Chem. A* 2015, 3, 23230–23239.

(31) Chen, C.; Nan, C.; Wang, D.; Su, Q.; Duan, H.; Liu, X.; Zhang, L.; Chu, D.; Song, W.; Peng, Q.; Li, Y. Mesoporous Multicomponent Nanocomposite Colloidal Spheres: Ideal High-Temperature Stable Model Catalysts. *Angew. Chem., Int. Ed.* 2011, 50, 3725–3729.

(32) Wang, F.; Li, W.; Feng, X.; Liu, D.; Zhang, Y. Decoration of Pt on Cu/Co double-doped CeO$_2$ nanospheres and their greatly enhanced catalytic activity. *Chem. Sci.* 2016, 7, 1867–1873.

(33) Wang, X.; Liu, D.; Song, S.; Zhang, H. Pt@CeO$_2$ Multicore@Shell Self-Assembled Nanospheres: Clean Synthesis, Structure Optimization, and Catalytic Applications. *J. Am. Chem. Soc.* 2013, 135, 15864–15872.

(34) Aguirre-Díaz, L. M.; Gándara, F.; Iglesias, M.; Snejko, N.; Gutiérrez-Puebla, E.; Monge, M. A. Tunable Catalytic Activity of Solid Solution Metal–Organic Frameworks in One-Pot Multicomponent Reactions. *J. Am. Chem. Soc.* 2015, 137, 6132–6135.

(35) Zhang, W.; Chi, Z.-X.; Mao, W.-X.; Lv, R.-W.; Cao, A.-M.; Wan, L.-J. One-Nanometer-Precision Control of Al$_2$O$_3$ Nanoshells through a Solution-Based Synthesis Route. *Angew. Chem., Int. Ed.* 2014, 53, 12776–12780.

(36) Sato, K.; Arai, M.; Valmalette, J.-C.; Abe, H. Surface Capping-Assisted Hydrothermal Growth of Gadolinium-Doped CeO$_2$ Nanocrystals Dispersible in Aqueous Solutions. *Langmuir* 2014, 30, 12049–12056.

(37) Zhou, Y.; Lawrence, N. J.; Wu, T.-S.; Liu, J.; Kent, P.; Soo, Y.-L.; Cheung, C. L. Pd/γ-Fe$_2$O$_3$-Nanorod Catalyst for CO Oxidation: Insights into the Origin of Their Regenerative Ability at Room Temperature. *ChemCatChem* 2014, 6, 2937–2946.

(38) Hu, Z.; Li, X.; Meng, D.; Guo, Y.; Guo, Y.; Lu, G. Effect of Ceria Crystal Plane on the Physicochemical and Catalytic Properties of Pd/ CeO$_2$ for CO and Propane Oxidation. *ACS Catal.* 2016, 6, 2265–2279.

(39) Zhang, S.; Chen, C.; Cargnello, M.; Fornasiero, P.; Gorte, R. J.; Graham, G. W.; Pan, X. Dynamic Structural Evolution of Supported Palladium–Ceria Core–Shell Catalysts Revealed by In Situ Electron Microscopy. *Nat. Commun.* 2015, 6, 7778.

(40) Kurnatowska, M.; Kepinski, L.; Mista, W. Structure Evolution of Nanocrystalline Ce$_6$Pd$_{32}$O$_{102}$ Mixed Oxide in Oxidizing and Reducing Atmosphere: Reduction-Induced Activity in Low-temperature CO Oxidation. *Appl. Catal., B* 2012, 117–118, 135–147.

(41) Satsuma, A.; Osaki, K.; Yanagihara, M.; Ohyama, J.; Shimizu, K. Activity controlling factors for low-temperature oxidation of CO over supported Pd catalysts. *Appl. Catal., B* 2013, 132–133, 511–518.

(42) Li, G.; Li, L.; Yuan, Y.; Shi, J.; Yuan, Y.; Li, Y.; Zhao, W.; Shi, J. Highly Efficient Mesoporous Pd/γ-Fe$_2$O$_3$ Catalyst for Low Temperature CO Oxidation Especially Under Moisture Condition. *Appl. Catal., B* 2014, 158–159, 341–347.

(43) Wang, S.-Y.; Li, N.; Zhou, R.-M.; Jin, L.-Y.; Hu, G.-S.; Lu, J.-Q.; Luo, M.-F. Comparing the CO Oxidation Activity of free PdO and Pd$_2$O$_3$.