Highly sensitive room-temperature CO gas sensors: Pt and Pd nanoparticledecorated In$_2$O$_3$ flower-like nanobundles

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We discovered a promising room-temperature CO sensing capability of Pt or Pd nanoparticlefunctionalized In$_2$O$_3$ flower-like nanobundles (NBs) composed of parallel nanoparticle-chains. The highly branched In$_2$O$_3$ flower-like NBs synthesized through the process of hydrolysis of InCl$_3$ in NaBH$_4$ aqueous solution and the subsequent heat treatment provided an extremely porous matrix required for high-performance gas sensors. The Pt and Pd nanoparticles uniformly deposited on the In$_2$O$_3$ NBs evidently play a key role in enhancing the sensitivity and response, and most importantly lowering the operating temperature to 25 $^\circ$C for the detection of CO gas. The present innovative hybrid nanostructures offer a potential platform for room-temperature oxide-based gas nanosensors. Moreover, the resulting sensing behaviours lead to a better understanding of using and designing such nanosensors for a number of gas sensory applications.

Introduction

Previous studies on the optimal design of oxide nanoparticle-based gas sensors have proved that the sensing response and sensitivity can be simultaneously and significantly enhanced when the particle size is close to twice the thickness of the carrier depletion layer. However, massive aggregation which occurs in either nanoparticle synthesis or the following film-casting processes, especially with tiny particles, will greatly reduce the surface areas and gas transporting paths and thus seriously suppress the sensing performance. A porous configuration in sensing layers is considered as the most fundamental strategy to efficiently provide larger active surface areas for gas sensors, but precisely and individually controlling the positions of each tiny nanoparticle for integrating the desirous porous structures using bottom-up techniques remains a challenge.

A practical and simple way to construct the porous sensing layers can be realized by synthesizing isolated and branched secondary nanoassemblies composed of numerous primary nanounits for the subsequent film formation procedures. This approach can generally create a high level of porous structures in sensing layers no matter what techniques are applied, for instance print-screen or dropping methods, and thus give more flexibility to manufacturing. As a result, the sensing performance should be mainly dominated by the architectures of the secondary nanoassemblies. Recently, various oxide nanoassemblies have been highlighted as promising candidate materials for gas sensing applications, such as hierarchical porous SnO$_2$, hierarchical In$_2$O$_3$ microspheres, and hierarchical assembly of hollow Cu$_2$O microspheres, and hierarchical porous ZnO nanostructures.

The operating temperature is another important issue for gas sensors. Typically, oxide sensors need to operate at a high temperature because reactions for both adsorption and desorption of dissociated gas species on oxide surfaces need to be thermally activated. A room-temperature sensor system could simplify device design by eliminating the heater component, save electrical power, be assembled on flexible polymer substrates, and most importantly, avoid triggering an explosion in an explosive environment. In recent years, many researchers have made tremendous effort towards the development of room-temperature oxide gas sensors in various ways. Among these latest studies, however, only a little effort has been made for detecting CO gas. To date, it has been experimentally proven that metallic catalysts such as Ag, Au, Pd, Pt, and Rh nanoparticles loaded onto the surfaces of the oxide supports serve as sensitizers or promoters, dramatically improving the sensing response, recovery time, and sensitivity and simultaneously decreasing the operating temperature. Therefore, it could be expected that a promising room-temperature sensor system for detecting CO may be obtained by combining oxide nanoassemblies constructed of desirably tiny oxide nanocrystals and catalytic noble-metal nanoparticles.

In this work, we newly designed, developed and then reproducibly synthesized novel three-dimensional (3-D) In$_2$O$_3$ flower-like nanobundles (NBs) composed of highly oriented nanoparticle-chains as one of the most promising hierarchical structures for manufacturing porous gas sensors. Based on this success, the innovative hybrid nanostructures were fabricated by decorating tiny Pt or Pd nanoparticles onto the In$_2$O$_3$...
flower-like NBs via low-temperature solution approaches and were then employed for reliably detecting CO gas at room temperature. To our knowledge, many hybrid nanostructures such as Pt–NiO nanotubes,26 Pt–ZnO nanowires,27 Pt–WO3 nanoparticles,28 Pd–SnO2 nanorods,29 and Pd–ZnO nanowires30 seem to be insensitive to CO gas. The present result is the first to show such a high CO sensing response and fast recovery/recovery speed using Pt or Pd nanoparticle-decorated In2O3 flower-like NBs operated at room temperature.

**Experimental section**

**Chemicals**

All chemicals used in this work were of analytical grade without further purification.

**Synthesis of In2O3 flower-like nanobundles**

First, 0.1 mmol of InCl3·4H2O was dissolved in 25 ml of distilled water. On adding 25 ml of NaBH4 solution (0.05 mM) to the prepared InCl3 solutions kept at 50 °C under stable stirring, white precipitates immediately formed. The precipitates were collected by centrifugation and were then repeatedly washed with distilled water to remove residual chemicals. After drying at 60 °C in air, the samples were annealed at 500 °C in air for 2 h to obtain In2O3 flower-like NBs.

**Decoration of In2O3 nanobundles with Pd and Pt nanoparticles**

The decoration engineering can be successfully achieved through the modified Wen34 and Pu35 methods. The In2O3 flower-like NBs were first irradiated by UV light (254 nm) for 3 h. Then the prepared In2O3 solutions kept at 50 °C under stable stirring, white precipitates immediately formed. The precipitates were collected by centrifugation and were then repeatedly washed with distilled water to remove residual chemicals. After drying at 60 °C in air, the samples were annealed at 500 °C in air for 2 h to obtain In2O3 flower-like NBs.

**Structural characterization**

The crystal structures of the prepared NBs were identified by X-ray diffraction (XRD, D2 Phaser, Bruker AXS) with Cu Kα radiation (λ = 1.5418 Å). The morphologies were characterized by field-emission scanning electron microscopy (FESEM, JEOL JSM-6700F). For high-resolution transmission electron microscopy (HRTEM, JEOL-2010) and selected area electron diffraction (SAED), a few drops of the as-synthesized products were deposited onto carbon-coated copper grids and dried at room temperature in vacuum. The Brunauer–Emmett–Teller (BET) surface areas of prepared In2O3 nanostructure were determined by nitrogen adsorption–desorption isotherms at 77 K (NOVA 1000e).

**Fabrication and performance of gas sensors**

To fabricate gas sensor devices, the Pd and Pt decorated NBs were ultrasonically re-suspended in distilled water and were then deposited onto an Al2O3 substrate with well patterned Pt electrodes. The prepared gas sensor was placed in a sealed chamber at room temperature to facilitate gas sensing tests. The sensing measurements were performed on an electrochemical analyzer (CH instruments, I600) by monitoring the change in the electrical current in air or in carbon monoxide (CO) gas. The gas sensitivity is defined as $S = I_d/I_a$, where $I_d$ and $I_a$ are the sensing electrical currents in air and in CO, respectively.

**Results and discussion**

Fig. 1a and b respectively show the SEM image and the corresponding XRD pattern of the In(OH)3 NBs successfully synthesized in the NaBH4 aqueous solutions. Obviously, numerous needle-shaped branches are conjoined at one centre point and are radially arranged to form a flower-like morphology, denoted as the flower-like NBs, without the assistance of surfactants and templates. In Fig. 1b, most traceable XRD peaks can be indexed by In(OH)3 (JCPDS 76-1464), excluding a few peaks around 27–29° which are coincident with some chloride and boron derivatives. In this case, the NaBH4 plays an important role in the formation of In(OH)3 NBs. With the addition of NaBH4 aqueous solutions, In3+ ions can be hydrolyzed to form hydroxide compounds, In(OH)3. Similar hydrolysis reactions have also been observed in other cases of metal ions in the presence of NaBH4 aqueous solutions.31 The possible growth mechanism of the flower-like In(OH)3 NBs is proposed below. Previous studies have demonstrated that In(OH)3 building blocks, usually one-dimensional (1-D) ones, tend to spontaneously construct oriented aggregates due to the existence of hydroxyl groups (–OH) on the surfaces of building blocks.34,35 When In(OH)3 nanochains come close to each other, a condensation reaction could occur between the –OH groups on the surfaces to build chemical bonds, o xo (O–) bridges, for the oriented attachment.34,35

By high-temperature heat treatments, the flower-like shape of the In(OH)3 NBs was maintained as shown in Fig. 2a, but In(OH)3 completely transformed into In2O3 as evidenced by the XRD pattern displayed in Fig. 2b. From the TEM image of single flower-like In2O3 NBs shown in Fig. 2c, it can be seen that numerous 1-D branches with similar lengths and widths are radially conjoined and most of the branches have relatively sharp tips. Looking closely at the tip part as shown in Fig. 2d reveals that great numbers of straight nanochains are closely bundled in parallel and each nanochain is composed of nanoparticles which are regularly assembled along the same direction. The broken rings observed in the SAED pattern (Fig. 2d, inset) indicate the
as 0.114 cm³ g⁻¹ and the total pore volume and average pore size are determined of an individual In₂O₃ NB, and (d) HRTEM lattice image and SAED pattern after annealing and (b) the corresponding XRD pattern. (c) TEM image of an individual In₂O₃ NB, and (d) HRTEM lattice image and SAED pattern of the tip of one branch of the NBs.

nature of the polycrystalline structure. Moreover, the distinct flower-like 3-D construction is greatly beneficial for the spontaneous formation of a very porous configuration in sensing layers when they randomly stack into films, as can be observed in Fig. 2a. The BET specific surface area is measured as 54.2 m² g⁻¹, and the total pore volume and average pore size are determined as 0.114 cm³ g⁻¹ and 33 nm, respectively, indicating that the present flower-like NBs contain numerous accessible channels and active sites. These features could contribute to the inward diffusion of targeted gas molecules and enhance the gas sensitivity significantly. As suggested above, the flower-like NB is a potential sensing nanostructure and can be a suitable oxide support to fabricate hybrid structures by decorating Pt or Pd nanoparticles for room-temperature CO sensing performance.

As can be observed in Fig. 3a and d, the wet decorating process does not seem to affect the morphology and dispersion of the preformed In₂O₃ NBs. Fig. 3b and c, as well as Fig. 3e and f, clearly evidence the successful decoration with numerous mono-

Fig. 2 (a) SEM morphology of randomly stacked In₂O₃ flower-like NBs after annealing and (b) the corresponding XRD pattern. (c) TEM image of an individual In₂O₃ NB, and (d) HRTEM lattice image and SAED pattern of the tip of one branch of the NBs.

dispersed tiny Pt and Pd nanoparticles, indicating that the decorating procedures have good reliability and flexibility. The average sizes of the Pt and Pd nanoparticles are ~3 nm and ~7 nm, respectively, and both sizes are considered suitable for exhibiting catalytic functions. The HRTEM image (Fig. 3c) focusing on one solidly attached Pt nanoparticle reveals an epitaxial relation between the Pt(111) and In₂O₃(222), suggesting the selective heteronucleation of Pt nanoparticles on specific sites of the In₂O₃ NBs. Similar to the case of Pd, the HRTEM lattice image of a single Pd nanoparticle (see Fig. 3f) also evidences a weak epitaxial Pd(111) on In₂O₃(222) growth. However, on looking closely at the larger Pd nanoparticle, multiple lattice domains can be observed within the Pd nanoparticle as well as along the heterointerface between Pd and In₂O₃.

The successful decoration of the well-dispersed Pt and Pd nanoparticles on the In₂O₃ supports is attributed to the UV-irradiation-assisted approach. One well-known decorating method that has been demonstrated for successfully decorating Au nanoparticles on TiO₂ supports basically comprises the following fundamental steps. First, the metal chloride precursor could form metal chloro-hydroxyl complexes by replacing Cl⁻ ligands with OH⁻ ones provided by NaOH. Second, the metal complexes would react with hydroxide on the surface of oxides and then attach onto oxides for the subsequent reduction process. However, it was found that the Pt and Pd nanoparticles tended to aggregate significantly if the UV irradiation was not applied before the decorating processes. The oxygen vacancies generated under the UV irradiation are beneficial for sticking hydroxyl groups (–OH) onto the surface of the In₂O₃ supports, and thus probably lead to uniform adsorption of the Pt and Pd complexes.

These remarkable structural parameters presented in the Pd–In₂O₃ and Pt–In₂O₃ flower-like NBs are expected to greatly contribute to the high sensing performance as many intensive studies have asserted that well-dispersed metallic nanoparticles on oxide supports would result in better sensing than agglomerative ones. Fig. 4a and b show the gas sensing response and recovery of the prepared Pd–In₂O₃ and Pt–In₂O₃ flower-like NBs, respectively, upon exposure to 50 ppm CO gas at 25 °C. Amazingly, the electrical currents immediately increase with the exposure to CO and rapidly recover to their initial state with the withdrawal of CO at room temperature, indicating that both sensors are very promising for the advanced heater-free nanosensors. As expected, the gas sensitivity (I/I₀) of the Pt–In₂O₃ sensors indeed reaches a very high value of ~2.8 and the corresponding response and recovery times are 60 s and 80 s, respectively. The Pd–In₂O₃ sensors have a further higher sensitivity of ~7.2 and faster response and recovery times of 50 s and 60 s, respectively. Both types of sensors show high sensitivity and remarkable response/recovery speeds at room temperature, whereas the bare In₂O₃ NBs present a negligibly low response even after 15 minutes of waiting with a much higher CO concentration of 200 ppm at the same temperature. Since these outstanding sensing features were not observed in the naked In₂O₃ NBs, it is evidently proven that the presence of catalytic Pt and Pd nanoparticles contributes to significantly lowering the operating temperature to 25 °C. The highly sensitive room-temperature sensors achieved by loading metal nanoparticles on In₂O₃ supports should involve two key mechanisms, namely...
The Pt and Pd nanoparticles would induce a well-known catalytic effect, which dramatically promotes dissociation of oxygen molecules in direct contact with the Pt and Pd catalysts, and thus largely increases the ionosorption of the dissociated oxygen species on the surface of the In$_2$O$_3$ NBs. In conclusion, the Pt or Pd nanoparticle-decorated In$_2$O$_3$ flower-like NBs have been newly designed based on our original structural considerations and successfully prepared via simple two-step approaches without the assistance of any surfactant or template with the goal of decreasing working temperature and enhancing performance in CO sensing. The innovative sensing platforms of both Pt– and Pt–In$_2$O$_3$ NBs demonstrated excellent response and recovery speeds as well as stable CO-concentration-dependent sensitivities at a very low operating temperature of 25 °C, in contrast to the insensitivity of the naked In$_2$O$_3$ NBs under the same analytical conditions. The potential combination of the well-dispersed tiny catalytic Pd or Pt nanoparticles as well as the distinct architectures of the In$_2$O$_3$ assemblies with extreme catalytic effects greatly withdraw electrons from In$_2$O$_3$ NBs and thus further expand the electron depleted regions, leading to higher Schottky barriers on the surface of In$_2$O$_3$ NBs.

When the Pt and Pd nanoparticles catalyze the oxidation of CO,*45–50 this causes the electrons trapped by the dissociated oxygen species to be released and then re-injected into the In$_2$O$_3$ NBs and thus the electrical resistance drastically decreases to create an outstanding sensing response. Based on the above considerations, it is reasonable to conclude that with the assistance of the catalytic metals, the oxide surfaces are strongly activated to drive the gas sensing reactions required at room temperature.

In the present sensing system, the geometry of the In$_2$O$_3$ flower-like NBs can offer great benefits in many respects. First, the In$_2$O$_3$ flower-like NBs are composed of numerous nanochains and nanocrystals which are tiny enough to be fully covered within the depletion regions formed. Such conditions are most desirable for an ultrahigh sensing response.*1–3 Second, the larger specific surface areas (54.2 m$^2$/g) provide more active sites not only for loading plenty of catalytic metals but also for enhancing adsorption and desorption of the dissociated gas species. Third, from the viewpoint of composition, In$_2$O$_3$ exhibits an excellent sensing performance for CO gas compared with other oxide materials.*44 All three factors also greatly contribute to the excellent sensing performance shown in this work.

The sensing performance, especially in a lower working temperature range, strongly depends on the metallic nanoparticles deposited instead of the oxide supports. Generally, metal nanoparticles with smaller sizes, narrow size distribution, and more uniform and isolated deposition on oxides have better results.*22 However, when carefully checking the relation between the nanostructures of the synthesized Pd and Pt nanoparticles and their slightly different sensing properties for detecting CO gas, it does not seem possible to infer a comprehensive rule covering different noble elements.

**Conclusions**

In conclusion, the Pt or Pd nanoparticle-decorated In$_2$O$_3$ flower-like NBs have been newly designed based on our original structural considerations and successfully prepared via simple two-step approaches without the assistance of any surfactant or template with the goal of decreasing working temperature and enhancing performance in CO sensing. The innovative sensing platforms of both Pt– and Pt–In$_2$O$_3$ NBs demonstrated excellent response and recovery speeds as well as stable CO-concentration-dependent sensitivities at a very low operating temperature of 25 °C, in contrast to the insensitivity of the naked In$_2$O$_3$ NBs under the same analytical conditions. The potential combination of the well-dispersed tiny catalytic Pd or Pt nanoparticles as well as the distinct architectures of the In$_2$O$_3$ assemblies with extreme catalytic effects greatly withdraw electrons from In$_2$O$_3$ NBs and thus further expand the electron depleted regions, leading to higher Schottky barriers on the surface of In$_2$O$_3$ NBs.

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The Pt– and Pd–In$_2$O$_3$ NBs show a higher sensitivity than Pt–In$_2$O$_3$. The sensing performance, especially in a lower working temperature range, strongly depends on the metallic nanoparticles deposited instead of the oxide supports. Generally, metal nanoparticles with smaller sizes, narrow size distribution, and more uniform and isolated deposition on oxides have better results.*22 However, when carefully checking the relation between the nanostructures of the synthesized Pd and Pt nanoparticles and their slightly different sensing properties for detecting CO gas, it does not seem possible to infer a comprehensive rule covering different noble elements.

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porosity in the stacking configuration plays a key role in these unusual findings. The present result evidently proves that the structural strategies used in this work would be one of the most promising guides for advanced nanoparticle-based gas sensors.

Notes and references