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In situ redox deposition of palladium nanoparticles on oxygen-deficient tungsten oxide as efficient hydrogenation catalysts†

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Noble metal/metal oxide support hybrid materials have attracted tremendous interest due to their wide applications in catalysis. Herein, we have developed a novel and surfactant-free method to prepare Pd/WO_{3-x} composite materials with clean surfaces. Oxygen-vacancy-rich WO_{3-x} nanowires (NWs) provide free electrons to reduce Pd²⁺, and surface-clean Pd nanoparticles (NPs) directly grow on WO_{3-x} surfaces through an *in situ* redox reaction between reductive WO_{3-x} and metal salt precursor (Na₂PdCl₄) in aqueous solution. The as-obtained Pd/WO_{3-x} nanocomposites show excellent catalytic activities for the hydrogenation of 4-nitrophenol (4-NP) and styrene. The apparent rate constant for 4-NP reduction is 0.045 s⁻¹, over the Pd/WO_{3-x} catalyst. The turnover frequency (TOF) value for styrene hydrogenation is 1074.5 h⁻¹, thus, exhibiting high catalytic performance. Moreover, the obtained Pd/WO_{3-x} catalyst exhibits good stability. Oxygen vacancies in WO_{3-x} NWs can accelerate electron transport and promote hydrogen adsorption and dissociation on the surface of the catalyst. The strong interaction between Pd NPs and WO_{3-x} support contributes to the excellent performance. Our work provides a novel and simple strategy to directly fabricate other-noble metal NP loaded oxygen-deficient metal oxides as highly efficient catalysts for chemical transformation.

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1. Introduction

The fabrication of noble metal/semiconductor metal oxide hybrid materials has aroused much attention because of their characteristic catalytic, optical and electrical properties.^{1,2} Among them, tungsten oxide-based materials have been widely studied due to potential applications such as gas sensors,³ photocatalysts,⁴ electrochromic devices⁵ and field-emission devices.⁶ Tungsten oxide is well documented for its non-stoichiometric character, where the crystal lattice can withstand a large number of oxygen vacancies.^{7,8} In particular, the presence of oxygen vacancies in tungsten oxide (WO_{3-x}, $x > 0$) receives considerable attention because of its unusual defect structure and unique characteristics. The existence of oxygen vacancies is reported to enhance catalytic activities by

narrowing the band gap and providing additional active catalytic sites.⁹ As a result, oxygen-deficient WO_{3-x} often displays critical impact on its physicochemical properties such as tuning optical absorption and increasing conductivity.

Moreover, recent reports have manifested that noble metal nanoparticle (NPs) loaded supports can function as effective catalysts in chemical and photochemical reactions.¹⁰ However, a serious problem in the application of supported catalysts is the agglomeration of noble metal NPs. Collectively, many researchers offer control over basic parameters such as particle size and shape to further enhance their properties by the passivation of specific surfactant or ligand molecules. These stabilizing molecules can be successfully removed by thermal and oxidative approaches, which usually requires a high temperature and often inevitably leads to an increase in the size of metal NPs.¹¹ It has already been proved beyond doubt that catalysis involving noble metal NPs depends on the size and the surface of the catalyst particle.¹² Therefore, direct deposition of noble metal NPs on oxygen-defective semiconductor *via* a facile method without any stabilizing agents is highly desired.

Surface oxygen vacancy sites are proved to have the reductive ability for the reduction of metal ions. Pan and co-workers reported the synthesis of TiO₂ with oxygen vacancies by a photocatalytic reaction, and noble-metal NPs (Ag, Pt, and Pd) can grow directly on TiO₂ surfaces *via* a redox reaction between reductive TiO₂ and metal salt precursors without any stabilizing

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agents or reducing agents.¹³ The obtained M-TiO₂ nanocomposites exhibit the enhancement in photocatalytic activities toward selective oxidation of benzyl alcohol and reduction of heavy metal Cr(VI) ions under visible light irradiation. Hu and co-workers prepared Au/TiO₂ nanocomposites through an *in situ* self-redox reaction between oxygen-defect-rich TiO_{2-x} and HAuCl₄ in aqueous solution.¹⁴ This process is named as a self-redox process without the use of any reducing agents or surfactant molecules. The Au loaded TiO₂ nanosheets display greatly enhanced photocatalytic performance in the photocatalytic degradation of toxic pollutants.

There are few reports on the preparation of tungsten oxide nanostructures with oxygen vacancies and noble metal NPs directly grow on tungsten oxide surfaces. Herein, we report a novel and simple route for synthesis of WO_{3-x} single crystal NWs with abundant oxygen vacancies. The main focus of this work is to find suitable conditions for an effective introduction of oxygen vacancies into metal oxides. It is reported that vacuum or hydrogen treatment can induce the surface disorder of metal oxides.^{15,16} Here, we develop a surfactant-free way to obtain oxygen-vacancy-rich WO_{3-x} NWs by one-step hydrothermal route. More importantly, it was reported that non-stoichiometric WO_{3-x} is thermodynamically stable at room temperature, and the quenching of oxygen vacancies was found to occur only upon annealing in air at a temperature higher than 400 °C.¹⁷ Pd NPs grow directly on WO_{3-x} surfaces through an *in situ* redox reaction between reductive WO_{3-x} and metal salt precursor (Na₂PdCl₄) in aqueous solution, and strong metal-support interaction occurs. The resulting Pd/WO_{3-x} nanocatalyst is used as catalyst for hydrogenation reactions such as the reduction of nitro-aromatic compounds to amines and the hydrogenation of styrene.

2. Experimental

2.1 Materials

Na₂PdCl₄ was purchased from Alfa Aesar. Ammonium tungstate hydrate, citric acid and NaBH₄ were obtained from Sinopharm Chemical Reagent Co., Ltd. All chemicals were used as received without further purification. Double distilled water was used in all experiments.

2.2 Preparation

To synthesize WO_{3-x} nanowires, 0.913 g of ammonium tungstate hydrate (0.3 mmol) and 0.183 g of citric acid were dissolved in distilled water under stirring. Subsequently, the mixture was transferred to a 100 mL Teflon liner and then heated at 160 °C for 12 h. After cooling to room temperature naturally, the deep-blue precipitates were collected and washed with distilled water and ethanol several times. The final products were dried in vacuum at 60 °C overnight. In the loading process, a facile method was used to load the palladium particles for direct growth on WO_{3-x} nanowires *in situ*. The synthesized WO_{3-x} (0.100 g) was dispersed in distilled water (20 mL), and an amount of Na₂PdCl₄ (10 mM) was added dropwise with constant stirring at room temperature overnight. Finally, the deep-gray solid was collected and dried under vacuum at 60 °C overnight.

2.3 Catalyst characterization

X-ray diffraction (XRD) patterns were recorded using a Rigaku/Max-3A X-ray diffractometer with Cu K α radiation (λ = 1.54178 Å). Field emission scanning electron microscopy observations (FE-SEM) were obtained using a JEOLJSM-6300F field-emission scanning electron microscope (15 kV). The transmission electron microscopy (TEM) images, high resolution transmission electron microscopy (HRTEM) images and elemental mapping were gathered on a JEOL JEM12100F field emission high resolution transmission electron microscope operated at 200 kV. X-ray photoelectron spectroscopy (XPS) measurements were done on a Perkin-Elmer RBD upgraded PHI-5000C ESCA system. The actual content of Pd was measured with a Thermo Scientific Plasma Quad 3 inductively-coupled plasma mass spectrometry (ICP-MS) after dissolving sample with a mixture of HCl and HNO₃ (3 : 1, volume ratio). Raman spectra were obtained at room temperature with a Perkin-Elmer 400F Raman spectrometer using a 514.5 nm laser beam. A Shimadzu spectrophotometer (Model 2501 PC) was used to record the UV-vis diffuse reflectance spectra of the samples in the region of 200 to 800 nm. The electron paramagnetic resonance (EPR) spectra were recorded on a JEOL JESFA200 EPR spectrometer (298 K, 9061 MHz, 0.998 mW, X-band).

2.4 Catalytic reduction of 4-nitrophenol

The catalytic properties of Pd/WO_{3-x} (Pd, 0.99 wt%) nanocomposites were investigated *via* the reduction of 4-nitrophenol to 4-aminophenol with NaBH₄ as the reductant under ambient temperature as a model reaction. First, 4-nitrophenol (150 μ L, 10 mmol L⁻¹) was added to NaBH₄ aqueous solution (8 mL, 0.1 mol L⁻¹), and the mixture was stirred for several minutes. Then Pd/WO_{3-x} nanocomposites aqueous suspension (50 μ L, 2.8 mg mL⁻¹) was added to the above solution, which was stirred until the bright yellow gradually changed to colorless. The reaction progress was monitored by measuring the UV-vis absorption spectra of the reaction solutions.

2.5 Catalytic hydrogenation of styrene

Pd/WO_{3-x} (5.5 mg), styrene (570 μ L, 5.0 mmol) and 1,3,5-trimethylbenzene (internal standard, 500 μ L, 3.6 mmol) were added into 10 mL ethanol. The air in the reactor was replaced by H₂. Then, the mixture was stirred in room temperature (about 30 °C) for the desired time. The product was analyzed by a gas chromatograph (GC) equipped with a flame ionization detector (FID). Gas chromatograms (GC) were obtained on Agilent HP-5, Gas Chromatograph with a SGE BP1 non-polar 100% dimethylpolysiloxane capillary column of (30 m \times 0.32 mm \times 0.25 μ m) dimensions. The method used for styrene hydrogenation is performed for 1 minute at 70 °C and a ramp of 10 °C min⁻¹ until 110 °C.

3. Results and discussion

The synthetic procedure of Pd/WO_{3-x} nanocatalyst is elucidated in Fig. 1. In the first step, WO_{3-x} nanowires (NWs) with abundant oxygen vacancies were synthesized from ammonium



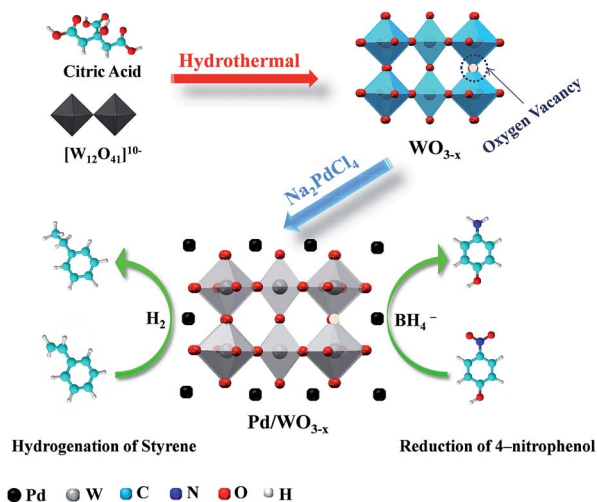


Fig. 1 Schematic illustration for the synthesis of oxygen-vacancy-rich WO_{3-x} NWs and $\text{Pd}/\text{WO}_{3-x}$ nanocomposites. The resulting $\text{Pd}/\text{WO}_{3-x}$ nanocatalyst is used for hydrogenation reactions of 4-nitrophenol (4-NP) and styrene.

tungstate hydrate and citric acid by a one-step hydrothermal treatment. Subsequently, the as-synthesized WO_{3-x} NWs and Na_2PdCl_4 were mixed in aqueous solution by stirring, Pd^{2+} ions were directly reduced by oxygen-vacancy-rich WO_{3-x} NWs, and surface-clean palladium nanoparticles (NPs) were deposited on WO_{3-x} surfaces at room temperature through an *in situ* redox reaction. The obtained $\text{Pd}/\text{WO}_{3-x}$ nanocatalyst was used to evaluate catalytic performance of the hydrogenation of styrene and 4-nitrophenol (see Experimental section).

The X-ray diffraction (XRD) analysis was performed to examine the crystalline structure of obtained WO_{3-x} NWs and $\text{Pd}/\text{WO}_{3-x}$, and the results are shown in Fig. 2a and S1.† The XRD pattern displays a pure phase of hexagonal crystalline structure with diffraction peaks assigned to tungsten trioxide (JCPDS no. 85-2459). The typical lattice constants were calculated to be $a = b = 7.326 \text{ \AA}$, $c = 7.669 \text{ \AA}$, confirming that WO_{3-x} was successfully synthesized by hydrothermal method. The diffraction peaks of Pd were not observed for $\text{Pd}/\text{WO}_{3-x}$ (Fig. 2a), likely due to its low weight content (2.3 wt% Pd) by inductively coupled plasma-atomic emission spectroscopy (ICP-AES) analysis.

The morphology and structural features of as-synthesized WO_{3-x} NWs were analyzed by scanning electron microscopy (SEM) and transmission electron microscopy (TEM). The SEM images shown in Fig. 2b and S2,† reveal WO_{3-x} NWs are uniform and very long. As estimated from the SEM image, the obtained WO_{3-x} NWs are 15–20 nm in diameters and 0.05–1.5 μm in lengths. TEM image further shows that the obtained WO_{3-x} is composed of a large number of radial NWs (Fig. S3a, ESI†). It is clearly seen from TEM images that the surfaces of WO_{3-x} NWs are decorated with Pd NPs (Fig. S3b, ESI†).

Clear lattice fringes measured with an interplanar lattice spacing of 0.367 nm for WO_{3-x} and 0.242 nm for Pd NPs can be unambiguously observed from the high-resolution TEM (HRTEM) images (Fig. 2c), which corresponds to the (110)

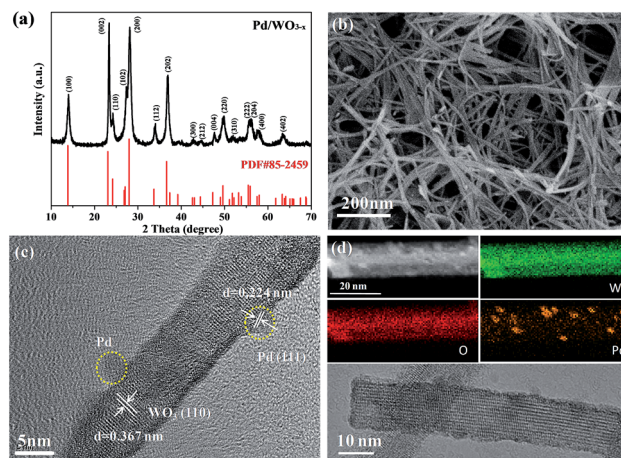


Fig. 2 (a) XRD pattern of the as-prepared $\text{Pd}/\text{WO}_{3-x}$. The standard card for hexagonal WO_3 (JCPDS no. 85-2459) is shown at the bottom. (b) SEM image of WO_{3-x} NWs. (c) HRTEM image of $\text{Pd}/\text{WO}_{3-x}$ nanocomposites. (d) STEM image and the corresponding elemental mappings of $\text{Pd}/\text{WO}_{3-x}$ (actual weight content of Pd, 2.3 wt%).

atomic plane of tungsten oxide and the (111) crystal plane of palladium, respectively. Simultaneously, the HRTEM images of WO_{3-x} also exhibit a wires-like morphology with an interfacial angle of 90° between (110) and (1–10) facets (Fig. S4, ESI†), corresponding to the lattice space of 0.367 nm and 0.320 nm, respectively. From these results, it is concluded that each hexagonal WO_{3-x} nanowire grew preferentially along the [110] direction. Meanwhile, the TEM and HRTEM images clearly exhibit that Pd NPs were well-dispersed on WO_{3-x} support with a narrow size distribution in the range of 2–5 nm. As expected, elemental mapping analysis (Fig. 2d) of the as-obtained $\text{Pd}/\text{WO}_{3-x}$ composites clearly displays W, O and Pd elements, confirming that the surfaces of WO_{3-x} NWs are decorated with small Pd NPs.

The compositions and valence states of WO_{3-x} NWs and $\text{Pd}/\text{WO}_{3-x}$ were investigated by X-ray photoelectron spectroscopy (XPS) measurements (Fig. 3). For WO_{3-x} NWs, the W 4f region exhibits two peaks at 37.5 eV and 35.4 eV, which are attributed to W^{6+} . The binding energies appeared at 36.5 eV and 34.6 eV are assigned to the lower oxidation state W^{5+} (Fig. 3a).^{18,19} W 4f peaks for $\text{Pd}/\text{WO}_{3-x}$ sample shift to lower binding energies as compared to bare WO_{3-x} (Fig. 3b) due to free electrons in oxygen vacancy abundant WO_{3-x} transfer to Pd. This result suggests the strong interaction occurs between Pd NPs and WO_{3-x} NWs.²⁰ The XPS results clearly demonstrate that W^{5+} was introduced into the final sample, consequently, oxygen vacancies were formed in WO_{3-x} NWs, which might have an effect on the catalytic activity. The content of W^{5+} was estimated to be about 16.5% in WO_{3-x} sample from XPS data. After the deposition of Pd NPs, the relative content of W^{5+} was reduced to about 11.9% from XPS analysis (Fig. S5, ESI†), suggesting Pd^{2+} ions were reduced by W^{5+} centers.

It is noted that the color of WO_{3-x} aqueous suspension changed from original blue to gray after adding a given concentration of Na_2PdCl_4 (Fig. S6, ESI†). This change in the color is attributed to the partial disappearance of oxygen



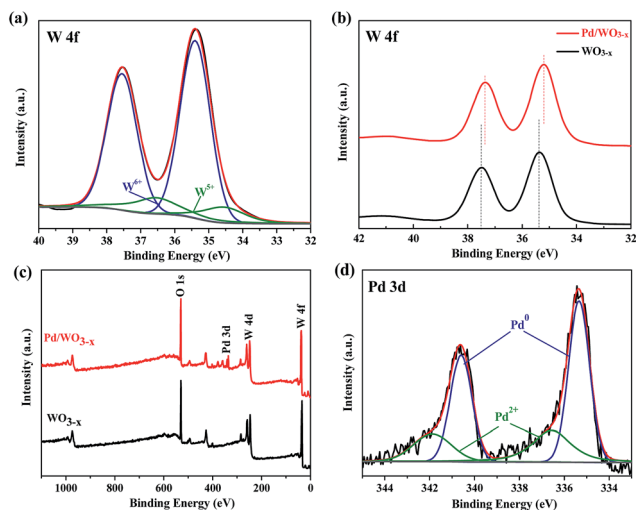


Fig. 3 (a) The W 4f spectrum of WO_{3-x} NWs. (b) The W 4f spectrum of $\text{Pd}/\text{WO}_{3-x}$ composites (red line) and WO_{3-x} (black line). (c) XPS survey scan spectra of WO_{3-x} NWs and $\text{Pd}/\text{WO}_{3-x}$ (Pd, 0.99 wt%) sample. (d) The Pd 3d spectrum of $\text{Pd}/\text{WO}_{3-x}$ sample.

vacancies and the formation of Pd NPs because oxygen vacancies provide free electrons for the reduction of Pd^{2+} ions. Furthermore, the color of the obtained WO_{3-x} had no significant change after at least six months, indicating that WO_{3-x} is stable in air. As shown in Fig. 3c, XPS survey spectrum of $\text{Pd}/\text{WO}_{3-x}$ sample clearly displays the presence of W, O and Pd elements, indicating the formation of $\text{Pd}/\text{WO}_{3-x}$. For high-resolution Pd 3d XPS spectrum of $\text{Pd}/\text{WO}_{3-x}$ (Fig. 3d), it can be seen that two peaks with binding energies of 335.3 eV and 340.6 eV are ascribed to $\text{Pd } 3d_{5/2}$ and $\text{Pd } 3d_{3/2}$ of Pd^0 , respectively.^{21,22} Moreover, the other two weak peaks appeared at 336.6 eV and 341.9 eV could be assigned to Pd^{2+} . The XPS results clearly reveal that most of Pd^{2+} ions were reduced to Pd^0 by oxygen-vacancy-rich WO_{3-x} NWs. Notably, the binding energies of Pd 3d signals for $\text{Pd}/\text{WO}_{3-x}$ sample has a positive-shift by about 0.3 eV compared to the standard peak position.²³ This shift in binding energies is influenced by the electronic modification of Pd by WO_{3-x} support and the size of metal particles,²⁴ indicating strong metal-support interaction (SMSI) occurs in $\text{Pd}/\text{WO}_{3-x}$, which leads to the enhanced catalytic performance of the obtained catalyst.

To further verify the formation of oxygen vacancies, photo-physical properties were characterized by UV-visible diffuse reflectance spectra (Fig. 4a). The commercial WO_3 only responds to visible light due to its intrinsic band gap (2.6 eV, Fig. 4b). For as-synthesized WO_{3-x} NWs, the strong absorption in the visible and near infrared (NIR) regions is clearly observed, thus providing clear evidence that the non-stoichiometric WO_{3-x} NWs contain a large number of oxygen vacancies.^{25,26} Moreover, in contrast to commercial WO_3 which appears as yellowish powders, the color of obtained WO_{3-x} NWs is deep blue because of the presence of abundant oxygen vacancies.²⁷ Raman spectroscopy was further performed to analyze the crystalline structure of WO_{3-x} NWs, which is sensitive to the surface oxygen vacancies (Fig. 4c). Raman spectrum of

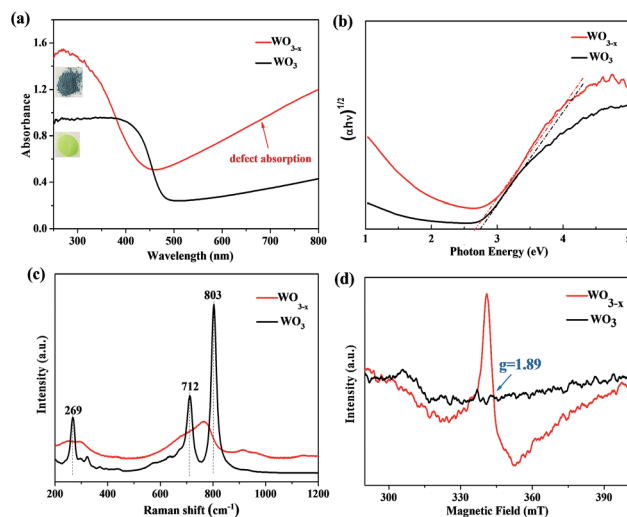


Fig. 4 (a) The UV-visible diffuse reflectance spectra for WO_{3-x} NWs (red line) and commercial WO_3 (black line). The insets in (a) show the corresponding photographs. (b) Curves of the Kubelka-Munk function plotted against the photon energy for WO_{3-x} and WO_3 . (c) Raman spectra of commercial WO_3 and as-synthesized WO_{3-x} NWs. (d) The X-band EPR spectra of deep blue WO_{3-x} NWs and commercial WO_3 as a reference sample recorded at $T = 298$ K.

commercial WO_3 is dominated by three characteristic vibration bands at 269, 712, and 803 cm^{-1} , corresponding to bending vibration of $\delta(\text{O}-\text{W}-\text{O})$ and the stretching vibration of $\nu(\text{W}-\text{O}-\text{W})$.²⁸ For WO_{3-x} NWs, Raman peaks are much broader and lower intensity as compared to commercial WO_3 . The broad characteristic peaks centered in the region of 200–400 cm^{-1} and 600–900 cm^{-1} result from the wide range of W–O–W bond lengths. This change is identified by the existence of $\text{W}^{5+}-\text{O}$ vibrations in WO_{3-x} sample,^{18,29} which is expected to increase the concentration of oxygen vacancies. To explore W^{5+} states in WO_{3-x} sample, we measured electron paramagnetic resonance (EPR) spectra performed at 298 K. As shown in Fig. 4d, a strong EPR signal observed at $g = 1.89$ is assigned to W^{5+} ,^{30,31} which is in agreement with the UV-vis spectra and XPS analysis. Interestingly, the EPR signal shows a significant decline after the deposition of Pd NPs (Fig. S7, ESI†), which indicates the concentration of oxygen vacancies decreases in WO_{3-x} NWs. These results, taken together, demonstrate the successful introduction of oxygen vacancies into tungsten oxide and *in situ* redox deposition of Pd onto WO_{3-x} surface with a strong interaction.

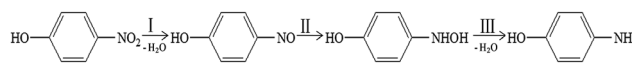
The reduction of nitro-aromatic compounds to amines is a significant process in synthetic organic chemistry of manufacturing many potent industrial intermediates.³² More recently, noble metal NPs like Au, Ag and Pd have been investigated to catalyze the well-known model reduction of 4-nitrophenol (4-NP) to 4-aminophenol (4-AP) owing to their efficient catalytic performance.^{33,34} Here, the catalytic activity of obtained $\text{Pd}/\text{WO}_{3-x}$ NWs was evaluated by the reduction of 4-NP to 4-AP in the presence of hydrogen source, NaBH_4 . When the reaction mixture contained 4-NP and NaBH_4 , the change in concentration was monitored by UV-vis absorption spectroscopy as



a function of time (Fig. 5). After adding an amount of NaBH_4 solution into 4-NP, the aqueous 4-NP solution undergone an immediate red-shift in the UV-vis absorbance band from 317 to 400 nm (Fig. S8a, ESI†), the change of color from light yellow to deep yellow could be attributed to the formation of 4-nitrophenolate ion.^{32,35} The intensity of the absorption peak of 4-NP at 400 nm was not reduced within 30 min in the presence of NaBH_4 without catalyst (Fig. S8b, ESI†). This result indicates that it is difficult for this reaction to proceed in the absence of catalyst. When a small amount of $\text{Pd}/\text{WO}_{3-x}$ catalyst was added into the reaction system, the absorbance peak of 4-NP at 400 nm rapidly decreased, suggesting the successive reduction of 4-NP. Meanwhile, a concomitant increased peak of 4-AP at 298 nm was observed, thus confirming the transformation from 4-NP to 4-AP (Fig. 5a). It is noted that, when $\text{Pd}/\text{WO}_{3-x}$ was added to the reaction system, the solution color quickly turned from deep yellow to colorless within 80 s. In addition, the rate constant k , calculated by the linear relationship between $\ln(C_t/C_0)$ and reaction time (Fig. 5b) for 4-NP reduction, was found to be 0.045 s^{-1} , corresponding to the first-order reaction kinetics. Additionally, bare WO_{3-x} NWs was also used to test the reaction under the same conditions. Almost no reaction was detected by monitoring UV-vis absorption (Fig. 5b), indicating that it is noble metal that effectively promotes the catalytic activity. The activity of our catalyst is comparable to those reported in the literature,^{36–38} as shown in Table S1 (ESI†).

The catalytic mechanism of the reduction of 4-NP is proposed in Scheme 1. NaBH_4 can react with water at room temperature to yield H_2 , then, $\text{Pd}/\text{WO}_{3-x}$ catalyst dissociates hydrogen, and Pd NPs act as electron relay system to transfer electrons donated by BH_4^- to the nitro groups of 4-NP, as a result, the nitro group was reduced to the nitroso group (I). Followed by formation of hydroxylamine (II) and then hydroxylamine was further reduced to the amino group (III) with the dehydration process, leading to the final 4-AP product.³⁹

To further investigate the catalytic activity of the resulting $\text{Pd}/\text{WO}_{3-x}$ nanocomposites, the catalytic hydrogenation of styrene was selected and carried out at room temperature and 1 atm H_2 . In terms of the hydrogenation of styrene, the noble metal based catalysts display a remarkable catalytic activity toward the hydrogenation of styrene.^{40,41} In this reaction,



Scheme 1 A plausible mechanism for the reduction of 4-NP by NaBH_4 over $\text{Pd}/\text{WO}_{3-x}$ catalyst.

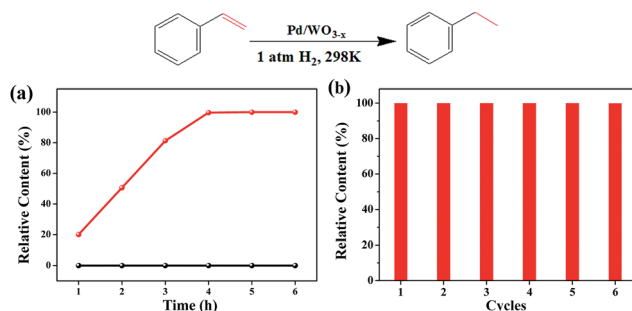


Fig. 6 (a) Typical conversion-time profile of ethylbenzene product from styrene hydrogenation over $\text{Pd}/\text{WO}_{3-x}$ catalyst (red line) and WO_{3-x} NWs (black line), respectively. (b) Bar diagram of recycling ability of $\text{Pd}/\text{WO}_{3-x}$ catalyst for hydrogenation reaction of styrene.

styrene is readily converted into ethylbenzene over catalyst by using hydrogen gas as the hydrogen source. Fig. 6a shows the relationship between conversion of styrene and time by using $\text{Pd}/\text{WO}_{3-x}$ (Pd, 2.3 wt%) catalyst. After 4 hours of reaction, the catalyst converted more than 99.5% of styrene to ethylbenzene without any detectable byproducts. Furthermore, we also evaluated the catalytic activity of bare WO_{3-x} NWs in the absence of noble metal. The result shows that almost no ethylbenzene was detected, indicating that it is difficult for the hydrogenation reaction to proceed without noble metal. The turnover frequency (TOF) value of $\text{Pd}/\text{WO}_{3-x}$ catalyst, defined as the number of moles of substrate converted per mol of palladium per hour, was calculated to be 1074.5 h^{-1} . The catalytic efficiency or TOF value of our $\text{Pd}/\text{WO}_{3-x}$ nanocatalyst is comparable to other previously reported Pd based catalysts,^{42,43} thus displaying a high activity. The reusability of $\text{Pd}/\text{WO}_{3-x}$ nanocatalyst in repeated catalytic reaction was performed in six consecutive runs (Fig. 6b). The results exhibit that the catalyst successfully recycled at least 6 times without obvious loss in activity. The high catalytic performance of $\text{Pd}/\text{WO}_{3-x}$ nanocatalyst is probably attributed to the following reasons. Firstly, oxygen deficient WO_{3-x} NWs directly reduce Pd^{2+} to metallic Pd NPs with clean surfaces deposited on WO_{3-x} support; secondly, strong metal-support interaction indeed contributes to the enhanced catalytic performance of supported Pd NPs; additionally, the smaller sized Pd NPs exhibit higher catalytic activities in general.

4. Conclusions

In summary, we have developed a novel and simple method to synthesize blue WO_{3-x} NWs with abundant oxygen vacancies. Notably, palladium NPs directly grow on WO_{3-x} surfaces through an *in situ* redox reaction between reductive WO_{3-x} and

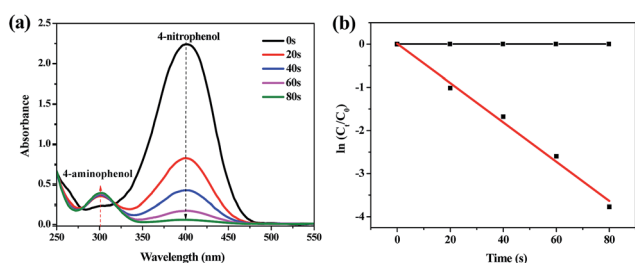


Fig. 5 (a) UV-visible absorption spectra of 4-NP reduced by NaBH_4 with $\text{Pd}/\text{WO}_{3-x}$ catalyst at room temperature. (b) Kinetic curves for the reduction of 4-NP catalyzed by $\text{Pd}/\text{WO}_{3-x}$ nanocatalyst (red line) and bare WO_{3-x} NWs (black line), respectively. C_t and C_0 are 4-NP concentrations at time t and 0, respectively.



Pd^{2+} ions in an aqueous solution. Thanks to this *in situ* redox deposition without any stabilizing agent or reducing agent, strong metal-support interaction occurs between Pd and WO_{3-x} . More importantly, the as-prepared Pd/ WO_{3-x} nano-catalyst shows excellent catalytic activity and good stability for the well-known model reduction of 4-nitrophenol to 4-aminophenol and the hydrogenation of styrene. The rate constant of Pd/ WO_{3-x} catalyst for 4-NP reduction reaction is 0.045 s^{-1} , and the turnover frequency value for the hydrogenation of styrene reaches as high as 1074.5 h^{-1} , comparable to that of recently reported Pd based catalysts. The significantly improved catalytic activity of the hydrogenation reaction is attributed to the small size of surface-clean Pd NPs and the strong interaction between metal and support. Owing to the synthetic simplicity and excellent properties, present work could provide important inspirations for developing other defect-type materials, and this novel route adopted here could be applied as a new strategy to prepare surface-clean metal nanoparticles loaded on support for various chemical transformations.

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