One-step Synthesis of Ordered Pd@TiO$_2$ Nanofibers Array Film as Outstanding NH$_3$ Gas Sensor at Room Temperature

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The one dimensional (1D) ordered porous Pd@TiO$_2$ nanofibers (NFs) array film have been fabricated via a facile one-step synthesis of the electrospinning approach. The Pd@TiO$_2$ NFs (PTND3) contained Pd (2.0 wt %) and C, N element (16.2 wt %) display high dispersion of Pd nanoparticles (NPs) on TiO$_2$ NFs. Adding Pd meshed with C, N element to TiO$_2$ based NFs might contribute to generation of Lewis acid sites and Brønsted acid sites, which have been recently shown to enhance NH$_3$ adsorption-desorption ability; Pd NPs could increase the quantity of adsorbed O$_2$ on the surface of TiO$_2$ based NFs, and accelerated the O$_2$ molecule-ion conversion rate, enhanced the ability of electron transmission. The response time of PTND3 sensor towards 100 ppm NH$_3$ is only 3 s at room temperature (RT). Meantime, the response and response time of the PTND3 to the NH$_3$ is 1 and 14s even at the concentration of 100 ppb. Therefore, the ordered Pd@TiO$_2$ NFs array NH$_3$ sensor display great potential for practical applications.

NH$_3$ sensor is of paramount importance in explosives detection, since it is a decomposition product of many explosive formulations and improvised explosive devices. It is also very useful in monitoring pollutants emitted by industrial and recycling processes.

Semiconducting metal oxides (SMO) are frequently used as gas-sensing materials. The suitability of a SMO material as a chemiresistor depends on several factors, many of which are correlated with each other. The most obvious factor is the chemical composition. Among the variety of appropriate materials, TiO$_2$, as an important non-toxic semiconductor, shows wide potential applications in the sensing field owing to their advantages of earth abundance, chemical and thermal stability$^{1-3}$. Thus, TiO$_2$-based gas sensors have also been explored quite extensively. Recent years, 1D ordered porous arrays have been the subject of extensive research due to their distinctive structural features and superior performances$^{6,7}$. However, there are still few applications of 1D ordered porous arrays in gas sensing. Chemiresistor arrays with pattern recognition techniques has been demonstrated to be powerful for addressing some of the challenging issues in chemical gas sensor$^8$. TiO$_2$ nanotubes and nano-fibers are very extensive used for sensing purposes. TiO$_2$ nanotubes synthesized by anodization$^{14,6,7,9-15}$ and electrospun$^{16-18}$ have higher ordered arrays, higher surface area, and desired location compared with electrospun TiO$_2$ nano-fibers, however, ordered template synthesis by anodization is a crucial and tedious technique compared with preparation methods concise and template-free electrospun TiO$_2$ nano-fibers and nanotubes. Not only that, the longer electrospun TiO$_2$ nano-fibers are in favour of preparing sensing device directly but also a lots of studies which detected gas sensitivity employing some apparatus (such as micro-hotplates$^{19-21}$ and so on) at high temperature have gained certain effect, operating at room temperature becomes more satisfied with commercial development prospect. In previous works, p-type TiO$_2$ has been researched$^{22-24}$ and In$_2$O$_3$/TiO$_2$ composite NFs were prepared in our group by electrospinning and exhibited better sensing performance for NO$_2$ at

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ies have provided insights into the interaction of Pd NPs with TiO$_2$, and our studies further investigated the combined effect of the noble metal with the main sensing material and the auxiliary elements of C, N is comparatively rare.

In this paper, we suggested a one-step method to fabricate 1D ordered porous Pd@TiO$_2$ NFs array film, with remaining small fraction of C, N element under insufficient oxygen calcining (see Fig. 1). The present studies have provided insights into the interaction of Pd NPs with TiO$_2$, and our studies further investigated the increased sensing properties on NH$_3$ which was the result of formation of acid sites. And Pd NPs as well as C, N element on the substrate of TiO$_2$ might play an important role in the generation of Lewis acid sites and Bronsted acid sites, which can greatly improve the NH$_3$ adsorption-desorption ability of gas sensing materials. Therefore, high-powered Pd@TiO$_2$ NFs NH$_3$ sensor displays great potential for practical applications.

**Results**

**Structure and morphology of samples.** The XRD patterns of pure TiO$_2$ and Pd@TiO$_2$ NFs are shown in Fig. 2a. It is obvious that the peaks at 20 values of 25.3, 37.8, 48.0, 55.1, 62.7 and 75.1° can be indexed to (101), (004), (200), (211), (204) and (215) crystal planes of anatase (JCPDS no. 71–1166) [37], and the peaks at 20 values of 27.4, 36.1, 41.2, 44.0, 54.3, 56.6, 64.0 and 69.0° can be indexed to (110), (101), (111), (210), (211), (220), (310) and (301) crystal planes of rutile (JCPDS no. 72–1148) [38], respectively. Meanwhile, XRD patterns of pPTND3 NRs is shown in Fig. S1. Notably, no typical diffraction peaks belonging to Pd (Pd or PdO) are observed in all of the samples, and it is indicated that Pd species are low mass loading and highly dispersed on the TiO$_2$ NFs.

Figure 2b,c show the N$_2$ adsorption-desorption isotherm and the BET pore-size distribution for samples. The isotherms of samples show a typical type IV with H3-type (PTND1 and PTND2) and H4-type (pure TiO$_2$, PTND3 and PTND4) hysteresis loop, which is characteristic of mesoporous material [39]. The BET surface areas and the dominated mesopores are 107.8 m$^2$·g$^{-1}$ and 5.8 nm for pure TiO$_2$, 82.5 m$^2$·g$^{-1}$ and 12.9 nm for PTND1; 86.7 m$^2$·g$^{-1}$ and 17.8 nm for PTND2; 100.8 m$^2$·g$^{-1}$ and 14.8 nm for PTND3; 89.0 m$^2$·g$^{-1}$ and 14.3 nm for PTND4, respectively. It is shows that the BET surface areas and the dominated mesopores results of Pd@TiO$_2$ NFs increase significantly than that of pure TiO$_2$. This illustrates that it is effective to improve cavity and interface structure with the addition of Pd NPs.

The morphology of calcined Pd@TiO$_2$ NFs are investigated by SEM. As shown in Fig. 2d,e and Fig. S2a,b,c, it can be clearly seen that 1D nanofibers of orderly arrangement are obtained on a large scale, which have typical lengths about 5–50 μm, and the diameter about 250 nm.

TEM investigations (Fig. 2f,g,h,i,j) are conducted for further analysis the structure of the PTND3 NFs. PTND3 NFs are consisted of Pd and TiO$_2$ polycrystalline nanoparticles as shown in Fig. 2h,i,j. During the calcination process, decomposition of the PVP polymer and Ti-O-Ti network results in the formation of porous and loosened structured of PTND3. The corresponding selected area electron diffraction (SAED) pattern shown in Fig. S3 confirms highly crystalline nature of anatase (101), (004) and rutile (101), (110) planes. HRTEM images in Fig. 2h,i,j confirm that the prepared porous PTND3 NFs have high crystalline nature with the lattice spacing of 0.350 (~0.354) nm corresponding to (101) plane of anatase, and 0.324 and 0.248 nm corresponding to (110) and (101) planes of rutile. Meanwhile, it can be clearly observed the existence of the lattice spacing of 0.225 nm corresponding to (111) planes of Pd NPs [41], which is illustrated that Pd NPs have been placed into the nanofibers successfully.

**IR spectroscopy (Fig. 3c,d,e) of nanofibers before calcination which precursor solution contains of PVP-H$_2$PdCl$_4$-TBT, thus, PVP-TBT, PVP-H$_2$PdCl$_4$ and PVP were measured. Contrast with the IR spectroscopy of pure PVP which contains amide group (N–C stretch of N–C=O (N–C$_2$) 1499 cm$^{-1}$, C=O (1639 cm$^{-1}$) and...
C–N stretch (N–C stretch of N=CH2 (N–C1, N–C2), 1277 and 1292 cm\(^{-1}\))\(^4\) (Scheme 2e), two differences could be clearly observed for that of the other samples 43. The carbonyl vibration band and hydrogen bonding band of PVP shifted to lower wavenumber (1659 to 1649 cm\(^{-1}\), 3452 to 3359 cm\(^{-1}\)) when the Pd\(^{2+}\) and (or) Ti-O-Ti network26 adhesive incorporated into the PVP (Fig. 3b,c,d). PVP can form hydrogen bonding between terminal hydroxyl groups with carbonyl side-groups of repeat units in PVP macromolecules, and metal ions due to the structure of polyvinyl skeleton with strong polar group (pyrrolidone ring) with lone pair of electrons from the nitrogen and oxygen atoms44 (Fig. 3b), this implied that the strength of the dipole - dipole interactions of the PVP units was reduced. Here, PVP acted as a template and played a self-assembly role in the solution. After calcination and before reduction, PdO and TiO\(_2\) NPs were taken shape in the absence of PVP. Therefore, the porous composites nanofibers were formed.

Gas Sensing Measurements. Gas response and response time are the basic parameters of gas sensors, rapid response to a target gas are demanded for practical application. Therefore, gas sensing performances of the PTNDs NFs sensor and the pPTND3 sensor were evaluated and compared its performance with those of the pure TiO\(_2\) sensor to NH\(_3\) at 26 °C (fabricated process of the thin film gas sensor in Supporting Information and the SEM image of the sensor coated with the composite thin film in Fig. S6). The tests show that there is no response to pure TiO\(_2\) sensor to 100 ppm NH\(_3\) (in Fig. S7). Figure 4a,b show the response and response time of PTND1, PTND2, PTND3 and PTND4 sensors to NH\(_3\) (see Fig. 4c, Fig. S8 and Table S3). The responses of four sensors to NH\(_3\) tend to become stronger with increasing NH\(_3\) concentration. Generally, adding appropriate amount of noble metals (such as Pd) to gas-sensing material can effectively reduce response times\(^4\),\(^5\). Obviously, the PTND3
Figure 3. (a) SEM image of electrospinning PTND3 NFs array after calcination, (b) The proposed crosslinking mechanism of the formation of electrospinning PTND3 NFs (PVP is displayed in red break line circle), (c) IR spectra of samples with wavenumber from 400 to 4000 cm\(^{-1}\) and (d) from 1200 to 1800 cm\(^{-1}\), which precursor solution contains of PVP-H\(_2\)PdCl\(_4\)-TBT, PVP-TBT, PVP-H\(_2\)PdCl\(_4\) and PVP, respectively. (e) IR spectra of PVP.

Figure 4. (a,b) The response time and gas response of samples to NH\(_3\), (c) Dynamic response of the PTND3 sensor to 0.1–100–0.1 ppm NH\(_3\) at 26 °C, the selectivity (d) and the stability (e) of PTND3 (inset in the top of (e) is Response/recovery curves of the sensor to 100 ppm NH\(_3\) at a certain interval) (RT: 26 °C, humidity of 30%).
sensor exhibits higher response and quickly response time than other sensors (PTND1, PTND2, PTND4 and pPTND3 see Table S3).

Figure 4c illuminates the response/recovery properties and response time of the PTND3 sensor to 0.1–100 ppm NH3 at 26 °C (the dynamic response diagrams and corresponding data of PTND1, PTND2, PTND4 and pPTND3 sensors to different concentrations of NH3 are in Fig S8, Table S3 and Table S4). Upon exposure to NH3, the resistance of PTND3 leads to a rapid increase, and after NH3 is removed, the resistance returns to the original value, which is the representative property of a p-type semiconductor. It is also observed that with the increase of NH3 concentration, the response of the PTND3 sensor is obviously increased, and the response time of the sensor is significantly reduced. The response time towards 100 ppm NH3 is about 3 s, and ~14 s to 0.1 ppm NH3. Meanwhile, the response of the PTND3 to NH3 is even at the concentration of 100 ppb. Not only that the PTND3 sensor shows a short-term reliability after two period tests, and an approximately linear response-concentration relationship in the range of 0.1–100 ppm NH3 to PTND3 sensor, as shows in Fig. 4c and in Fig. S9, which indicated that PTND3 has a potential application in combustible and noxious NH3 detection.

Since the selectivity and a long-term stability are also important parameters of gas sensor. The sensor must have rather high selectivity and stability for its application. Figure 4d depicts the selectivity of PTND3 sensor at 26 °C. The sensor shows high selectivity for NH3, and could distinguish an extremely small amount of 100 ppb NH3 compared with CO, CH4, H2, H2S, and NO2. The excellent selectivity further confirms that PTND3 sensor can be used as a promising material for NH3 sensor.

Figure 4e shows the response/recovery curves of the sensor to 100 ppm NH3 at 26 °C within 35 days. The tests were carried out in the same interval (7 days) under the same conditions. The PTND3 sensor shows a stable gas response with 6.97% after repeated tests. It is found that the signal has no obvious changes to repeat many times, which illustrated superior stability of PTND3 sensor.

It is true that humidity is prone to change significantly and rapidly in most environments. And the gas response of the PTND3 sensor had been tested under different humidity in order to measure its practicality at different environmental humidity. Fig. S10 shows the curves of the gas response for the PTND3 sensor to NH3 at 26 °C under different humidity. It is observed that with the increase of humidity from 20% to 60%, there is no obvious variation of the response of the PTND3 sensor, and the response of the sensor is significantly reduced when the humidity is increase to 70%. It illustrates that the PTND3 sensor has higher stability when the humidity is about 20%–60%.

In addition, there is no response for pure TiO2 sensor to 100 ppm NH3 (in Fig. S7), and the gas-sensing properties of PTND3 sensor (PTND3 was calcined under the mixed atmosphere of N2 and air, and it has 2% Pd) are better than pPTND3 sensor (pPTND3 was calcined under air, and it has 2% Pd too) (see Table S3 and Table S4). This demonstrated that, for PTND3 and pPTND3 sensor, the influenced factors for gas sensing properties was not just the electronic sensitization and chemical sensitization of Pd NPs27,36,47–50 in Pd@TiO2 NFs.

Discussion

The influence factors on gas sensing property. The elemental compositions of pPTND3 and PTND3 NFs are further researched because of calcining under the different atmosphere. Figure 5a and d describe the TG curves of pre-cal pPTND3 and pre-cal PTND3 NFs (TG treatment conditions and calcined products of above mentioned samples see in Table S2). Respectively, TG curve of the pre-cal PTND3 NFs was measured under air in the temperature range of 25–900 °C. Three stages of weight losses can be seen from Fig. 5a. The first stage of weight losses (ca. 13.3%) below 140 °C is mainly attributed to the loss of ethanol and water. The second loss stage (ca. 57.4%) from 270 °C to 460 °C is primarily due to the decomposition of PVP side chains. The third one (ca. 18.9%) in the range of 460–560 °C mostly results from the decomposition of main chain of PVP51,52. The total weight loss is about 89.6%. And at the same calcined condition, pure PVP was completely breakdown (as shown in Fig. S11). It can be roughly deduced that the 10.4% of the total weight of sample calcined is TiO2 and PdO.

TG curve of the pre-cal PTND3 NFs was measured under the mixture atmosphere of N2 and air in the temperature range of 25–900 °C. Three stages of weight losses in the range of 25–940 °C is about 89.5% (remaining 10.5% of TiO2 and PdO). The third one in the range of 490–940 °C chiefly results from the decomposition of main chain of PVP. Thus, there is some of C and N element (~16.2%) in remnant in PTND3 NFs after calcined at 600 °C under the N2 and air atmosphere and reduced (the TG curve of pure PVP calcined under the mixture atmosphere of N2 and air in Fig. S12).

IR and XPS spectra of samples (pure TiO2, PTND3 and pPTND3) adsorbed and desorbed NH3 were investigated in order to find the interactions between the samples and NH3, at 26 °C.

After pPTND3 adsorbed NH3 at 26 °C (in Fig. 5b,c) compare with pure TiO2 and apure TiO2 (see Fig. S13), it can be observed that, besides the peak of Ti-O, its IR spectra displays additional features corresponding to two distinct NH3 species: bands at 3340 and 1600 cm−1, they are assigned to adsorb NH3 on Lewis acid sites53,54.

In Fig. 5e,f, the IR spectra of PTND3 not only display bands at 3340 and 1600 cm−1 which are assigned to adsorb NH3 on Lewis acid sites, but also have bands at 3230 and 1470 cm−1 (see Fig. 5f) which are attributed to adsorb NH3 on Bronsted acid sites55. That means, both Lewis acid sites and Bronsted acid sites are existing in PTND3. This demonstrates that the synergy of Pd NPs and 16.2% of elements of C, N in PTND3 might play an important role in the generation of acid sites. Furthermore, to the curve of dPTND3 (Fig. 5f), NH3 species are disappeared basically.

In order to further research the effect of O, C and N element on the property of Pd@TiO2 NFs, XPS measurements were applied. As shown in Fig. 6a,b,c and in Fig. S14a,b, for pure TiO2, PTND3, aPTND3, pPTND3, and apPTND3, the O 1s spectrum can be recounted into three peaks: 529.5, 530.5 and 532.4 eV, which are attributed to lattice oxygen, oxygen defect and adsorbed oxygen. The oxygen defect of pPTND3, aPTND3, pure TiO2, pPTND3 and PTND3 are 24.7%, 25.0%, 25.6%, 37.4% and 39.8%, respectively. The increased contribution percentage of oxygen defect in PTND3 NFs and pPTND3 NFs can be considered that the effect of deformation of introducing Pd NFs and the remains of
Figure 5. (a,d) TG curve of pre-cal pPTND3 and pre-cal PTND3; (b,c) IR spectra curves of apPTND3 (pPTND3 adsorbed NH3) and dpPTND3 (pPTND3 desorbed NH3); (e,f) IR spectra curves of PTND3, aPTND3 (PTND3 adsorbed NH3) and dPTND3 (aPTND3 desorbed NH3) (RT: 26 °C, humidity of 30%, synthesis and treatment conditions of above mentioned samples see in Table S1).

Figure 6. Curve-fitted XPS spectra of O 1s (g–i), N 1s (j–l) and C 1s (m–o) of pure TiO2, PTND3 and aPTND3 (RT: 26 °C, humidity of 30%, synthesis and treatment conditions of above mentioned samples see Table S1).
elements of C, N in TiO2 might enhance the oxygen defect on the surface of TiO2 grains. Since more the oxygen defects, more will be the adsorption of gas25. In the meantime, the oxygen defect might directly or indirectly promote formation of Lewis acid sites or Bronsted acid sites, and improve their gas sensing performance.

As shown in Fig. 6d,e,f, N (1s) in the samples which calcinated under the mixture atmosphere of N2 and air was studied by means of XPS spectrum. The N (1s) XPS spectrum of pure TiO2 has only a little peak at 399.4 eV which is amino generally considered in non-reactive molecules. The N element which exists on the surface of PTND3 is amino iminic nitrogens -N = (397.9 eV), amino (399.4 eV) and -NH - (400.1 eV) states (see Fig. 6e). After NH3 adsorption, the weak NH4+ (401.8 eV) peak could be observed in aPTND3 (see Fig. 6f), expect amino (399.4 eV) and -NH - (400.1 eV) peaks56–58. It is consistent with the result of IR results, namely that Bronsted acid sites of PTND3 were formed because of existence of Pd NPs and elements of C and N, compared with pPTND3 and pure TiO2 (pPTND3, which was oxidized thoroughly, has no elements of N, and pure TiO2 has no elements of Pd). And ammonia can be adsorbed on the Bronsted acid sites forming NH4+ species and then the ability of adsorption NH3 will increase. Therefore, the response of PTND3 sensor enhanced compared with pPTND3 sensor.

In addition, the C 1s XPS spectrum of pure TiO2 can be deconvoluted into two peaks at 284.5 eV and 285.3 eV (Fig. 6g), which are attributed to the typical signals of C=C (sp2) and C-C (sp3), and PTND3 and pPTND3 are revealed from the C=C (sp2) at 284.5 eV, C-C (sp3) at 285.3 eV, C-O at 286.3 eV and COO at 288.5 eV (in Fig. 6h and Fig. S15a). Meanwhile, there are clearly exhibited C=C (sp2), C-C (sp3), C-O and C-NH x components in the sample of aPTND3 and pPTND3, which located at 284.5, 285.3, 286.3, and 288.0 eV54,56,61, respectively (in Fig. 6i and Fig. S15b). The C-NH x band located at 288.0 eV may be assigned to the adsorbed NH3 on the acid sites which formed on the surface of PTND3 and pPTND3 and resulted from synergy effect between Pd and C (or N) element. And this means that the response of PTND3 (in particular) and pPTND3 to NH3 was promoted up compared with pure TiO2.

Temperature-programmed desorption (TPD) is a surface science technique which has been used extensively to study surface reactions and molecular adsorption and desorption62. NH3-TPD were carried out to further investigate the acid strength of samples. Fig. S16 depicts the NH3-TPD patterns of both pure TiO2, PTND1, PTND2, PTND3, PTND4 and pPTND3. As shown in Fig. S16, the NH3 desorption peaks of pure TiO2 are too weak to be clearly observed, which suggested that pure TiO2 is shortage of acid sites. However for all of the Pd@TiO2, a larger peak was observed from 110 °C to approximately 450 °C compared with pure TiO2. Thus, it seemed that there was a broad distribution of weak and moderate acidity in the samples which contains Pd. As refer to the IR and XPS results (Figs 5 and 6), this implies that the peaks of Pd@TiO2 could be explained that the acidic sites were generated due to interacting between TiO2 and Pd. Furthermore, the increasing area of desorption peaks indicates that the acid strength is improved with the increasing of Pd amount, but the difference is not significant for Pd (2.0%) and Pd (2.5%). Meantime, there was weaker desorption peak at low temperature in the pPTND than that of PTND because of lackage of Bronsted acid sites of pPTND63. Therefore, the strong acidity decreased as the following order: PTND1 > PTND2 ≈ PTND3 > PTND4 > pPTND3 > TiO2.

Gas sensing mechanism. The NH3 sensing properties of the PTND3 sensor have been significantly improved compared with pure TiO2 and pPTND3 depend upon the presence and interaction with Pd NPs, N and C elements in TiO2 NFs. And the gas-sensing mechanism of PTND3 sensor was probed into on the relationship between its structure and properties.

Firstly, a pure TiO2 (600 °C for 4 h under the mixed atmosphere of N2 and air) sensor could generate oxygen ions (O2−) when it is placed in an open atmosphere at RT. Nevertheless, the tests showed that there was no response to pure TiO2 sensor to 100 ppm NH3 (in Supporting Information, Fig. S7). This illustrates that it is hard to directly interact between NH3 molecules and oxygen ions (O2−), and there is no obviously effect between TiO2 and remaining C (or N) element.

And, secondly, pPTND3 (Pd 2.0%, 600 °C for 4 h under air atmosphere) sensor could produce a large number of oxygen ions (O2−), when it is placed in an open atmosphere at RT. As we known, the Pd NPs on the surface of metal oxide material can form electronic sensitization mechanism, that is, oxygen molecules can rapidly dissociate and form oxygen ions (O2−) because of the catalytic action of Pd NPs77,31,33,34,36. Not only that, the response of the pPTND3 sensor to the NH3 is 1.43 at the concentration of 100 ppm, it illustrates that interaction between NH3 molecules and oxygen ions happened with the presence of Pd. But at the same time, pPTND3 and PTND3 have identical Pd and oxygen ions (O2−) content but different NH3 response (the response of the PTND3 sensor to 100 ppm NH3 is 6.97), it explains that it is hard to directly interact between NH3 molecules and oxygen ions (O2−), in addition, remaining C and N elements in PTND3 not contained in pPTND3 also play an active role on the performance of gas sensing in addition to Pd. In the meantime, from IR spectrum, pPTND3 displays bands at 3340 and 1600 cm−1 which are Lewis acid sites which is easily to adsorb NH3 molecules, and PTND3 also has bands at 3230 and 1470 cm−1 (see Fig. 5f) which are Bronsted acid sites which can attribute to adsorb NH3 too. Thereby, it deduces that Pd NPs and TiO2 NPs could form Lewis acid sites and the interaction between Pd NPs, N and C elements in TiO2 NFs contributes to forming of Bronsted acid sites. So far, the gas-sensing mechanism of PTND3 sensor is investigated.

Initially, as shown in Fig. 7a, d, when a PTND3 sensor is placed in an open atmosphere at RT, oxygen molecules capture electrons from the surface of TiO2 NPs of nanofibers forming oxygen ions (O2−), and the Pd NPs catalytically activate oxygen molecules into oxygen ions (O2−) (electronic sensitization mechanism), subsequently oxygen ions (O2−) diffuse into the TiO2 NPs of nanofibers around Pd NPs and form an activity region. This process greatly increases the quantity of oxygen ions (O2−) compared with pristine TiO2 NPs. Meanwhile, the interaction of Pd NPs, C and N elements in TiO2 NPs of nanofibers could form Lewis acid sites and Bronsted acid sites in the activity region around Pd NPs.

On exposure to NH3 at RT (Fig. 7b,e), NH3 molecules can be easily adsorbed on Lewis acidic sites and Bronsted acidic sites of the activity region on PTND3. Under the impact of chemical sensitization mechanism of
Pd NPs, oxygen ions ($O_2^-$) can promptly interact with NH$_3$ and readily release more electrons back to conduction band of TiO$_2$ NPs of nanofibers (see reactions 2), which leading to an decrease in the carrier concentration of p-type TiO$_2$ NPs active layer. Hence the resistance of the PTND3 sensor swiftly increases.

Equations (1–2) are the corresponding reactions$^{64,65}$.

$$O_2(TiO_2\text{adsorbed}) + e^- \rightarrow O_2^-$$  \hspace{1cm} (1)

$$4NH_3(\text{ads}) + 3O_2^- \rightarrow 2N_2 + 6H_2O + 6e^-$$ \hspace{1cm} (2)

**Conclusion**

The present study has demonstrated a one-step method to fabricate ordered porous Pd@TiO$_2$ NFs array film. In particularly, PTND3 ordered nanofibers show a high sensing response and fast response time towards different concentration NH$_3$. Most importantly, the PTND3 ordered nanofibers sensor present superior stability to 100 ppm after used 35 days.

The enhanced gas sensing could be ascribed to two factors (i) The electronic sensitization mechanism and chemical sensitization mechanism of Pd NPs on the TiO$_2$ nanofibers, which could be more favorable to produce oxygen ions and accelerated the molecule-ion conversion rate, enhanced the reaction activity of oxygen ions ($O_2^-$) with NH$_3$ compared with pristine TiO$_2$ NFs; (ii) The synergistic effect between Pd NPs, C and N element in TiO$_2$ NFs could form Lewis acid sites and Brønsted acid sites which could enhance NH$_3$ adsorption-desorption ability and activity, and thus favour the rapid-response between NH$_3$ and oxygen ions. Therefore, the Pd@TiO$_2$...
NFs sensor exhibited excellent gas-sensing property at RT. Pd@TiO₂ NFs array film would be effectively used in commercial gas-sensing and environmental monitoring applications.

Experimental Techniques

Sample preparation. All chemicals used were of analytical grade without further purification. The Pd@TiO₂ NFs were synthesized by calcination and reduced using NaBH₄ aqueous solution. In a typical process, 7.2 g poly (N-vinylpyrrolidone) (M₉₀ = 58,000) was dissolved in 15 mL ethanol, then 1 mL 0.005 mol/L H₂PdCl₄ solution and a certain amount of tetra butyl titanate (TBT, C₆H₁₃O₂Ti, M₉₀ = 340.32) was added to obtain a transparent precursor solution of different quality percentage Pd. The quality percentage of Pd is 3.0, 2.5, 2.0 and 1.5 wt%, respectively. The mixed solution was stirred vigorously at RT for 5 h and then transferred into a 10 mL syringe with a capillary tip (0.8 mm in diameter) for spinning. A high voltage of 18 kV was applied at the spinneret as a direct-current power supply (BGG, DFS-01, Beijing Kaiweixin Technique Co., China), and the nanofibers were collected using the aluminium foil at RT. Then the nanofibers blanked were taken off the aluminium foil, and treated at 600 °C for keeping 4 h under the mixed atmosphere of N₂ and air (in a proportion of 1 to 1) with a heating rate of 3 °C min⁻¹. The nanofibers array film was obtained. Then, 0.05 g nanofibers array film was added into 20 mL 0.01 mol/L NaBH₄ aqueous solution at one time. The color of solution changed from yellow to dark-brown gradually. The reduction reaction lasted for 10 h under impregnation. Subsequently, the sample was filtered with washing by distilled water, and dried at 60 °C for 10 h. Pd@TiO₂ NFs with 3.0, 2.5, 2.0 and 1.5 wt% of Pd were obtained and marked as PTND1, PTND2, PTND3, and PTND4, respectively. Here, pPTND3 was the sample which PTND3 was calcined under air atmosphere. In addition, pure TiO₂ was the sample which Pd@TiO₂ NFs sensor was obtained and marked as PTND3 to PTND4, respectively. Here, pPTND3 was the sample which PTND3 was calcined under air atmosphere. Moreover, in order to compare the effects after the different samples adsorbed or desorbed NH₃, aPTND3 adsorbed 100 ppm NH₃ at RT for 30 min, and dPTND3 was the sample which PTND3 adsorbed 100 ppm NH₃ at RT for 30 min, and then PTND3 was treated with He of 30 mL · min⁻¹. Then, the gas response was defined as the ratio |Rₐ − R₅| / R₅, where Rₐ and R₅ are thin film resistance, measured in air atmosphere and NH₃ gas, respectively. The response time is defined as the time taken for the sensor to attain 90% of the maximum change in resistance on exposure to the target gas.

Characterization. The crystal phase of samples was characterized by X-ray powder diffraction (XRD, D/ MAX-III-B–40 KV, Japan, Cu-Kα radiation, λ = 1.5406 Å). The morphology and structures of the nanofibers were observed by scanning electron microscope (SEM, HITACHI S-4800) with an acceleration voltage of 200 kV. The Brunauer-Emmett-Teller (BET) surface area of the products was measured by N₂ adsorption-desorption (TriStar II 3020). The sample was dried for 10 h at 150 °C under vacuum before the measurement. X-Ray photoelectron spectra (XPS) analysis was performed with a VGESCALAB MK II using MgKα (1253.6 eV) achromatic X-ray radiation under high vacuum of 5 × 10⁻⁹ Pa. Infrared spectra (IR) were recorded on a Nicolet Antaris Fourier transform spectrometer between 400 and 4000 cm⁻¹. Spectra were obtained after 20 accumulations with a resolution of 4 cm⁻¹. Thermogravimetric (TG) curves were obtained with a Shimadzu TGA-50 using a platinum crucible with a sample mass of 10 mg. The heating rate was 10 °C min⁻¹ between 25 and 1200 °C, in air atmosphere or mixture atmosphere of air and N₂ (50 mL · min⁻¹). Temperature-programmed desorption (TPD) experiments were carried out with a TP-5000 multi-purpose adsorption instrument using Chemisorp 2705 unit (Micromeritics Instrument. Co., USA) equipped with thermal conductivity detector (TCD) to detect NH₃ adsorption and desorption behavior.

Gas sensing test. The fabrication and measurements of gas sensors were made from the as-synthesized composites by the method we reported before. ²⁹ An alumina substrate (7 mm × 5 mm × 0.38 mm, content 99.6%) with two interdigitated Au electrodes on its top surface was used. Each Au electrode contained 50 fingers, and the distance between two fingers was 20 µm. The Au electrodes on Al₂O₃ substrate was cleaned by diluted HCl, distilled water and acetone. Nanofibers array film was placed on the Au electrodes under the microscope, then calcined at 350 °C for 1 h in N₂ and air (or air) to obtain nanofibers array film sensor.

The gas sensor was installed inside a test chamber with an inlet and an outlet in. The electrical resistance measurements of the sensor was carried out at RT (26 °C) and the relative humidity (RH) around 30%. The gas concentration was controlled by injecting a volume of target gas into the chamber in a certain amount of time with a small fan, and then the chamber was purged with air to revert to the original baseline. The resistance change of the thin film sensor has been measured to confirm the interactions of the sensor and the NH₃ gas. The gas response was defined as the ratio |R₂ − R₅| / R₅, where R₂ and R₅ are thin film resistance, measured in air atmosphere and NH₃ gas, respectively. The response time is defined as the time taken for the sensor to attain 90% of the maximum change in resistance on exposure to the target gas.

TPD analysis. Typically, the sample of 50 mg loaded in the quartz tube was firstly pretreated with He of 30 mL · min⁻¹ at 250 °C for 0.5 h, and subsequently the sample was saturated with NH₃ (5% in He) in a flow rate of 30 mL · min⁻¹ for about 30 min at 100 °C. After that, the physically adsorbed NH₃ was purged by high purity He at the same temperature. The chemisorbed NH₃ was measured by heating the sample in He (30 mL · min⁻¹ from 100 °C to 600 °C with a heating rate of 10 °C · min⁻¹).

Data availability statement. The authors declare that the data in the manuscript are true and reliable.

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Author Contributions
Hongyuan Wu, Haitao Huang and Keying Shi conceived the experiments, Hongyuan Wu and Haitao Huang conducted the experiment, Hongyuan Wu and Jiao Zhou fabricated samples and characterized gas sensing properties, Hongyuan Wu, Haitao Huang, Li Li and Keying Shi analysed data and wrote the manuscript. Dahai Hong, Muhammad Ikram and Afrasiab Ur Rehman deposited and characterized samples. Muhammad Ikram and Afrasiab Ur Rehman reviewed the paper.

Additional Information
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