

# Ammonia Detection at Low Temperature by Tungsten Oxide Nanowires <sup>†</sup>

Sandrine Bernardini <sup>1,\*</sup>, Florent Pourcin <sup>2</sup>, Nassirou Nambiema <sup>1</sup>, Olivier Margeat <sup>2</sup>, Khalifa Aguir <sup>1</sup>, Christine Videlot-Ackermann <sup>2</sup>, Jörg Ackermann <sup>2</sup> and Marc Bendahan <sup>1</sup>

<sup>1</sup> Aix Marseille Univ, Université de Toulon, CNRS, IM2NP, Marseille, France; nambiema.nassirou@gmail.com (N.N.); khalifa.aguir@im2np.fr (K.A.); marc.bendahan@im2np.fr (M.B.)

<sup>2</sup> Aix Marseille Univ, CNRS, CINaM, Marseille, France; pourcin@cinam.univ-mrs.fr (F.P.); olivier.margeat@univ-amu.fr (O.M.); videlot@cinam.univ-mrs.fr (C.V.-A.); ackermann@cinam.univ-mrs.fr (J.A.)

\* Correspondence: sandrine.bernardini@im2np.fr; Tel.: +33-491-288-971

<sup>†</sup> Presented at the Eurosensors 2018 Conference, Graz, Austria, 9–12 September 2018.

Published: 10 December 2018

**Abstract:** Ammonia detection at low temperatures below 150 °C is attractive to be well suited for flexible substrates in terms of thermal strain and to specific environment not allowing high temperature such as explosive one. In commercial gas sensors, tungsten trioxide is the mostly used semiconducting metal oxide after tin dioxide. We report herein the efficiency of tungsten trioxide nanowires deposited on rigid substrate by drop coating from colloidal solution. This study provides an interesting approach to fabricate ammonia sensors on conformable substrate with significant properties for applications in environmental monitoring devices.

**Keywords:** ammonia; gas sensor; tungsten oxide nanowires; environmental monitoring

## 1. Introduction

Poor air quality is a global concern. Around us, ammonia (NH<sub>3</sub>) is a natural toxic and flammable gas present throughout the atmosphere. It is used in manufacturing of fertilizers, explosives, cleaning fluids and in pharmaceutical applications. It is a colorless gas with a characteristic pungent smell and it can affect environment and human health. The National Institute for Occupational Safety and Health (NIOSH) Recommended Exposure Limit (REL) for ammonia is 25 ppm (18 mg/m<sup>3</sup>) averaged over an eight-hour workday and a Short Term Exposure Limit (STEL) of 35 ppm (27 mg/m<sup>3</sup>) during any 15 min period in the day. Due to these reasons, ammonia monitoring has received considerable attention. Numerous methods have been developed and are in use for ammonia measurement [1]. Since 1962, it has been demonstrated that absorption or desorption of a gas on a metal oxide surface changes the material conductivity [2]. Then, Metal Oxide Semiconductors (MOS) have been applied increasingly as conventional chemical sensing materials because of their simplicity in measurement setup, their miniaturization facility for portable instruments, their high sensitivity and their relatively low cost [3]. Conductometric gas sensor based on WO<sub>3</sub> sensing material is demonstrated to respond to NH<sub>3</sub> for gas leak detection indoor and outdoor air quality monitoring [4]. Nevertheless, high working temperatures (275–375 °C) and recovery behaviors limit the widespread use of this material. However, when the active layers have a

large surface area, it means a high surface/volume ratio, the surface adsorbs as much of the target gas as possible and gives a stronger and easily measurable electrical signal [5]. This work aims to investigate the gas-sensing performances of tungsten trioxide ( $\text{WO}_3$ ) nanowires (NWs) at low temperature.

## 2. Materials and Methods

Our conductometric gas sensor consists of Ti/Pt interdigitated electrodes (100 nm thick and 50  $\mu\text{m}$  width, respectively), deposited on Si/SiO<sub>2</sub> by magnetron radiofrequency sputtering and a sensitive  $\text{WO}_{3-x}$  based NWs on the top of them. A heating source can be provided underneath if needed inside the chamber. Ultrathin  $\text{WO}_{3-x}$  NWs were synthesized by a solvothermal method using slight modifications from our previous work [6]. We will briefly remind here. From 110 mg of  $\text{Na}_2\text{WO}_4$  in 3 mL DI water, and 3 mL of 7%  $\text{HNO}_3$  aqueous solution, a  $\text{H}_2\text{WO}_4$  deposition was formed instantaneously. Then, 16 mL of oleylamine was added to this  $\text{H}_2\text{WO}_4$  deposition. To get clear solutions, the mixture was sonicated at room temperature for 20 min, transferred into a 45 mL autoclave, and purged with argon for 15 min. Afterward, the autoclave was put into an electronic oven at 220 °C during 12 h. The product was then deposited by adding 40 mL of ethanol and washed twice with absolute ethanol. Then, 10mg/mL of the resulting  $\text{WO}_{3-x}$  NWs were dispersed in toluene. Finally, this solution with the nanomaterials was deposited by drop-casting 10  $\mu\text{L}$  of the solutions (about 1 mg/mL) onto the electrodes. For this study, UV/O<sub>3</sub> treatment during 15 min at 80 °C have been done before and after  $\text{WO}_{3-x}$  NWs deposition to remove hydrocarbon contamination on the surface which became more hydrophilic with higher surface tension [7]. To be able to test the sensors up to 130 °C a heating was performed at 150 °C during 30 min on some samples.

Gas sensing investigates was performed at various temperatures under obscurity. A 1 V DC voltage was applied to the samples while the electrical resistance was monitored using a Keithley Model 2450 Source Meter (Tektronix, USA, Source Measure Unit). As the main purpose was first to check the  $\text{WO}_{3-x}$  NWs gas-sensing properties using  $\text{NH}_3$  as target gas at low temperature, humidity was avoided using dry air as carrier gas. Because  $\text{WO}_3$  is n-type semiconductor and  $\text{NH}_3$  a reduction gas, the sensor response is defined as the following equation:

$$R = R_{\text{dry air}} / R_{\text{NH}_3} \quad (1)$$

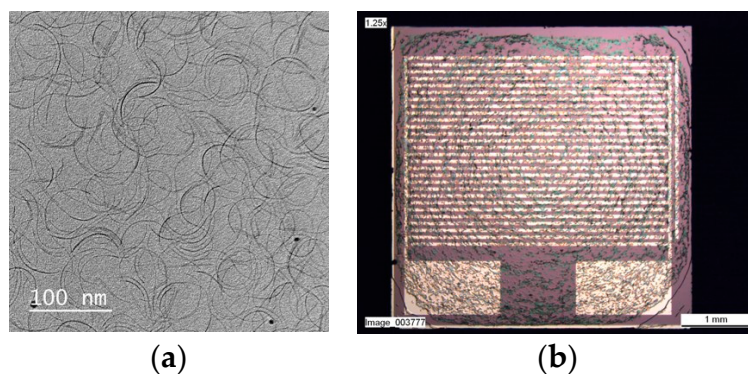
where  $R_{\text{dry air}}$  is the resistance under dry air and  $R_{\text{NH}_3}$  is the resistance under  $\text{NH}_3$  exposure.

The measurements have been performed with dry air as both the reference and the carrier gas, maintaining a constant total flow of 500 Standard Cubic Centimeters per Minute (SCCM) via mass flow controllers. The exposition time was fixed at 30 s.

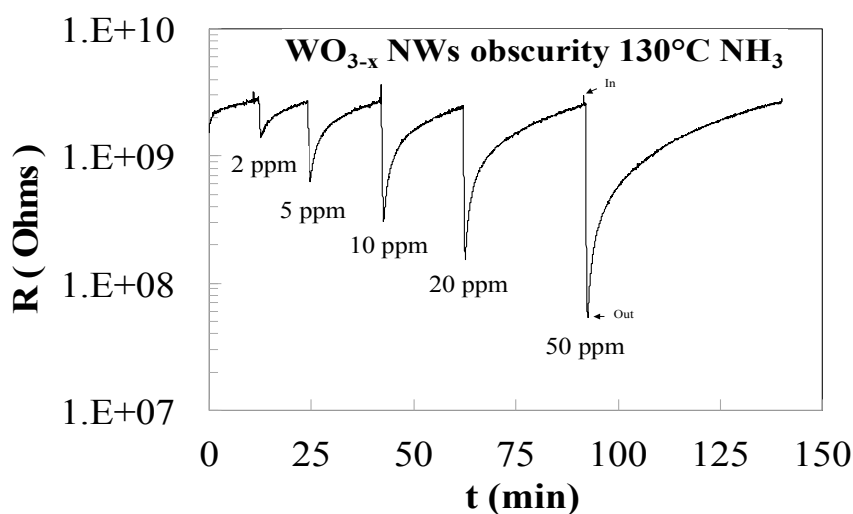
## 3. Results and Discussion

The  $\text{WO}_{3-x}$  NWs drop casted from a diluted solution on a mesh-coated carbon film presented a homogeneous length (about 100 nm) and shape dispersions (Figure 1a) observed by High-Resolution Transmission Electron Microscope (HR-TEM) JEOL-JEM 3010 (USA). Figure 1b presents the  $\text{WO}_{3-x}$  NWs homogeneous layer formed on Si-SiO<sub>2</sub> substrate. No response was measurable at 25 °C and 50 °C due to the high material resistivity and the high signal/noise ratio. Sensor responses were observed from 75 °C up to 130 °C since post-heating process was carry on at 150 °C.

Figure 2 shows the representative dynamic gas response of  $\text{WO}_{3-x}$  NWs to  $\text{NH}_3$  exposure with the concentrations of 2, 5, 10, 20 and 50 ppm at a working temperature equals to 130 °C. The resistance decrease induced by  $\text{NH}_3$  exposures on Figure 2 is in agreement with the behavior of a reduction gas on an n-type semiconductor. The mean idea is that the conductivity is controlled by oxygen vacancies which are generated and destroyed at the gaseous interface by reaction with oxygen.



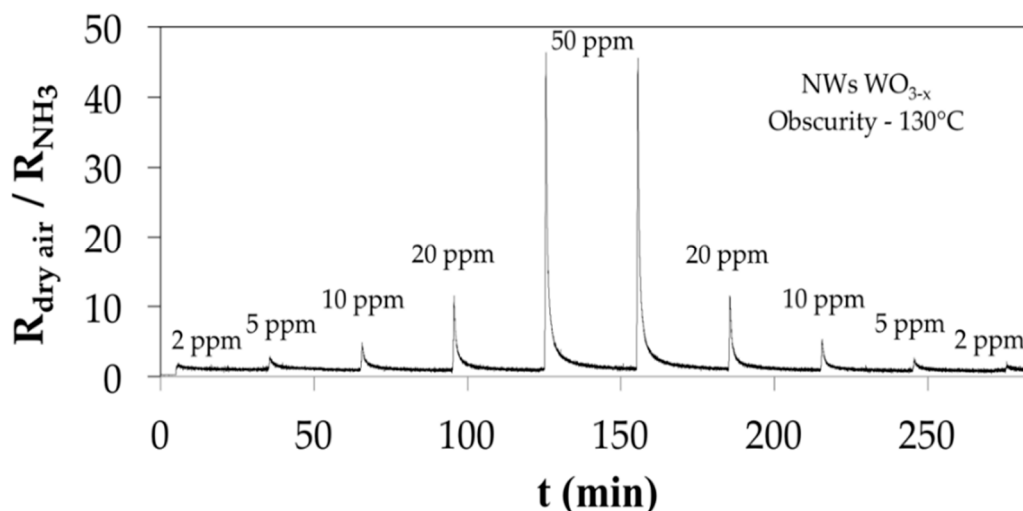
**Figure 1.** (a) TEM image of  $\text{WO}_{3-x}$  NWs and (b) Si-SiO<sub>2</sub> substrate picture with  $\text{WO}_{3-x}$  NWs dropped on it.



**Figure 2.** Resistance variation of  $\text{WO}_{3-x}$  NWs with  $\text{NH}_3$  exposure with the concentrations of 2, 5, 10, 20 and 50 ppm at a working temperature equals to 130 °C.

Before  $\text{NH}_3$  exposure, oxygen ( $\text{O}_2$ ) in the air will adsorb and react with  $\text{WO}_3$  on its interface. The charge carrier concentration and therefore the conductivity change in response to the oxygen vacancy concentration variations at the interface. In this process,  $\text{O}^{2-}$  will capture electrons from  $\text{WO}_3$  and transform into negatively-charged species ( $\text{O}^{\sigma-} = 1/2, 1 \text{ or } 2$ ). During  $\text{NH}_3$  exposure, it will react with  $\text{O}^{\sigma-}$  to produce nitrogen or nitrogen oxides and electrons will return to  $\text{WO}_3$  [8]. The reactions at the interface develop a charge at the interface, which is balanced by a space charge of electrons within the oxide nanowire, near the interface. The surface-to-volume ratio of the nanowires will also increase the conductivity change and sensor sensitivity. The sensor response as a function of the  $\text{NH}_3$  concentration was plotted in Figure 3.

The response shows a nearly linear increase with the  $\text{NH}_3$  concentration rise. As illustrated in Figure 3, it was found a fast response less than 20 s and a time to return to the baseline around 5 min for an ammonia concentration of 2 ppm at 130 °C. This return time at this low temperature is interesting for using it several times in a short period. According to the sensor response curve presented in Figure 3, the  $\text{WO}_{3-x}$  nanowires-based sensing device exhibits good reversible response at 130 °C without need of light excitation. The repeatability tests, by increasing and decreasing  $\text{NH}_3$  concentration exposures show a small deviation, an important response, a good reversibility, fast response and recovery without sensor saturation from 2 ppm to 50 ppm  $\text{NH}_3$  gas.



**Figure 3.** Sensor response curve of  $\text{WO}_{3-x}$  NWs based sensing device post-annealed at 150 °C and working at 130 °C in obscurity.

#### 4. Conclusions

This study presents the efficiency of tungsten trioxide nanowires  $\text{WO}_{3-x}$  deposited on rigid substrate by drop coating from colloidal solution to detect ammonia below the Recommended Exposure Limit Values (25 ppm) as low as 2 ppm at 130 °C without need of light excitation. This working temperature is compatible with thermal strain of flexible substrate. It is an attractive way to fabricate ammonia sensors on flexible substrate with interesting properties in terms of sensitivity and response times for applications in environmental monitoring devices. Further studies will be carry on selectivity versus other environmental gases and light illumination impact.

#### References

1. Timmer, B.; Olthuis, W.; Van den Berg, A. Ammonia sensors and their applications—A review. *Sens. Actuators B* **2005**, *107*, 666–677, doi:10.1016/j.snb.2004.11.054.
2. Seiyama, T.; Kato, A.; Fujiishi, K.; Nagatani, M. A new detector for gaseous components using semiconductive thin films. *Anal. Chem.* **1962**, *34*, 1502–1503, doi:10.1021/ac60191a001.
3. Fine, G. F.; Cavanagh, L.M.; Afonja, A.; Russell, B. Metal oxide semi-conductor gas sensors in environmental monitoring. *Sensors* **2010**, *10*, 5469–5502, doi:10.3390/s100605469.
4. Wang, X.; Miura, N.; Yamazoe, N. Study of  $\text{WO}_3$ -based sensing material for  $\text{NH}_3$  and NO detection. *Sens. Actuators B* **2000**, *66*, 74–76, doi:10.1016/S0925-4005(99)00410-4.
5. Wei, W.; Li, W.; Wang, L. High-selective sensitive  $\text{NH}_3$  gas sensor: A density functional theory study. *Sens. Actuators B* **2018**, 502–507, doi:10.1016/j.snb.2018.02.108.
6. Liu, J.; Margeat, O.; Dachraoui, W.; Liu, X.; Fahlman, M.; Ackermann, J. Gram-Scale, Synthesis of Ultrathin Tungsten Oxide Nanowires and their Aspect Ratio-Dependent Photocatalytic Activity. *Adv. Funct. Mater.* **2014**, *24*, 6029–6037, doi:10.1002/adfm.201401261.
7. Luo, S.; Wong, C.P. Effect of UV/ozone treatment on surface tension and adhesion in electronic packaging," in IEEE Transactions on Components and Packaging Technologies, **2001**, *24*, 1, 43-49, doi:10.1109/6144.910801.
8. Jiménez, I.; Centeno, M.A.; Scotti, R.; Morazzoni, F.; Arbiol, J.  $\text{NH}_3$  interaction with chromium-doped  $\text{WO}_3$  nanocrystalline powders for gas sensing applications. *J. Mater. Chem.* **2004**, *14*, 2412–2420, doi:10.1039/b400872c.

