





# Proceedings 2D SnS<sub>2</sub>—A Material for Impedance-Based Low Temperature NO<sub>x</sub> Sensing? <sup>+</sup>

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**Abstract:** The sensor signal of tin disulfide (SnS<sub>2</sub>), a two-dimensional (2D) group-IV dichalcogenide, deposited as a film on a conductometric transducer is investigated at 130 °C. The focus is on the detection of the total NO<sub>x</sub> concentration. Therefore, the sensor response to NO and NO<sub>2</sub> at ppm- and sub-ppm level at low operating temperature is determined. The results show that the sensing device provides a high sensor signal to NO and NO<sub>2</sub> even at concentrations of only 390 ppb NO<sub>x</sub>. Both nitrous components, NO and NO<sub>2</sub>, yield the same signal, which offers the opportunity to sense the total concentration of NO<sub>x</sub>.

**Keywords:** 2-dimensional SnS<sub>2</sub>; conductometric sensor; total NO<sub>x</sub> sensing; sub-ppm NO<sub>x</sub>; low temperature

### 1. Introduction

NO<sub>x</sub> sensing at low temperatures is still a difficult task, especially for air-quality monitoring in stationary or transportable air quality monitoring devices [1]. In the past years, strict emission and immission limits for NO<sub>x</sub> have been set up, for instance by the EU immission legislation Directive 2008. Currently, the emissions of NO<sub>x</sub> by traffic in urban regions are a widely discussed topic since they exceed the regulatory limits. To enforce the limits, the detection of low NO<sub>x</sub> concentrations at ppm- and sub-ppm level is required. Reliable and long-term stable sensing devices for the lowest NO<sub>x</sub> concentrations are necessary to meet the strict requirements of, for instance, the European legislation, with regard to quality of the data especially in real-time air quality monitoring [2]. In literature, various NO<sub>x</sub> gas sensing technologies are discussed [1,3–5]. The sensors have to be accurate, selective, long-term stable, and should have low NO<sub>x</sub> detection limits. Especially for air-quality monitoring, the sensors for low-temperature NO<sub>x</sub> sensing with a low power consumption are beneficial [1,2].

In this work a new material class based on 2D transition metal dichalcogenides (TMD) is discussed as sensing materials for NO<sub>x</sub> sensors [6,7]. Due to the special structure of SnS<sub>2</sub>, the charge transfer between *physisorbed* NO<sub>2</sub> gas molecules and the 2D SnS<sub>2</sub> material allows for NO<sub>2</sub> sensing with high NO<sub>2</sub> sensitivity and selectivity at low operating temperatures. In [6], the NO<sub>2</sub> sensor response of SnS<sub>2</sub> is described at 120 °C. In the present work, tin disulfide (SnS<sub>2</sub>) flakes are investigated as functional materials for NO<sub>x</sub> sensing at 130 °C with focus on the NO sensing performance of SnS<sub>2</sub>.

#### 2. Materials and Methods

The transducer composed of an alumina substrate with a screen-printed interdigitated-electrode (IDE) structure (Au-IDE: 100  $\mu$ m/100  $\mu$ m). The sensitive film of 2D SnS<sub>2</sub> was synthesized by a wet chemical route and drop-casted on the IDE-structure [6]. The structure and morphology of the SnS<sub>2</sub> film was analyzed by scanning electron microscopy (SEM) and transmission electron microscopy (TEM).

The sensor response, the complex impedance |Z| of the SnS<sub>2</sub>-film, was determined at 130 °C in a synthetic gas test bench. As base gas, synthetic air with 2 vol.% water was used and NO and NO<sub>2</sub> were added in a concentration range between 390 ppb and 2 ppm. The added NO<sub>x</sub> concentration was analyzed downstream the sensing device by a chemiluminescence detector (CLD).

#### 3. Results and Discussion

The microstructure of the SnS<sub>2</sub> film is shown in Figure 1. The SEM image (Figure 1a) of the deposited film shows SnS<sub>2</sub> particles with a hexagonal shape which appear to from flake like structures. This is proven by the TEM image (Figure 1b). The shape of the particle is a hexagonal plate, with an average diameter of 100 nm and a thickness of less than 10 nm. This planar 2-dimensional flake structure was selected due to its very high active surface area resulting in a high adsorption capability for physisorbed NO<sub>x</sub> molecules.

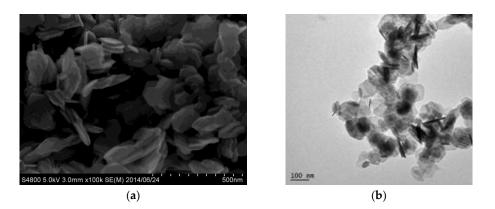
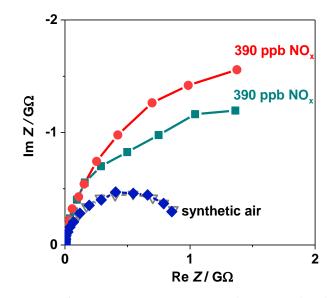


Figure 1. (a) SEM image of SnS<sub>2</sub> particles; (b) TEM image of SnS<sub>2</sub> particles.

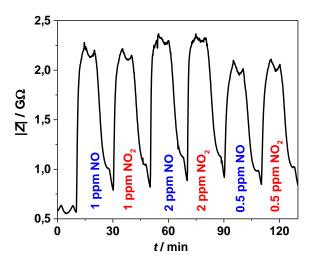
Initial impedance spectra of a SnS<sub>2</sub> sensor, shown in Figure 2 in the form of Nyquist-plots (frequency between 1 MHz and 1 Hz, root-mean-square value of the amplitude 200 mV, temperature 130 °C), present a semi-circular behavior with a high sensor signal when exposed to 390 ppb NO<sub>x</sub>. The sensor signal in synthetic air is very stable (shown are two measurement curves). The complex impedance increases strongly in presence of NO<sub>x</sub>, even at a NO<sub>x</sub> concentration in the sub-ppm range.

As stated in [6] for NO<sub>2</sub> exposure, the strong resistance increase can be explained by the effect that the adsorbed NO<sub>2</sub> gas molecules act as electron acceptors. Charge is transferred from the  $SnS_2$  flakes to the adsorbed NO<sub>2</sub> and the  $SnS_2$  flakes deplete with charge carriers. The reduced number of free electrons leads to the increasing resistance.

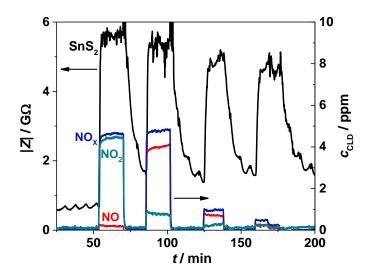
For further measurements, we selected a constant frequency of 1 Hz. The sensor was exposed to varying NO and NO<sub>2</sub> concentration steps, and the resulting |Z| is presented in Figures 3 and 4. The impedance of the SnS<sub>2</sub> sensor is around 600 M $\Omega$  in synthetic air, and increases when exposed to 1 ppm NO or NO<sub>2</sub> to 2.2 G $\Omega$  with the same sensor signal for NO and NO<sub>2</sub>. The oscillating of the sensor signal is due to temperature fluctuations of the furnace (around 10 °C). The response time is quite good, but the signal recovery is relatively slow. The sensor seems to be a total NO<sub>x</sub> sensing device.



**Figure 2.** Impedance spectra of a SnS<sub>2</sub> sensor at 130 °C in synthetic air and with 390 ppb NO<sub>x</sub> in the Nyquist-plot representation; spectra determined with  $U_{eff}$  = 200 mV and between 1 Hz and 10 MHz.



**Figure 3.** Complex impedance |Z| signal of the SnS<sub>2</sub> sensor determined at f = 1 Hz at 130 °C during NO<sub>x</sub> exposure to 1 ppm, 2 ppm and 0.5 ppm NO res. NO<sub>2</sub>.



**Figure 4.** Complex impedance |Z| of the SnS<sub>2</sub> sensor determined at f = 1 Hz at 130 °C during NO<sub>x</sub> exposure and the added NO<sub>x</sub> concentration analyzed by CLD downstream the sensor device.

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To investigate this more in detail, Figure 4 includes the NO and NO<sub>2</sub> concentrations determined by a CLD gas analyzer. Comparing the first two NO<sub>x</sub> peaks, almost the same sensor response is visible for 5 ppm total NO<sub>x</sub> (5 ppm NO<sub>2</sub> res. 4 ppm NO with 1 ppm NO<sub>2</sub>). A huge sensor signal can determined even for NO<sub>x</sub> concentrations below 1 ppm.

# 4. Conclusions

The  $SnS_2$  sensors show a huge  $NO_x$  gas response even for low concentrations that needs to be investigated with respect to the behavior as a total  $NO_x$  sensor. The developed sensing device provides high impedance values. The impedance changes strongly when exposed to low concentrations of NO or  $NO_2$  and the sensor seems to be suitable for sub-ppm level  $NO_x$  detection. The dependence of the resistance on the thickness of the  $SnS_2$  film is an interesting task for further investigations. Additionally, the concentration dependent sensor response has to be analyzed and the response and the recovery time need to be improved.

Conflicts of Interest: The authors declare no conflict of interest.

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