

Abstract

Pulsed Temperature Operation of SnO₂-Based Gas Sensors [†]

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[†] Presented at the XXXV EUROSENSORS Conference, Lecce, Italy, 10–13 September 2023.

Abstract: We herein demonstrate the pulsed-mode temperature operation of chemical sensor devices based on thin SnO₂ films, which were synthesized by magnetron sputtering. The gas-sensitive films were integrated on SiN-based micro-hotplate (μ hp) chips, which enable operation temperatures up to 500 °C. We compared the gas sensor performance in constant temperature mode with pulsed temperature mode operation towards the test gases carbon monoxide and toluene. In contrast to constant temperature, the pulsed temperature mode operation reveals additional information about the type of test gas.

Keywords: chemical sensors; metal oxides; nanoparticles; temperature modulation

1. Introduction

Nowadays, conductometric chemical sensors based on metal oxides like SnO₂, ZnO, CuO_x, and WO_x are the most promising and investigated types of solid-state sensors [1,2]. Gas sensor devices are of high importance for many applications, where energy-autonomous sensor systems suitable for IoT applications could achieve wide-area pollution monitoring and mapping. Power consumption of such sensor devices, however, is a key performance aspect that is decisive for their application success. Pulsed temperature operation offers much lower power consumption (<1 mW) as compared to conventional DC-mode temperature operation (>30 mW). Moreover, pulsed temperature operation reveals target gas-dependent features in the response, which can be exploited to increase the selectivity of the sensor devices.

2. Methods and Materials

SiN-based μ hp chips, which provide a heating structure for operating temperatures up to 500 °C, and electrodes for contacting the SnO₂ films were processed by photolithography with a negative resist mask. Next, the SnO₂ sensing layer with a thickness of 50 nm was deposited by reactive magnetron sputtering of a Sn target in Ar + O₂. Afterwards, the sensors were functionalized with metallic nanoparticles (Ag, Ti, and Cu) synthesized by magnetron sputter inert gas condensation [3] to improve sensitivity and selectivity. The sensor devices were characterized in an automatized setup with synthetic air (80% N₂, 20% O₂, and humidity 50%) as a background gas and a constant flow rate of 1000 sccm. The target gases are carbon monoxide (CO) and toluene (C₇H₈).

3. Discussion

A typical resistance measurement of a sensor device is exemplified in Figure 1a, and the temperature cycling schematics are shown in the lower graph. First, the sensors are heated up to 500 °C for 15 min (DC-mode), the test gas (indicated as a grey column) is



Citation: Egger, L.; Reiner, L.; Togni, A.; Mitterer, C.; Köck, A. Pulsed Temperature Operation of SnO₂-Based Gas Sensors. *Proceedings* **2024**, *97*, 211. <https://doi.org/10.3390/proceedings2024097211>

Academic Editors: Pietro Siciliano and Luca Francioso

Published: 6 May 2024



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inserted for 5 min, and the sensors' resistance decreases. Next, the devices are cooled down to room temperature for 10 min; this is followed by three subsequent heating pulses (pulsed mode—1 min), where only the second heating pulse is applied in presence of the test gases. This procedure is executed three times for 5, 10, and 20 ppm test gas concentration. Figure 1b shows in detail the entirely different resistance gradient behavior (emphasized by the blue circle) in pulsed mode operation for 5 ppm CO and C₇H₈.

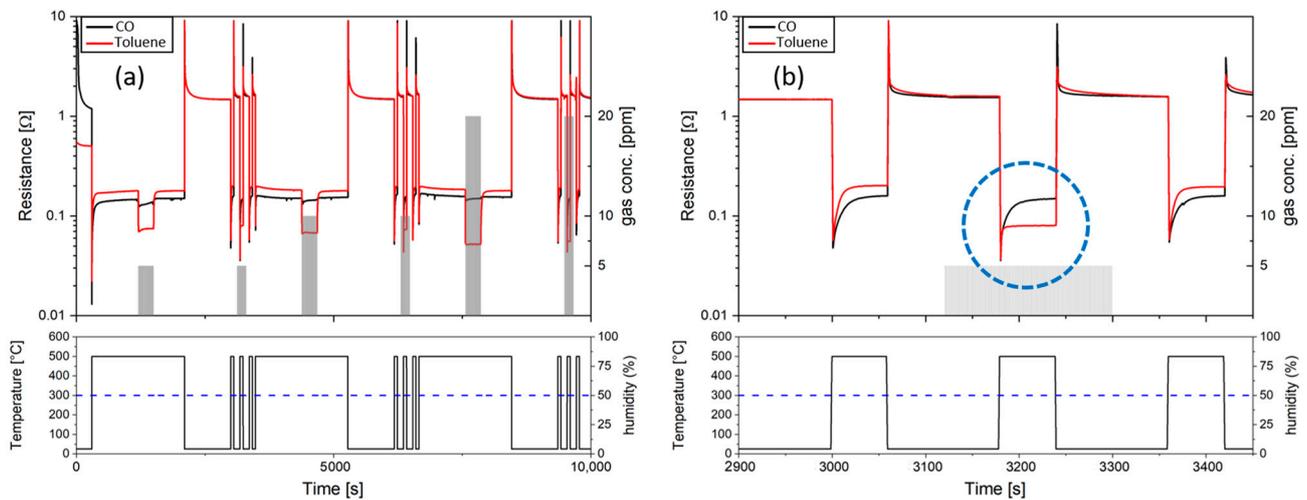


Figure 1. (a) Typical resistance measurement in DC- and pulsed-mode T operation for CO and C₇H₈ (5, 10, and then 20 ppm). (b) Comparison of pulsed-mode T operation for 5 ppm of CO and C₇H₈.

The sensor response (= resistance decrease) in presence of the test gases is comparable to the DC-mode. However, it is obvious that the shape of the response curves (see blue circle in Figure 1b) is entirely different for CO and C₇H₈. Thus, we conclude that pulsed T-mode operation reveals particular information about the test gas that cannot be derived from DC-mode operation.

Author Contributions: Conceptualization, L.E. and A.K.; methodology, A.T. and C.M.; validation, L.E. and L.R.; formal analysis, L.E. and L.R.; investigation, L.E.; writing—original draft preparation, L.E. and A.K.; writing—review and editing, L.E., A.K., A.T. and C.M.; supervision, A.K. and C.M. All authors have read and agreed to the published version of the manuscript.

Funding: This project has received funding from the ECSEL Joint Undertaking (JU) under grant agreement No 876362. The JU receives support from the European Union's Horizon 2020 research and innovation program and Austria, Belgium, Czech Republic, Finland, Germany, Italy, Latvia, Netherlands, Poland, Switzerland. The authors gratefully acknowledge the financial support under the scope of the COMET program within the K2 Center "Integrated Computational Material, Process and Product Engineering (IC- MPPE)" (Project No 886385). This program is supported by the Austrian Federal Ministries for Climate Action, Environment, Energy, Mobility, Innovation and Technology (BMK) and for Labour and Economy (BMAW), represented by the Austrian Research Promotion Agency (FFG) and the federal states of Styria, Upper Austria and Tyrol.

Institutional Review Board Statement: Not applicable.

Informed Consent Statement: Not applicable.

Data Availability Statement: No new data were created.

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

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