Response dynamics of metal oxide gas sensors working with temperature profile protocols

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Abstract

In this work we present the analysis of gas sensors working in modulated temperature mode with temperature varying according to exponential law. We integrate conductometric gas sensor based on semiconducting metal oxide layers and an ad-hoc developed electronics to present a sensing system based on a single sensor featuring a degree of selectivity arising from the exploitation of response dynamics features. In particular, a set of parameters is used to summarize the deviation of the response shape from the single exponential law.

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1. Introduction

Measurements protocols using temperature profiles [1] are attractive because of the large amount of information (features) they allow to extract, compared with isothermal protocols. From a chemical-physical point of view, a larger information arises because of the different thermodynamic conditions explored in temperature cycles. At the same time, the full exploitation of these protocols requires the use of control electronics featuring the capability to work with a fast clock and read the sensor conductance through the wide range that would be spanned during temperature cycles. Despite commercial solutions exist, their high cost makes the sensing system (sensor and control/readout electronics) quite expensive,

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in contrast with the philosophy underlying sensor development. To this aim, the authors developed a cheap readout electronics suitable to perform fast measurements (10 ms) over a wide range of values (10 pS - 100 μS), [2]. In a previous work, we used a first prototype of the system to control an array of four metal oxide (MOX) based chemiresistors to measure their static response and exploit the selectivity of the array to discriminate key-aromas developed during bread baking [3]. The electronic system has been further developed to control the sensor temperature in order to develop an autonomous device (sensor and electronics) fulfilling the requirements of reduced cost, fast readings, wide range of measurable conductance. In this work, we use this system to analyze the dynamic response of a WO3-based gas sensor exposed to different concentrations of ethanol and methane in different humidity conditions. The use of temperature protocols allows obtaining different features summarizing the static and dynamic response of the device to the different test-atmospheres.

2. Experimental

An ad-hoc electronic circuit has been designed for the complete and flexible management of resistive chemical sensors, such as metal oxide devices. Fig. 1 (a) shows the front-end used for the estimation of the sensor parameters; in particular the system is capable to estimate the sensor conductance \( G_S \) as well as a parasitic capacitance \( C_S \), modeled in parallel with \( G_S \). It is based on a conductance-to-time conversion (CTC) scheme, which allows a wide range of conductances to be estimated without using scale factors. To obtain a short measuring time (10 ms), an interpolation method, by means of the least means square algorithm, is used whenever the CTC approach requires a longer time to be accomplished [2].

Fig.1 (b) shows the block scheme of the complete system, detailing the stage devoted to the thermal management of the sensor [4]. The sensor heater is driven by a user-programmable voltage \( V_{tot} \), furnished by means of a digital to analog (D/A) converter followed by a current boost. The current \( I_h \) flowing through the sensor heater is estimated by means of the voltage drop \( V_{sh} \) on the shunt resistor \( R_{sh} \), and then acquired by means of an analog to digital (A/D) converter. Thus, the actual heater voltage \( V_h \) can be easily computed and, together with the estimated value of \( I_h \), allows both the heater resistance \( R_H \) and the heater power \( P_H \) to be reckoned. Every measuring cycle (10 ms), a new value of \( V_{tot} \) can be provided and a new estimation of the heater parameters is carried out, in a synchronous way with \( G_S \) and \( C_S \) estimations. Thus, it is possible to apply various thermal profiles and to finely track both the sensor and the heater behavior.

Finally, collected data are sent every cycle via a serial link to a PC, where a human interface application in LabVIEW environment allows the user to set the measurement parameters, to create the desire thermal profile and to save the sensor data for off-line analysis.

![Fig. 1: (a) scheme of the sensor interface; (b) scheme of the complete system, with a detail of the sensor heater management stage.](image-url)
The WO$_3$ layer used as sensing element has been prepared over a 2 mm x 2 mm x 0.25 mm alumina substrate by a modified thermal evaporation method detailed in [5]. Interdigitated electrical contacts spaced by 200 μm were deposited by Sputtering DC over the oxide layer. The device was provided by a Pt meander (deposited by Sputtering DC) acting both as heating element and temperature probe on the substrate back side.

Measurements were carried out by flow through method working with a constant flow of 300 sccm in a thermostatic sealed chamber at room pressure under controlled humidity conditions (RH=0-50% @ 20°C). Controlled gas mixtures were obtained by using mass flow controllers mixing flows from certified bottles.

3. Results and Discussion

The response dynamics of the WO$_3$ sensor working in temperature-modulated mode during the exposure to different atmospheres are compared in Fig. 2 (a).

![Fig. 2](image_url)

The observed dynamics arise both from a direct conductance modulation induced by the temperature profile of the device as well as from the temperature-dependent interactions occurring among the semiconducting WO$_3$ layer and the gaseous environment. In particular, the adopted temperature-protocol consists of a voltage square-wave applied to the heating element. This induces a sequence of exponential rise and decay of the device temperature in the range from 200 to 400 °C. The device temperature is tracked by measuring the heating element resistance ($R_H$, Fig. 2(b)), whose value is proportional to its temperature. The time dependence of the temperature has been found to behave according to a single exponential (time-constant) law, so that, in a vertical log scale (Fig. 2 (d)), the normalized heater
resistance $R_{H}^*$ calculated according to Eq. (1), is linear:

$$R_H = R_{H,f} - \left( R_{H,f} - R_{H,0} \right) e^{-t/\tau}, \quad R_H^*(t) = \frac{R_{H,f} - R_H(t)}{R_{H,f} - R_{H,0}} = e^{-t/\tau}$$

where $0$ and $f$ subscripts stand for initial and final respectively.

Independently of the surrounding atmosphere, time constants of 2.1 s and 2.9 s have been extrapolated for the heating and cooling cycles respectively.

The shape of the sensing layer conductance $G_S$ rise and fall during the heating and cooling cycles is analyzed through a normalized conductance, $G_S^*$, calculated according to Eq. (1). $G_S^*$ allows to analyze the response dynamics according to their deviation from the single exponential law. According to Fig. 2 (c), deviations are observed for all the tested gases and concentrations, but ethanol, with $G_S^*$ reaching its minimum before the end of the heating (cooling) cycle. Ethanol, differently, induces a peculiar response shape quite close to the basic model depicted in Eq (1) for most of the cycle.

The shape of $G_S^*$ can be summarized through a set of parameters such as: the time constant that can be extrapolated from the initial stage of heating/cooling cycles ($\tau_0$), the time and value of the $G_S^*$ minimum ($t_{\text{min}}$ and $G_{S,\text{min}}^*$). In the case of ethanol, $t_{\text{min}}$ coincides with the end of the cycle and $G_{S,\text{min}}^*$ is null (according to the definition provided by Eq (1)).

The approach provides a set of parameters featuring a marked analyte dependence (and a weak concentration dependence) that can be coupled with typical response amplitude parameters (e.g. $\Delta G_S$), which exhibit enhanced concentration dependence, but a reduced analyte dependence. For example, during the sensor exposure to the air background, $\tau_0$ extrapolated during the warming cycle ranges from 20 to 25 s, depending on relative humidity (0-50%), while, in presence of ethanol, the same parameter ranges between 5 and 15 s (whichever the RH value, 0-50%), providing a useful parameter to separate ethanol from the humidity background.

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