

Chemical Stability Of Sr(Ti,Fe)O_{3-δ} Exhaust Gas Sensors

Thomas Schneider¹, Ellen Ivers-Tiffée¹,
Harumi Yokokawa²

¹Institut für Werkstoffe der Elektrotechnik IWE, Universität
Karlsruhe (TH), Adenauerring 20b, 76131 Karlsruhe, Germany

²Fuel Cell Group, Energy Electronics Institute
National Institute of Advanced Industrial Science and
Technology (AIST), AIST Tsukuba Central 5, Tsukuba 305-
8565, Japan

Oxygen sensors for automotive applications control the air-fuel ratio in order to reduce fuel consumption. New control strategies for direct injection engines or lean burn engines operating with air excess ($\lambda > 1$) require more advanced sensor concepts [1].

As a future option, research groups evaluate the features of a temperature independent resistive-type oxygen sensor based on semiconducting metal oxides.

Acceptor doped Sr(Ti,Fe)O_{3-δ} has been qualified as a resistive type oxygen sensor material [2], its electrical conductivity reflects the equilibrium between the oxygen partial pressure p_{O_2} in the atmosphere and the bulk stoichiometry at temperatures typically above 700 °C. In contrary to donor doped compounds, Sr(Ti,Fe)O_{3-δ} shows a long term stable and temperature independent characteristic [3].

The dependence of electrical conductivity on the oxygen partial pressure is well understood on a defect chemistry basis [4].

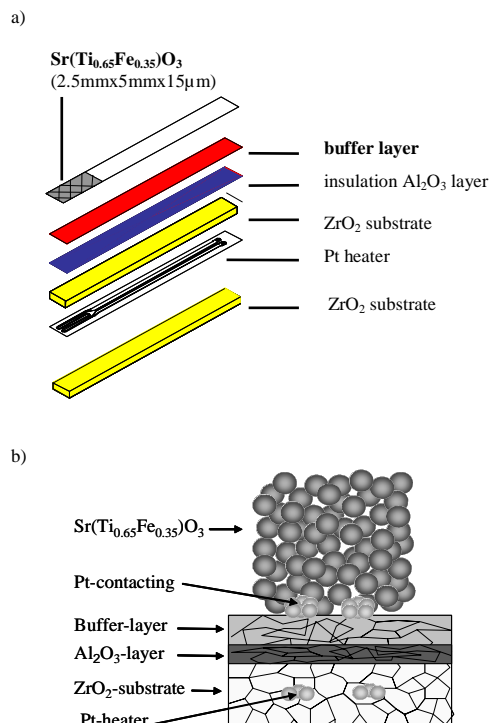


Fig. 1: Design concept of a planar exhaust gas sensor. Sr(Ti,Fe)O_{3-δ}: resistive type oxygen sensor material

For the development of a planar type sensing element, Sr(Ti,Fe)O_{3-δ} has to be applied as a thick film on a zirconia substrate, which has integrated Pt heating elements as well as Pt contacts. Fig. 1 shows the design concept and the multi-layer structure of a planar exhaust gas sensor [5].

Temperature independence (at T=750...900°C, $p_{O_2}=10^{-6}$...1 bar) and fast response times ($t_{90}=6.5$ ms at 900 °C), both key issues of Sr(Ti,Fe)O_{3-δ}, have to be maintained over the entire lifetime of the sensing element. In this work, financially supported by the BMBF (PTJ-NMT 03N3102), the stability of the interface Sr(Ti,Fe)O_{3-δ}/Al₂O₃ was investigated by interdiffusion experiments and by the calculation of chemical potential diagrams (CPD) using the software MALT [6,7,8,9].

CPDs have been calculated for the solid solution end members SrTiO₃ and Sr₂Fe₂O₅, both in contact with the insulating Al₂O₃ layer, for T=750...900°C and p_{O_2} , as shown in figures 2 and 3.

The results support the development of a buffer layer, which inhibits the interdiffusion and the formation of secondary layers at the interface Sr(Ti,Fe)O_{3-δ}/Al₂O₃

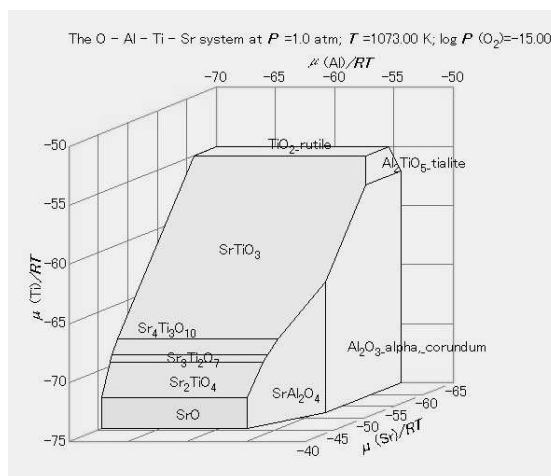


Fig. 2: Chemical potential diagram of Sr-Ti-O-Al system at T=800 °C and $p_{O_2}=10^{-15}$ bar

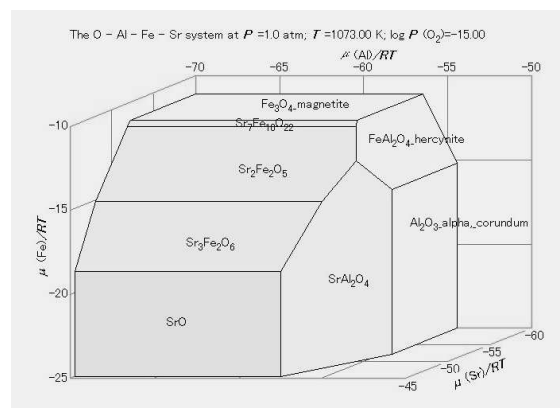


Fig. 3: Chemical potential diagram of Sr-Fe-O-Al system at T=800 °C and $p_{O_2}=10^{-15}$ bar

- [1] E. Ivers-Tiffée et al., *Electrochimica Acta* **47** (2001) 807-814
- [2] R. Moos et al., *Sensors and Actuators B* **67** (2000), 178-183
- [3] W. Menesklou et al., *MRS Symp. Proc.* **604** (2000), 305
- [4] W. Menesklou et al., *Sensors and Actuators B* **59** (1999), 184-189
- [5] J. Riegel et al., *Solid State Ionics* **152-153** (2002) 783-800
- [6] H. Yokokawa et al., *Thermochimica Acta* **245** (1994), 45-55; see <http://www.kagaku.com/malt/index.html>
- [7] H. Yokokawa, *Annu. Rev. Mater. Res.* (2003), 581-610
- [8] H. Yokokawa, *Journal of Phase Equilibria*, Vol. 20 No. 3 (1999)
- [9] H. Yokokawa et al., *Calphad* Vol. 26 No. 2 (2002), 155-166