Electronic Supplementary Material (ESI) for Journal of Materials Chemistry C. This journal is © The Royal Society of Chemistry 2023

Supporting Information

Gas-Sensing Performance of Core-shell SnO₂-based Chemiresistive

MEMS Sensor for H₂S Detection under Vacuum

Wenbo Pi^{a, ‡}, Xi Chen^{a,‡}, Qiuyun Fu^a, Zixiao Lu^b, Honglang Li^b, Zaiqi Tang^c, and Wei Luo*,^{a,d}

^aSchool of Integrated Circuits, Huazhong University of Science and Technology, Wuhan 430074, PR China
^bCAS Center for Excellence in Nanoscience, National Center for Nanoscience and Technology, Beijing 100190, PR China
^cSysmo Technologies Co., LTD, Beijing 100020, PR China
^dResearch Institute of Huazhong University of Science and Technology in Shenzhen, Shenzhen 518000, PR China.
^{*}These authors contributed equally to this work.
*E-mail: luowei@mail.hust.edu.cn Characterization of silver interdigital electrode SnO_2 gas sensor after introducing H_2S gas.

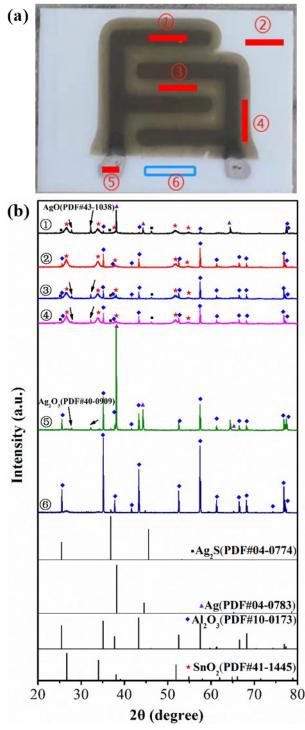


Figure S1. (a) Photo of the silver interdigital electrode SnO_2 gas sensor after introducing H_2S gas at ambient temperature and pressure, (b) Micro-area XRD pattern of the silver interdigital electrode SnO_2 gas sensor exposed to H_2S gas.

When 1 ppm of H_2S gas was introduced to the SnO_2 gas sensor through the silver interdigital electrode, an exponential decrease in the resistance of the SnO_2 gas sensor was observed. It is noteworthy that after the gas sensing reaction, the SnO_2 sensitive

film above and around the silver interdigital electrode of the gas sensor turned brownish-yellow, while the color of the SnO₂ sensitive film away from the silver interdigital electrode remained unchanged, as shown in Figure S1(a). The phase composition of each part of the silver interdigital electrode SnO₂ gas sensor exposed to H₂S was studied using a micro-area X-ray diffractometer, and the XRD spectrum is shown in Figure S1. The results showed that there were small amounts of Ag₂O and Ag₂S around the Ag electrode, which implied that there were two possible processes for this gas-sensitive reaction: one is the direct reaction between Ag and H₂S gas to generate Ag₂S, another is the reaction between the Ag₂O and H₂S. Both gas-sensitive processes are expected to be applied in the detection of H₂S under low-temperature and high-vacuum conditions.

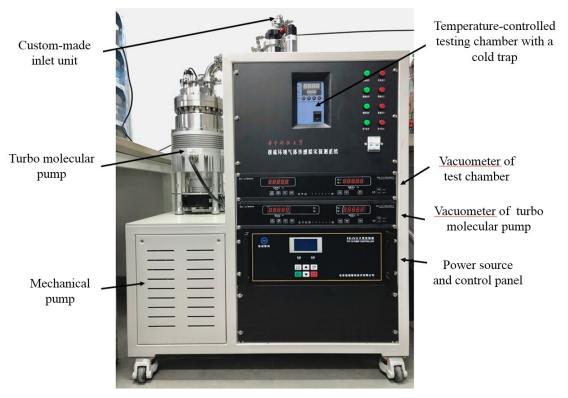


Figure S2. Gas sensor test system under low temperature and high vacuum

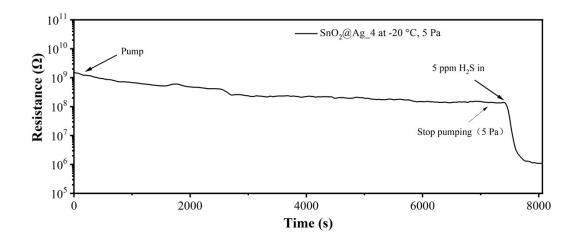


Figure S3. The response curve of SnO₂@4Ag gas sensor to 5 ppm H₂S in a low-temperature and low-vacuum condition (-20 °C, 5 Pa)

Then, $SnO_2@4Ag$ gas sensor was tested for its gas sensing performance towards 5 ppm of H₂S gas in a low-temperature and low-vacuum test system (-20 °C, 5 Pa). The experimental results, as shown in Figure S3, indicated that upon injecting 5 ppm of H₂S gas into the system, the resistance of the $SnO_2@4Ag$ gas sensor immediately decreased, and the response value reached above 100, indicating its capability of detecting H₂S gas in such a low-temperature and low-vacuum environment (-20 °C, 5 Pa).

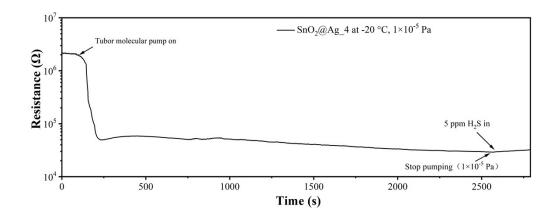


Figure S4. The response curve of SnO₂@Ag_4 gas sensor to 5 ppm H₂S in a low-temperature and high-vacuum condition (-20 °C, 1×10^{-5} Pa)

As shown in Figure S4, upon injecting 5 ppm of H_2S gas into the test system (-20 °C, 1×10^{-5} Pa), the SnO₂@Ag_4 gas sensor showed no response, and its resistance only had a slightly increasing trend, which was completely different from the experimental phenomenon observed under low-temperature and low-vacuum conditions.

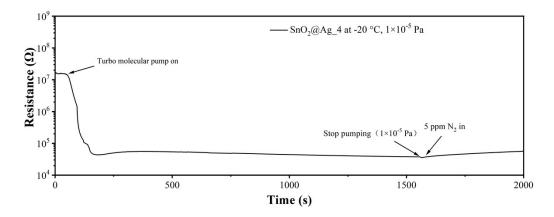


Figure S5. The response curve of SnO₂@Ag_4 gas sensor to 5 ppm N₂ in a low-temperature and high-vacuum condition (-20 °C, 1×10^{-5} Pa)

As shown in Figure S5, upon introducing an equal amount of N_2 , the resistance of the sensor also showed a slightly increasing trend, which was consistent with the experimental phenomenon observed in Figure S4. This indicates that under high-vacuum conditions, SnO₂ and Ag did not react with H₂S.

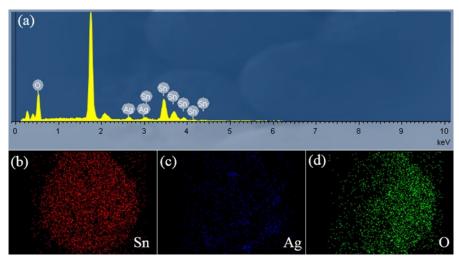


Figure S6. (a) EDS spectra, and (b-d) elemental mapping of $SnO_2@Ag_2O_2$

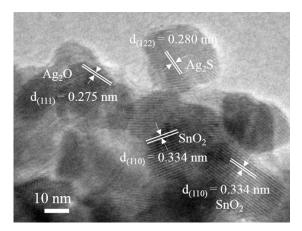


Figure S7. High-resolution TEM image of SnO_2@Ag_O_2 after exposure to $\rm H_2S$

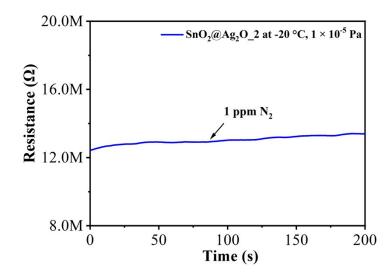


Figure S8. Gas sensing response curve of $SnO_2@Ag_2O_2$ gas sensor towards with 0.4 mL N_2

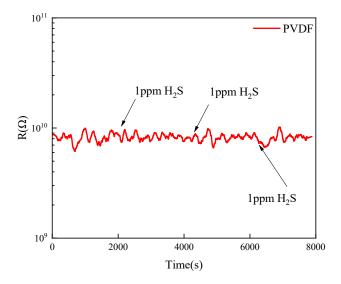


Figure S9 The PVDF for H_2S gas detection under vacuum $(1 \times 10^{-5} \text{ Pa})$.

Table 1

Method	Spacecraft Model	Mass (kg)	Power (W)	Size (cm × cm × cm)	Sensitivity and Limitation
Mass Spectrometry	Phoenix Mars Lander[1]	5.7	13	24 × 23× 18	0.7 to 4, 7–35, 14–70 and 28–140 amu;H ₂ O, CO ₂ , H ₂ , O ₂ , NH ₃ , SO ₂
	Huygens[2]	17.3	28	$\Phi 19.8 \times 47$	$2\sim141$ amu
	Pioneer Venus project[3]	3.81	12	$\Phi 0.2 imes 7.5$	1-64 amu; CO CO ₂ N ₂
Visible Spectrometer	LADEE[4]	3.6	13	-	Mean noise-equivalent power ~45 R/nm
Our method		<10g	<1 W	0.5	H ₂ S, 100 pb

1. Journal of the American Society for Mass Spectrometry Volume 19, Issue 10, October 2008, Pages 1377-1383

2. Niemann, H. B., Atreya, S. K., Bauer, S. J., et al. (2003). The gas chromatograph mass spectrometer for the Huygens probe. In The Cassini-Huygens Mission (pp. 553-591). Dordrecht: Springer Netherlands

3. Hoffman, J. H., Hodges, R. R., Wright, W. W., et al. (1980). Pioneer Venus sounder probe neutral gas mass spectrometer. IEEE Transactions on Geoscience and Remote Sensing, 18(1), 80-84.

4. Elphic, R.C. et al. (2015). The Lunar Atmosphere and Dust Environment Explorer Mission. In: Elphic, R., Russell, C. (eds) The Lunar Atmosphere and Dust Environment Explorer Mission (LADEE). Springer, Cham. Space Science Reviews