## **Electronic Supplementary Information**

## A Mechanical Actuated SnO<sub>2</sub> Nanowire for Small Molecules Sensing

Hongbin Feng,<sup>*a,b*</sup> Jin Huang,<sup>*b*</sup> and Jinghong Li\*<sup>*b*</sup>

 <sup>a</sup>Department of Chemistry, University of Science and Technology of China, Hefei 230026, China. <sup>b</sup>Department of Chemistry, Beijing Key Laboratory for Analytical Methods and Instrumentation, Tsinghua University, Beijing 100084, China. Fax: +86 10 6279 5290; Tel: +86 10 6279 5290; E-mail: jhli@mail.tsinghua.edu.cn

## Experimental

*Preparation of the*  $SnO_2$  *nanowires*: SnO\_2 nanowires were grown by the vapor–liquid–solid (VLS) process consists of a horizontal quartz reaction chamber, silicon substrate, a DC controller, a gas supply and control system, and a rotary pump system. The source material, high-purity (99.99%) powders of Sn, was first loaded in an alumina boat. Growth substrate of Si (100) was covered with a 10-nm-thick Au film, and then was placed on 2/3 height of the boat. The alumina boat was positioned at the center of a quartz reaction chamber that was inserted into a horizontal tube furnace. The furnace was heated from room temperature to 850 °C at a rate of 20 °C min<sup>-1</sup> under a flow of N<sub>2</sub> (160 sccm) with a trace amount of oxygen. The growth time was 1 h at 850 °C. The furnace was then cooled to room temperature at a rate of 5 °C min<sup>-1</sup>.

Structure and optical analysis of  $SnO_2$  nanowires: After nanowire growth, morphological investigations were performed by field-emission SEM (JSM 7401F). For further structural studies, the nanowires were removed from the Si growth substrate by sonication in isopropyl alcohol and then deposited on carbon-coated copper grids for TEM (JEM-1200EX) characterization. The Renishaw RM2000 microscopic confocal Raman spectrometer (Gloucestershire, United Kingdom) was used for Raman measurements with 514.5 nm excitation wavelengths at 25% energy and a laser spot size of 2.0  $\mu$ m.

*The calculation of the SnO2 nanowire:* The resonance frequency of the fork with the  $SnO_2$  FW is about 1700 Hz higher (Figure S2a, b). This increase in the resonance frequency is related to Young's modulus of the  $SnO_2$  FW, which is given by

$$E = \frac{4\pi^2 \left(f^2 - f_0^2\right) m_{eff} l}{A}$$
(1)

where *l* and *A* are the length and cross-sectional area of the SnO<sub>2</sub> FW, respectively, *f* and  $f_0$  are the resonance frequencies of the tuning fork with a SnO<sub>2</sub> FW and with the FW severed, respectively,  $m_{eff}$  is the effective mass for the oscillation beam of the tuning fork.<sup>20</sup> The Young's moduli of a single SnO<sub>2</sub> a nanowire was calculated as 178.2 *GPa*. This demonstrates a simple method of determining the mechanical properties of a semiconductor nanowire.



Figure S1. EDS analysis taken from the body of  $SnO_2$  nanowire. C and Cu signals in the spectra come from copper TEM grid coated with carbon film.



**Figure S2.** Frequency dependence of the oscillation amplitude of a microfabricated tuning fork with a single  $SnO_2$  nanowire (a), and of the same device after cutting the  $SnO_2$  wire (b). The corresponding resonance frequencies are *f* and *f*<sub>0</sub> respectively.



Figure S3. The resonance response of the sensor device with a broken  $SnO_2$  wire upon exposure to ethanol vapors in different concentrations.



**Figure S4.** The shifts in resonance responses of the  $SnO_2$  FW-based sensor between air and the 35 ppm ethanol vapor at 25 °C, showing the response time (a) and recovery time (b), respectively.



Figure S5. Resonance frequency shift as a function of ethanol concentration at 25, 30, and 35% RH on SnO<sub>2</sub> nanowire device.



Figure S6. Raman spectrum of the SnO<sub>2</sub> nanowires.



Figure S7. The mechanism of the gas adsorption on the SnO<sub>2</sub> wire.



**Figure S8.** Oscillation amplitude vs. frequency plots showing different resonance peaks of the MTF with a  $SnO_2$  wire between air and different vapor concentrations of ethanol,  $CH_2Cl_2$  (a), and heptane (b).



**Figure S9.** The resonance frequency shift of the  $SnO_2$  nanowire QTF was made as a function of formaldehyde concentration.



Figure S10. The frequency shift of the same  $SnO_2$  nanowire device exposed to several different organic vapors at 15 ppm.