

Electronic Supplementary Information for
**Catalytic-induced Sensing Effect of Triangular CeO₂
Nanoflakes for Enhanced BTEX Vapor Detection with
Conventional ZnO Gas Sensors**

Ding Wang,^a Yue Yin,^a Pengcheng Xu,^{*b,c} Feng Wang,^a Ping Wang,^a Jingcheng Xu,^a
Xianying Wang^{*a} and Xinxin Li^{b,c}

^a School of Material Science & Engineering, University of Shanghai for Science and Technology, Shanghai, 200093, China. E-mail: xianyingwang@usst.edu.cn

^b State Key Lab of Transducer Technology, Shanghai Institute of Microsystem and Information Technology, Chinese Academy of Sciences, 865 Changning Road, Shanghai 200050, China. E-mail: xpc@mail.sim.ac.cn

^c University of Chinese Academy of Sciences, Beijing 100049, China

S1. Preparation of CeO₂ nanomaterials

CeO₂ nanomaterials with different morphologies were prepared by hydrothermal method. In a typical process, the Ce(NO₃)₃ solution was first obtained by dissolving 0.068 g of Ce(NO₃)₃·6H₂O in 5 mL of deionized water. In addition, 9.6 g NaOH was dissolved in 35 mL deionized water. And then, the Ce(NO₃)₃ solution was dropped into NaOH solution with continuously stirring at room temperature. After stirring for 30 minutes, the mixture was hydrothermally treated at 100 °C in a 50 mL Teflon-lined autoclave for 24 h. The precursor was then washed several times with deionized water

and ethanol and dried for 10 h at 80 °C. The final CeO₂ nanorods were obtained by heat treatment for 3 h at 500 °C. The CeO₂ microsphere was obtained by the similar hydrothermal method, except that 60 mmol urea was used instead of NaOH. CeO₂ micro-polyhedrons was prepared by the similar method to CeO₂ nanorods with the 180 °C hydrothermal treatment.

S2. Materials Characterization

To characterize the morphology of the as-prepared samples, FE-SEM (FEI, Quanta FEG 450, USA) and Tecnai G220S-Twin transmission electron microscope were used with an accelerating voltage of 120 and 200 kV, respectively. The crystalline structure was analysed by using XRD (Breker, D8 Advance, Germany) with Cu-K α ($\lambda = 0.15418$ nm) radiation in the range of 10-70° at room temperature. The Raman spectra measurements were conducted by using Raman Microscopy (Horiba, LabRAM HR Evolution, France) with an excitation wavelength of 532 nm. On-line mass spectrum (on-line MS) was implemented on a Pfeiffer ThermoStar mass spectrometer.

S3. Gas sensor measurement

The gas sensing performance of gas sensor was measured by a static test system (Elite tech co. LTD) and the test gases were obtained by static distribution method. For the reducing gases and n-type MOS, the response of gas sensors is defined as $S=R_a/R_g$, where R_a is the gas sensor's resistance in air atmosphere and R_g is the resistance of gas sensor in target molecules contained atmosphere. The response and recovery time are defined as the time taken by the gas sensor to achieve the resistance changes ranging from R_a to $R_a-90%$ (R_a-R_g) and from R_g to $R_g+90%$ (R_a-R_g) in the case of adsorption and desorption of target gases, respectively. The exact values of temperature and relative humidity (RH) are

recorded by using a digital thermo-hygrometer (Dretec, model O251) which is prior placed on the testing chamber.

S4. Online MS experimental setup

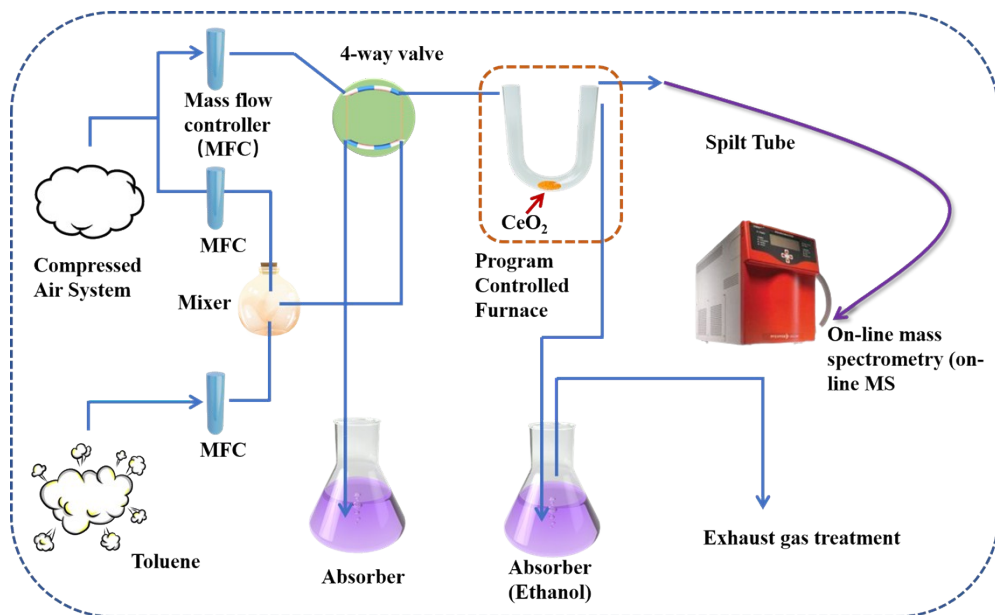


Fig. S1 Experimental setup for sensing product identification.

S5. Characterization results of CeO₂ materials

Table S1 Physicochemical characteristics of CeO₂ nanomaterials

Samples	Morphology ^a	Crystallite size ^b (nm)	I ₅₉₇ /I ₄₆₀ ^c (%)
CeO ₂ -MSs	Microsphere	12.3	1.1
CeO ₂ -MPHs	Micro-polyhedrons	30.9	3.1
CeO ₂ -NRs	Nanorod	17.5	4.7
CeO ₂ -NFs	Nanoflakes	10.3	11.7

^a The morphology of CeO₂ was obtained by SEM images in the Fig. 2(a-d), ^b the crystallite size of CeO₂ was estimated by using Scherrer equation based on XRD patterns in Fig. 2(e) of the manuscript, ^c the values of I₅₉₇/I₄₆₀ was calculated by Raman spectra in Fig. 4(b).

