Supporting Information

Humidity-Independent Oxide Semiconductor Chemiresistors Using Terbium-Doped SnO₂ Yolk-Shell Spheres for Real-Time Breath Analysis

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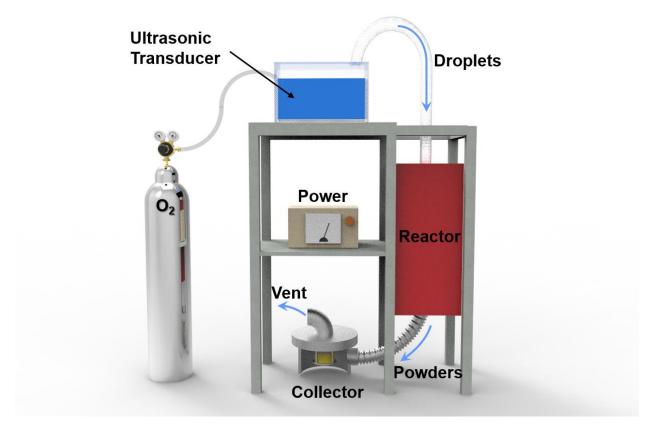


Figure S1. Schematic of the ultrasonic spray pyrolysis process for synthesizing yolk-shell spheres.

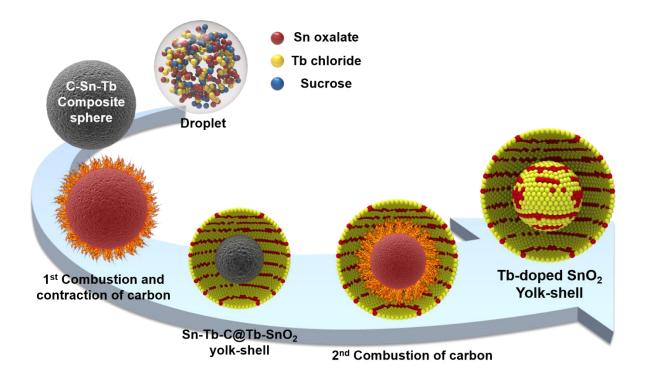


Figure S2. Formation mechanism of Tb-doped SnO₂ yolk-shells via spray pyrolysis.

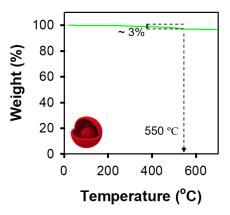


Figure S3. TG curve of as-prepared precursors of 5Tb-SnO₂ spheres.

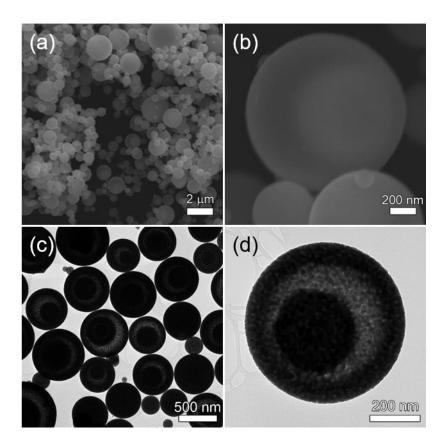


Figure S4. (a,b) SEM and (c,d) TEM images in pure SnO₂ yolk-shell spheres.

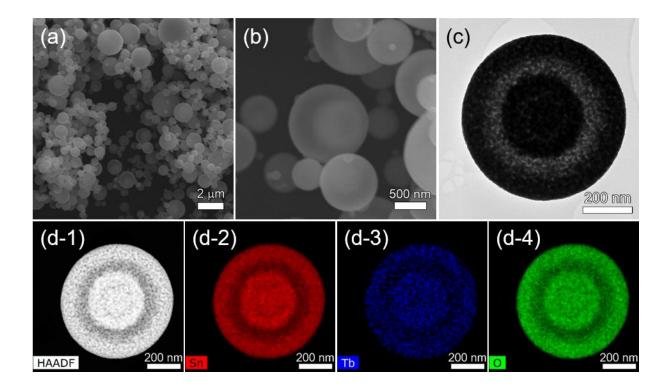


Figure S5. (a-c) SEM images and TEM images of 1Tb-SnO₂ yolk-shell spheres; (d) EDS elemental mapping of Sn, Tb, and O in 1Tb-SnO₂ yolk-shell spheres.

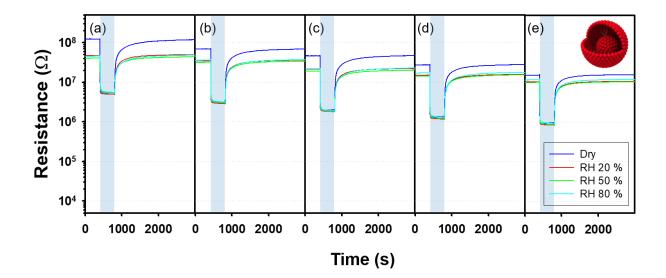


Figure S6. Gas-sensing transients of the 5Tb-SnO₂ sensor to 20 ppm of acetone at (a) 350 °C, (b) 375 °C, (c) 400 °C, (d) 425 °C and (e) 450 °C in dry and humid conditions (RH = 20, 50, 80%).

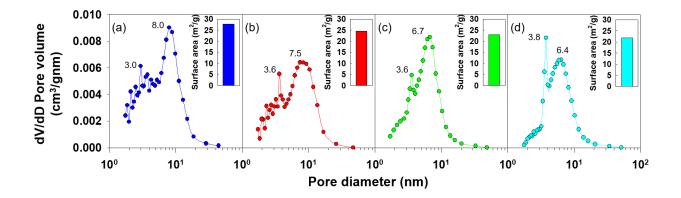


Figure S7. Pore-size distributions and BET specific surface areas of (a) pure SnO₂, (b) 1Tb-SnO₂,
(c) 5Tb-SnO₂, and (d) 15Tb-SnO₂ yolk-shell spheres measured by N₂ desorption isotherm.

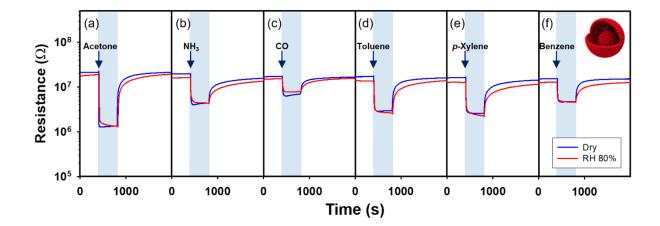


Figure S8. Gas-sensing transients of the 5Tb-SnO₂ sensor to 20 ppm of (a) acetone, (b) NH₃, (c) CO, (d) toluene, (e) *p*-xylene, and (f) benzene at 450 °C in dry and humid conditions (RH = 80%).

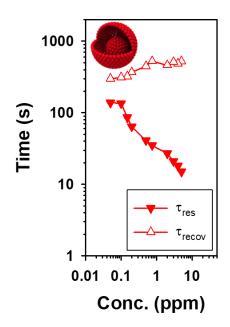


Figure S9. Response and recovery times of $5\text{Tb}-\text{SnO}_2$ sensor upon exposures to acetone and air as a function of acetone concentration at 450 °C in humid condition (RH = 80%).