

## SIIC027

### THE ANALYSIS OF TIN(IV) OXIDE PREPARED VIA HYDROTHERMAL TREATMENT AND ETHANOL GAS SENSING PROPERTIES

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#### **Abstract:**

Tin(IV) oxide SnO<sub>2</sub> is a semiconductor material with a wide bandgap energy (3.6 eV). It is particularly interesting, because it has high thermal and chemical stability and high surface to-volume ratio. Tin(IV) oxide nanostructure can be synthesized using various techniques such as sol-gel, electrospinning, thermal evaporation, and co-precipitation. They produce nanostructure, however, they involve expensive instrumentation and high operational temperature. Among them, the hydrothermal synthesis is widely used nowadays because of its simplicity process. This study was carried out to analyze the data of SnO<sub>2</sub> nanostructures which were prepared via hydrothermal treatment at different heat treatment durations (6, 12, 18, and 24 hours) under constant temperature, 180 °C. The effect of different heat treatment durations on the structure, size and morphology was investigated by X-ray diffraction (XRD), Ultraviolet-visible spectroscopy (UV-vis), Field emission scanning electron microscope (FESEM), and Fourier-transform infrared spectroscopy (FTIR). It was found that the formation of SnO<sub>2</sub> nanorods for all samples grew into closely packed flower-like nanostructures. The diameter of the SnO<sub>2</sub> nanorods reduced from 43.30 to 24.96 nm for heat treatment durations of 6 to 24 hours, respectively. For HRTEM analysis the diameter of SnO<sub>2</sub> nanorods for 24 hours was approximately 22.89 ± 4.84 nm. For UV-vis analysis, it was observed that the band gap energy increased (3.52, 3.56, 3.57 and 3.6 eV). In the second part of the study, the best prepared SnO<sub>2</sub> sample was tested on ethanol gas at different operating temperatures (200 °C - 450 °C). The highest sensor response ( $R_o/R_g \sim 1645.265$ ) was obtained at temperature 450 °C. Therefore, the operating temperature with the highest sensor response was selected as the operating temperature.

#### **Keywords:**

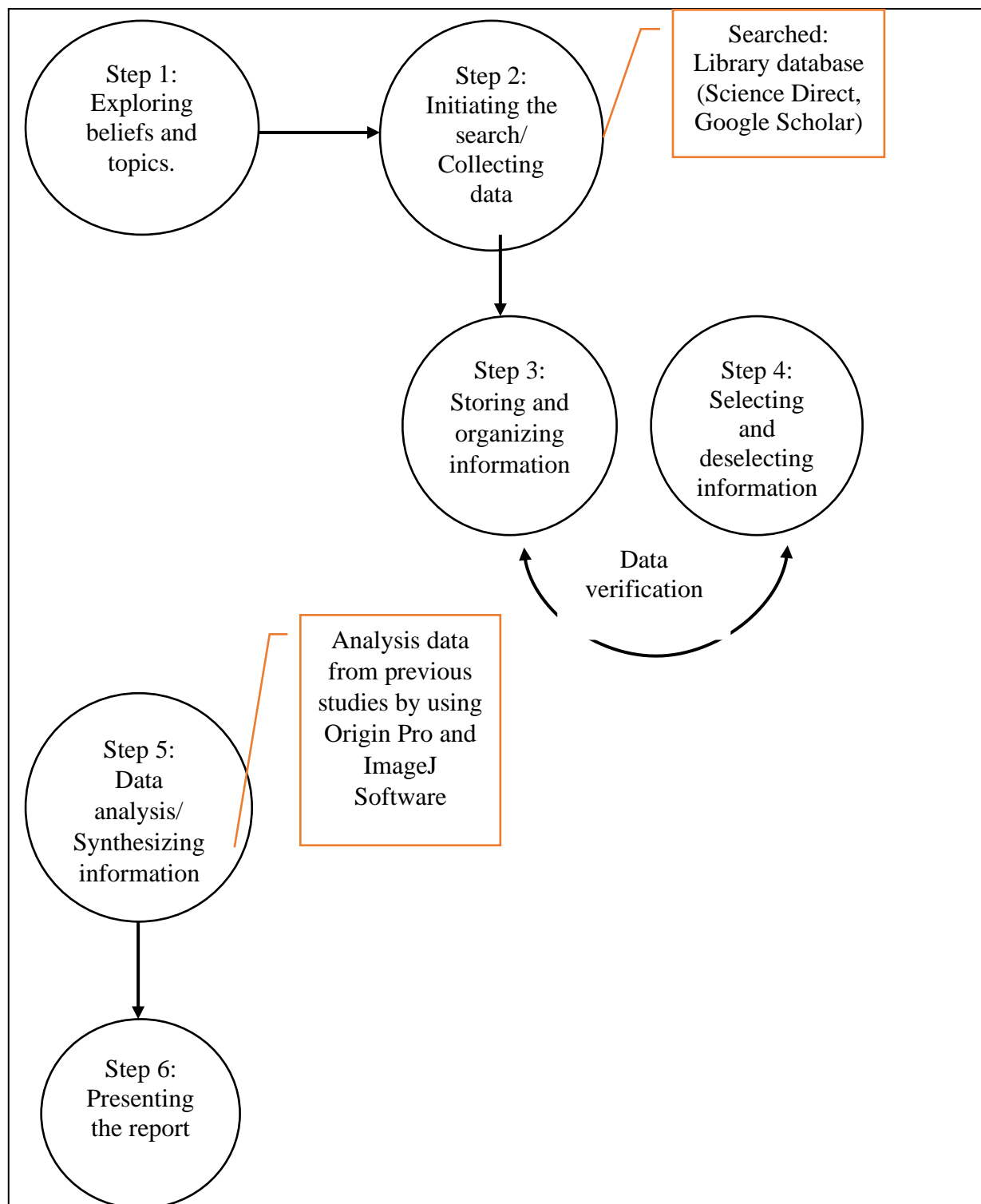
Hydrothermal, Tin(IV) oxide, Heat treatment duration; Ethanol gas sensor; Operating temperature

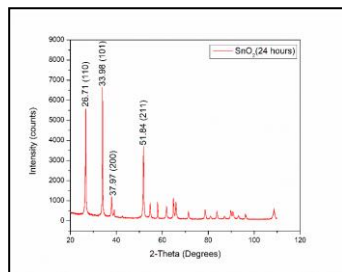
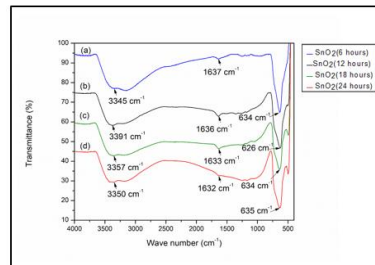
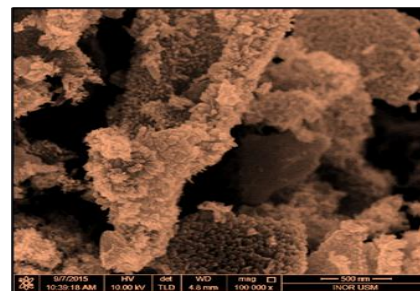
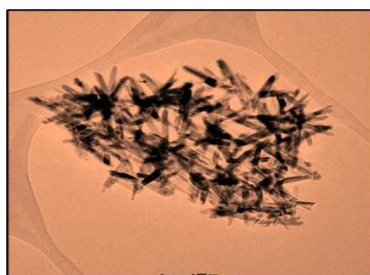
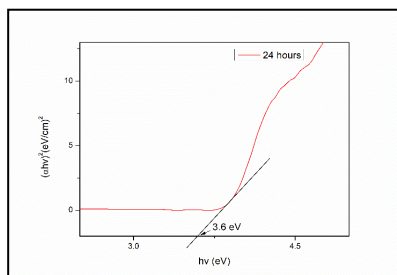
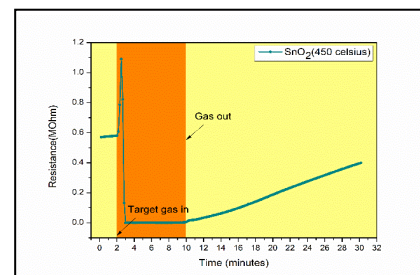
#### **Objectives:**

- To analyse the data of SnO<sub>2</sub> nanostructures prepared by hydrothermal method at different heat treatment durations and compare the data with previous studies.

- To determine the performance of ethanol gas sensing at different operating temperatures and propose a plausible mechanism for ethanol gas sensing.

### Methodology



**Results:****SnO<sub>2</sub> for 24 hours - XRD****SnO<sub>2</sub> - FTIR****SnO<sub>2</sub> for 24 hours - FESEM****SnO<sub>2</sub> for 24 hours - HRTEM****SnO<sub>2</sub> for 24 hours - UV-vis****SnO<sub>2</sub> nanorods - 450 °C****Conclusion:**

The XRD results show that all the diffraction peaks for SnO<sub>2</sub> can be assigned to rutile tetragonal structure and the dominant peaks were observed for all the samples (6, 12, 18, and 24 hours) at (110), (101), and (211). Based on the FTIR analysis, the functional group formed in the SnO<sub>2</sub> samples were O-H stretching, O-H bending, and O-Sn-O stretching. Next, the diameter of SnO<sub>2</sub> samples in FESEM analysis were decreased from 43.30 nm to 24.96 nm with increase in heat treatment duration (6-24 hours). For UV-vis analysis, it was observed that, the band gap energy increased (3.52, 3.56, 3.57, and 3.6 eV) as the size of SnO<sub>2</sub> nanostructures decreased. The ethanol gas sensors possess the best response performance at the operating temperature 450 °C with a response of ( $R_o/R_g \sim 1.6 \times 10^3$ ) and response time 3 minutes. As the operating temperature increased to 450 °C, atomic ions species such as O<sup>2-</sup> and O<sup>-</sup> are dominated. Then, the oxygen ions absorbed on the surface of the SnO<sub>2</sub> leads to the structure of an electron-depleted surface layer, which results in the reduction in the electronic conductivity of SnO<sub>2</sub>.