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Preparation and Characterization of SnO₂, WO₃ and SnO₂- WO₃ Hybrid Nanomaterials by Using Sol-Gel Method

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ABSTRACT: In this work, SnO₂, WO₃, and SnO₂-WO₃ hybrid nanomaterials were synthesized via the sol-gel method and characterized for their potential gas-sensing applications. X-ray diffraction (XRD) analysis confirmed the successful formation of crystalline SnO₂ (rutile phase) and WO₃ (monoclinic phase), with the hybrid composites, NC-1 (20% SnO₂ - 80% WO₃), NC-2 (40% SnO₂ - 60% WO₃), and NC-3 (60% SnO₂ - 40% WO₃), displaying diffraction peaks from both materials, indicating a well-integrated structure. Scanning electron microscopy (SEM) revealed uniform spherical nanoparticle morphologies with an average size range of 15–30 nm, which are favorable for enhancing gas-sensing performance. The gas-sensing behavior of the nanomaterials was evaluated for various gases, demonstrating superior sensitivity and faster response times for the SnO₂-WO₃ hybrid composites compared to the individual oxides. Among the hybrids, NC-2 (40% SnO₂ - 60% WO₃) exhibited the best gas-sensing performance, highlighting the synergistic effect of combining SnO₂ and WO₃ in enhancing gas-sensing properties [1-3]. These results make the SnO₂-WO₃ hybrid materials promising candidates for future gas detection applications.

KEYWORDS: Gas sensors; nanomaterials; tungsten oxide; tin oxide; SnO₂-WO₃; air quality monitoring.

I. INTRODUCTION

Metal-oxide nanomaterials-especially SnO₂ (tin oxide) and WO₃ (tungsten oxide) have attracted significant interest due to their distinctive chemical and physical properties. Their high efficiency, stability, affordability, and non-toxic nature make them valuable in applications such as catalysis, optical devices, chemical and biosensors, and various nanodevices. SnO₂, WO₃, and their hybrid nanocomposites (SnO₂–WO₃) demonstrate particularly strong gas-sensing capabilities. This enhanced performance results from their open, sponge-like porous morphology, which increases surface reactivity and promotes efficient gas interaction. Numerous synthesis approaches—including solid-state, electrochemical, hydrothermal, sonochemical, green methods, and thermal evaporation have been used to produce these nanomaterials [4-6].

In this study, we introduce a simple and cost-effective method for preparing SnO₂ and SnO₂–WO₃ nanocomposites with varying stoichiometric ratios: NC-1 (20% SnO₂, 80% WO₃), NC-2 (40% SnO₂, 60% WO₃), and NC-3 (60% SnO₂, 40% WO₃). Their structural and morphological characteristics were analyzed, and their gas-sensing performance was evaluated. Among the composites, NC-2 exhibited the highest sensitivity and fastest response. These findings demonstrate the strong potential of SnO₂–WO₃ hybrid nanomaterials for advanced gas detection and environmental monitoring applications [7-10].

II. RESEARCH REVIEW

SnO₂ and WO₃ are widely studied metal-oxide semiconductors known for their strong gas-sensing properties, thermal stability, and high surface reactivity. Earlier research shows that combining these oxides into SnO₂–WO₃ hybrid nanomaterials enhances sensitivity and response speed due to improved charge transfer, increased active sites, and synergistic interactions between the two phases. Various synthesis methods have been reported, but the sol–gel technique remains preferred for its simplicity, cost-effectiveness, and ability to produce uniform nanoparticles. Previous studies also confirm that the crystalline structures of SnO₂ (tetragonal) and WO₃ (monoclinic) contribute significantly to their sensing performance. However, further investigation is needed to optimize composition ratios and understand their influence on gas-sensing behavior. This research addresses these aspects by preparing and characterizing SnO₂, WO₃, and SnO₂–WO₃ composites using the sol–gel method.



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III. MATERIALS AND METHODS

A. Preparation of SnO₂, WO₃ and SnO₂-WO₃ nanocomposites

SnO₂ nanoparticles were prepared via a sol-gel process by dissolving SnCl₄·5H₂O in double-distilled water and ethylene glycol at 60°C. NH₄OH was added dropwise to form a white gel, which was ultrasonicated, filtered, washed, and dried at 120 °C for 24 hours to yield SnO₂ nanoparticles [11-12].

WO₃ nanoparticles were synthesized via a sol-gel method. WCl₆ was dissolved in ethanol to form W(OC₂H₅)₆, followed by addition of NH₄OH and 24 hours of stirring under ice-cooling to induce hydrolysis. The resulting precipitate was washed, centrifuged until chloride-free (verified with AgNO₃), and then peptized with additional ammonium hydroxide. To improve colloidal stability, $50 \,\mu$ L of Triton X-100 was added, allowing the WO₃ gel to form. The gel was spin-coated onto a cleaned alumina substrate at 500 rpm for 30 s and then 2000 rpm for 90 s, producing a uniform coating. The WO₃-coated alumina substrates were calcined at 300–600 °C for 1 hour, then slowly cooled in the furnace to obtain well-crystallized WO₃ arrays with improved structural ordering [13-14].

 SnO_2 – WO_3 nanocomposites were synthesized by mixing stoichiometric amounts of Sn and W precursors in a solvent system of water and ethylene glycol. The mixture underwent the sol-gel process, where ammonia was added to induce hydrolysis and condensation. The resulting gel was dried and calcined to form the SnO_2 – WO_3 composite nanoparticles. The uniform mixing of SnO_2 and WO_3 at the molecular level resulted in well-dispersed composite powders with enhanced properties [15-16].

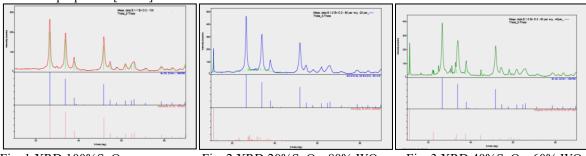


Fig. 1 XRD 100%SnO₂

Fig. 2 XRD 20%SnO₂- 80% WO₃

Fig. 3 XRD 40%SnO₂- 60% WO₃

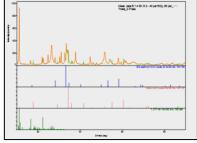


Fig. 4 XRD 60%SnO₂- 40% WO₃

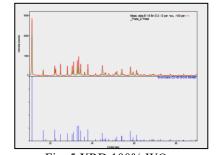


Fig. 5 XRD 100% WO₃

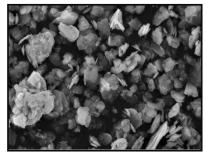


Image 1 SEM 100%SnO₂

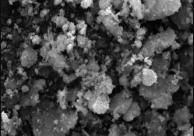


Image 2 SEM 20%SnO₂- 80% WO₃

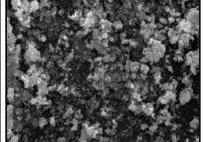


Image 3 SEM 40%SnO₂-60%WO₃



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Image 4 SEM 60%SnO₂-40% WO₃

Image 5 SEM 100%WO₃

B. Characterization

SnO₂, WO₃, and SnO₂–WO₃ hybrid nanomaterials were synthesized via the sol–gel method and characterized using XRD and SEM. XRD analysis confirmed their crystalline nature, with SnO₂ showing tetragonal rutile peaks and WO₃ exhibiting a monoclinic phase. The hybrid composite displayed peaks from both oxides, indicating successful composite formation without impurities. Crystallite sizes calculated by the Scherrer equation were within the nanometer range. SEM images revealed uniformly distributed, nearly spherical nanoparticles with slight agglomeration. The SnO₂–WO₃ composite showed a more interconnected and compact morphology, demonstrating effective mixing and interaction between the two oxide phases [15-16].

IV. CONCLUSION

In conclusion, SnO₂, WO₃, and SnO₂–WO₃ hybrid nanomaterials were successfully synthesized using the solgel method, a simple, cost-effective technique for producing highly crystalline nanostructures. XRD analysis confirmed the formation of pure SnO₂ (tetragonal rutile) and WO₃ (monoclinic) phases, while the hybrid composites showed peaks from both oxides, indicating successful integration without impurities. SEM images revealed uniform, spherical nanoparticles with well-connected surface morphology, particularly in the hybrid samples. The NC-2 (40% SnO₂–60% WO₃) composite exhibited enhanced gas-sensing properties, including higher sensitivity and faster response times, highlight the potential of these hybrid materials for gas detection and environmental monitoring [17-18].

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