

Visible-Light-Active N-Doped TiO₂ coupled with Bi₂WO₆ Z-Scheme Heterojunctions for Indoor VOC and CO₂ Abatement

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Highlights

- N-doped TiO₂ and Bi₂WO₆/N-TiO₂ Z-scheme heterojunctions enabled visible-light activation for VOC/CO₂ abatement
- Solution combustion and sol-gel doping strategies yield tailored decreasing bandgap and charge separation.
- Hydrothermally synthesized Bi₂WO₆ coupling enhances stability and visible-light utilization in defective nanostructures
- Synthesis parameters directly control crystallinity, surface area, and optical properties for indoor pollution remediation

1. Introduction

Indoor air pollution from volatile organic compounds (VOCs) like ethylene and CO₂ threatens human health, deteriorating respiratory issues and contributing to the global warming [1]. Photocatalytic degradation under visible light offers a sustainable, low-energy solution, with TiO₂ standing out for its stability and oxidizing power but limited by its wide bandgap and UV dependence [2]. This study develops visible-light-active TiO₂-based photocatalysts via non-metal doping and Z-scheme heterojunctions, targeting efficient VOC degradation and CO₂ reduction under realistic indoor conditions. Nitrogen (N) doping narrows the bandgap through valence band modulation, while bismuth tungstate (Bi₂WO₆) coupling promotes charge separation and photocorrosion resistance, addressing key limitations for practical deployment [3].

2. Methods

Undoped and N-doped TiO₂ were synthesized via hydrothermal (HYD), solution combustion (SCS), and sol-gel (SG) synthesis, with N: Ti molar ratios from 3:1 to 1:1 using urea and ammonium hydroxide as N sources. Bi₂WO₆, Bi₂WO₆/TiO₂, and Bi₂WO₆/N-TiO₂ composites were prepared using hydrothermal synthesis to form Z-scheme heterostructures. Materials were characterized by XRD (crystallinity, phase, crystallite size), N₂ physisorption (BET surface area, pore volume), and DR-UV-Vis (bandgap, visible-light absorption). Photocatalytic oxidation of ethylene (100ppm in air) was evaluated with a continuous-flow gas-phase reactor under Xe arc lamp irradiation, with GC analysis for performance screening to be presented.

3. Results and discussion

XRD confirmed crystalline anatase TiO₂ phases across samples, with minor FWHM shifts indicating preserved structure post-doping and coupling where Bi₂WO₆ also maintained its orthorhombic phase. UV-Vis revealed bandgap narrowing in N-TiO₂ (enhanced visible absorption) and further reduction in Bi₂WO₆/N-TiO₂, enabling efficient photoexcitation. BET showed TiO₂_HYD with the highest surface area, decreasing in N-TiO₂_SG and Bi₂WO₆ composites due to heterojunction formation—balancing adsorption capacity with charge dynamics. These traits position Bi₂WO₆/N-TiO₂ as optimal for visible-light VOC/CO₂ conversion, with synthesis variations (e.g., SCS for bulk efficiency) offering tunable defect-rich sites for superior electron-hole separation versus pristine TiO₂. Ethylene conversion for synthesized samples with respect to pristine TiO₂, correlating to optimized bandgap and defects; full conversion and mineralization (CO₂ yield) will be presented.

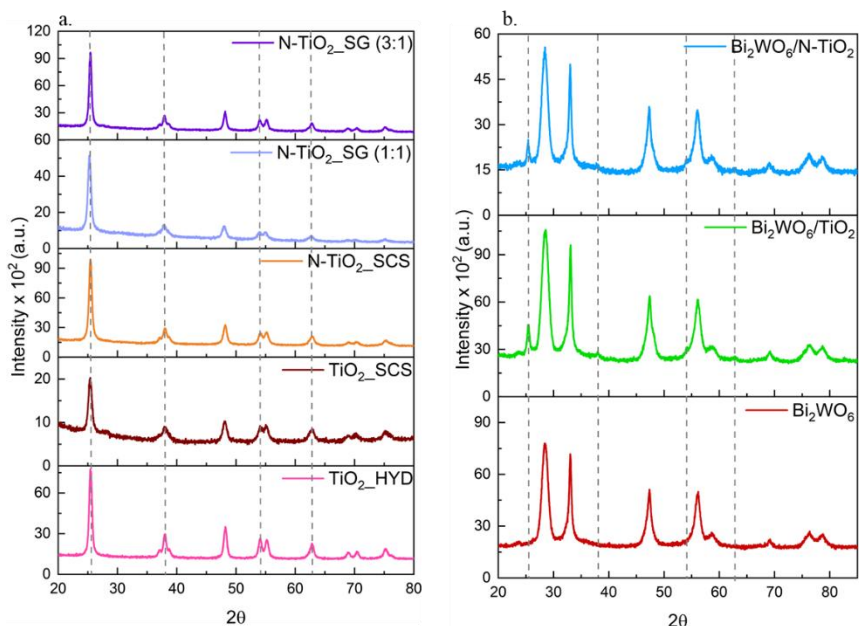


Figure 1. a. XRD of $\text{TiO}_2_{\text{HYD}}$, $\text{TiO}_2_{\text{SCS}}$, $\text{N-TiO}_2_{\text{SCS}}$, $\text{N-TiO}_2_{\text{SG}}$, b. XRD of Bi_2WO_6 , $\text{Bi}_2\text{WO}_6/\text{TiO}_2$ and $\text{Bi}_2\text{WO}_6/\text{N-TiO}_2$

4. Conclusions

N-doping and Bi_2WO_6 Z-scheme engineering transformed TiO_2 into a versatile visible-light photocatalyst, where the synthesis routes shaped the critical optical and textural properties for indoor air purification. Ongoing reactor screening will quantify ethylene conversion, CO_2 mineralization, and stability, promising breakthroughs in sustainable indoor air pollution control.

References

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Keywords

Indoor air pollution; visible-light photocatalysis; N-doped TiO_2 ; Bi_2WO_6 coupled TiO_2 ; Z-scheme heterojunction; VOC oxidation; CO_2 reduction

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