

Co₃O₄-ZnO/rGO catalyst preparation and rhodamine B degradation by sulfate radical photocatalysis

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Cite this as: Zhanmei ZHANG, Yi ZHANG, Xilin CHEN, Ziran HUANG, Zuqin ZOU, Huaili ZHENG, 2023. Co₃O₄-ZnO/rGO catalyst preparation and rhodamine B degradation by sulfate radical photocatalysis. *Journal of Zhejiang University-SCIENCE A (Applied Physics & Engineering)*, 24(8):710-721.

<https://doi.org/10.1631/jzus.A2200490>

sulfate-radical-mediated photocatalysis (SR-photo)

Under the influence of light and the catalyst in the SR-photo system, PMS can also produce other active species, such as sulfate ($\text{SO}_4^{\bullet-}$) and hydroxyl radicals ($\bullet\text{OH}$). Improve the effectiveness of photocatalytic oxidation reactions and reduce the recombination of electron hole pairs during the photocatalytic process

The combination of p-type Co_3O_4 and n-type ZnO creates a favorable internal electric field for photocatalysis, improves the efficiency of the photo reaction, and can effectively activate PMS. The presence of rGO further enhances the catalytic performance

Preparation of $\text{Co}_3\text{O}_4\text{-ZnO/rGO}$

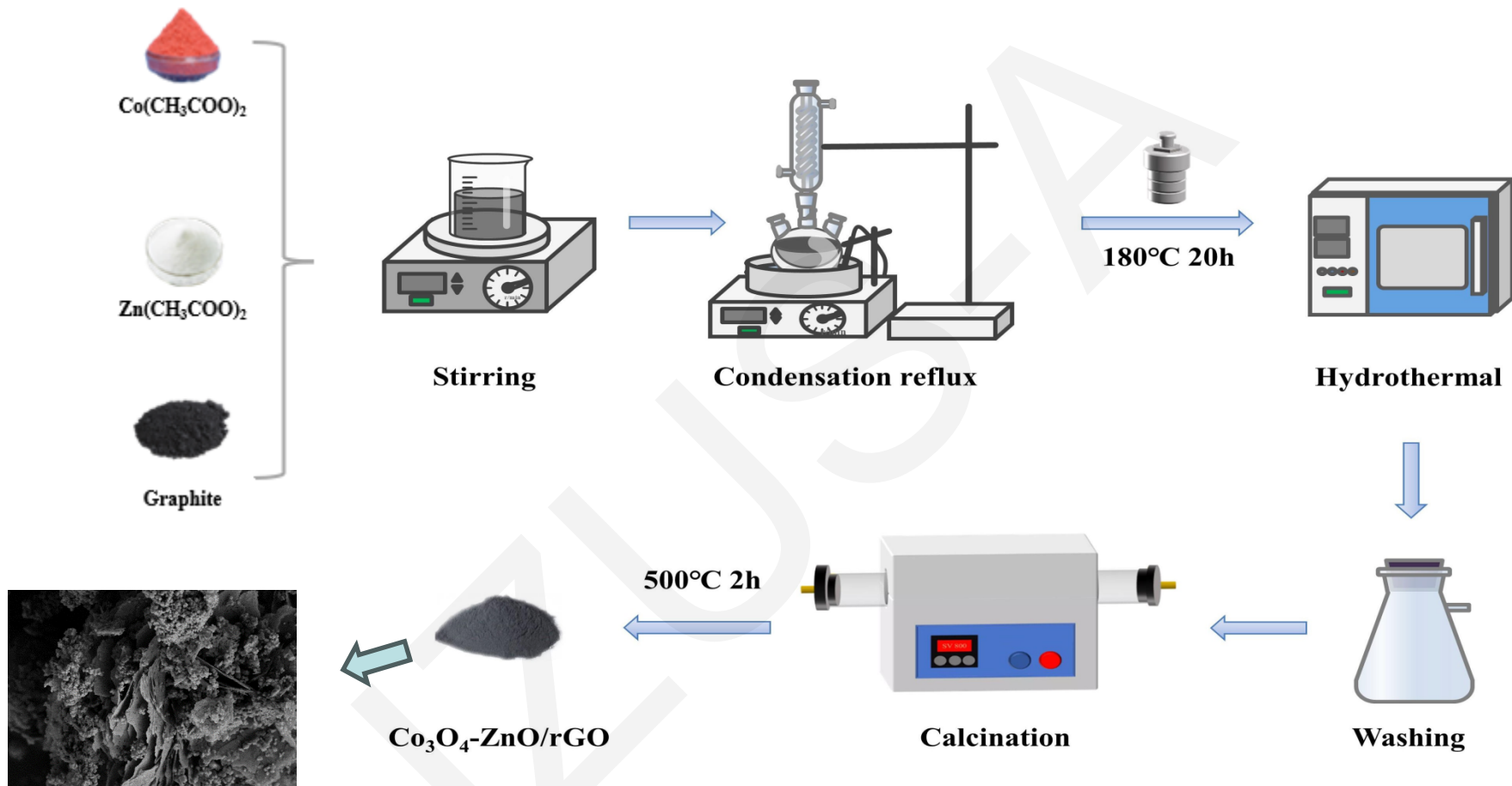


Fig. 1. Synthesis path of $\text{Co}_3\text{O}_4\text{-ZnO/rGO}$ heterojunction

Degradation of RhB by UV/Co₃O₄-ZnO/rGO/PMS

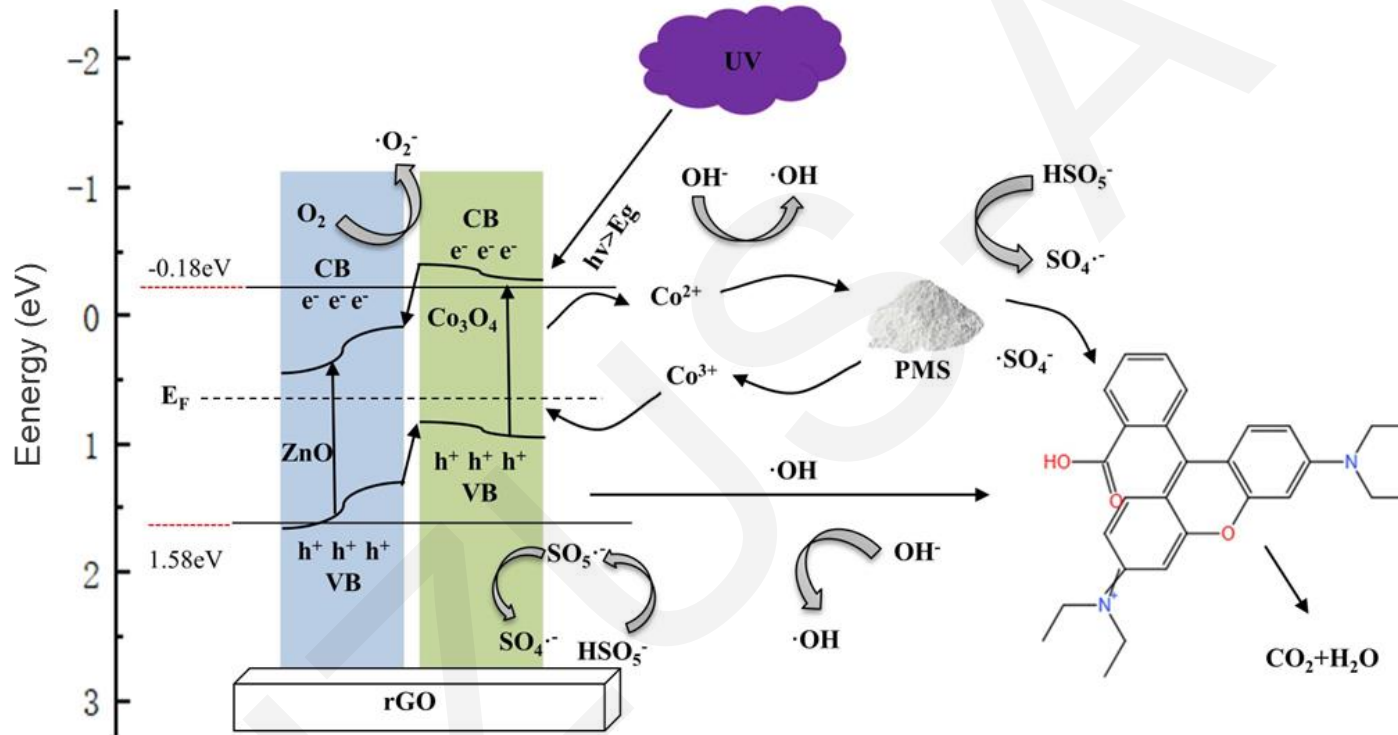


Fig. 2. mechanism of RhB degradation in the UV/Co₃O₄-ZnO/rGO/PMS

Conclusions

- The distinct particle/sheet structure of $\text{Co}_3\text{O}_4\text{-ZnO/rGO}$ offers a large surface area with numerous active sites for the reaction, which enables the extension of the carrier lifetime and speeds up the transport and separation of electron-hole pairs, resulting in superior photocatalytic efficiency and the ability to activate PMS compared to stand-alone Co_3O_4 or ZnO .
- PMS and photocatalysis exhibit good synergy. PMS better suppresses electron-hole pair recombination by capturing electrons, and the photogenerated carriers generated by illumination also strengthen the redox cycle between Co(II) and Co(III)
- In the presence of major radicals, such as $\text{SO}_4^{\cdot-}$ and $\cdot\text{OH}$, the UV/ $\text{Co}_3\text{O}_4\text{-ZnO/rGO/PMS}$ system cleared 90.4% of the RhB in 40 min.