

Electronic Supplementary Information

Fabrication of highly visible-light-responsive $\text{ZnFe}_2\text{O}_4/\text{TiO}_2$ heterostructures for the enhanced photocatalytic degradation of organic dyes

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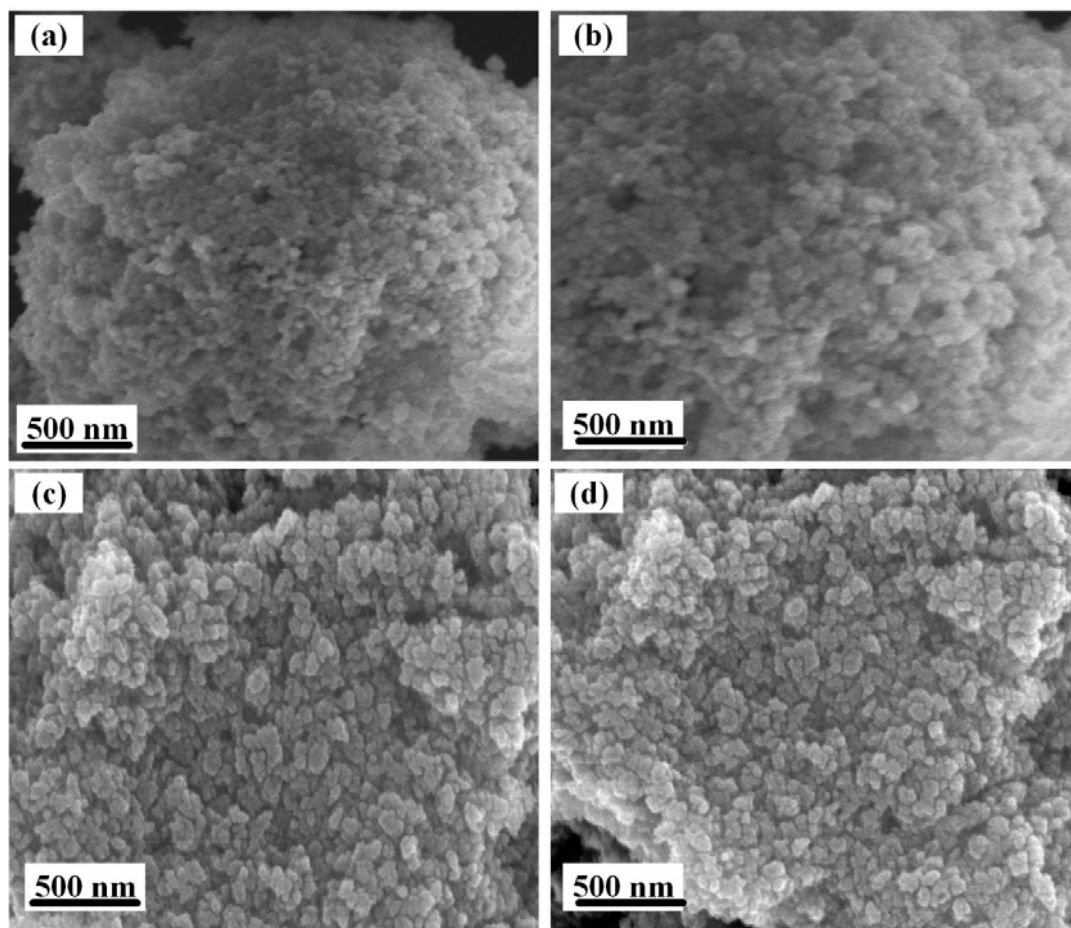


Fig. S1. SEM images of ZnFe₂O₄-TiO₂ nanocomposites at various ZnFe₂O₄ loadings of (a) 0.2, (b) 0.5 (c) 1 and (d) 2 wt%.

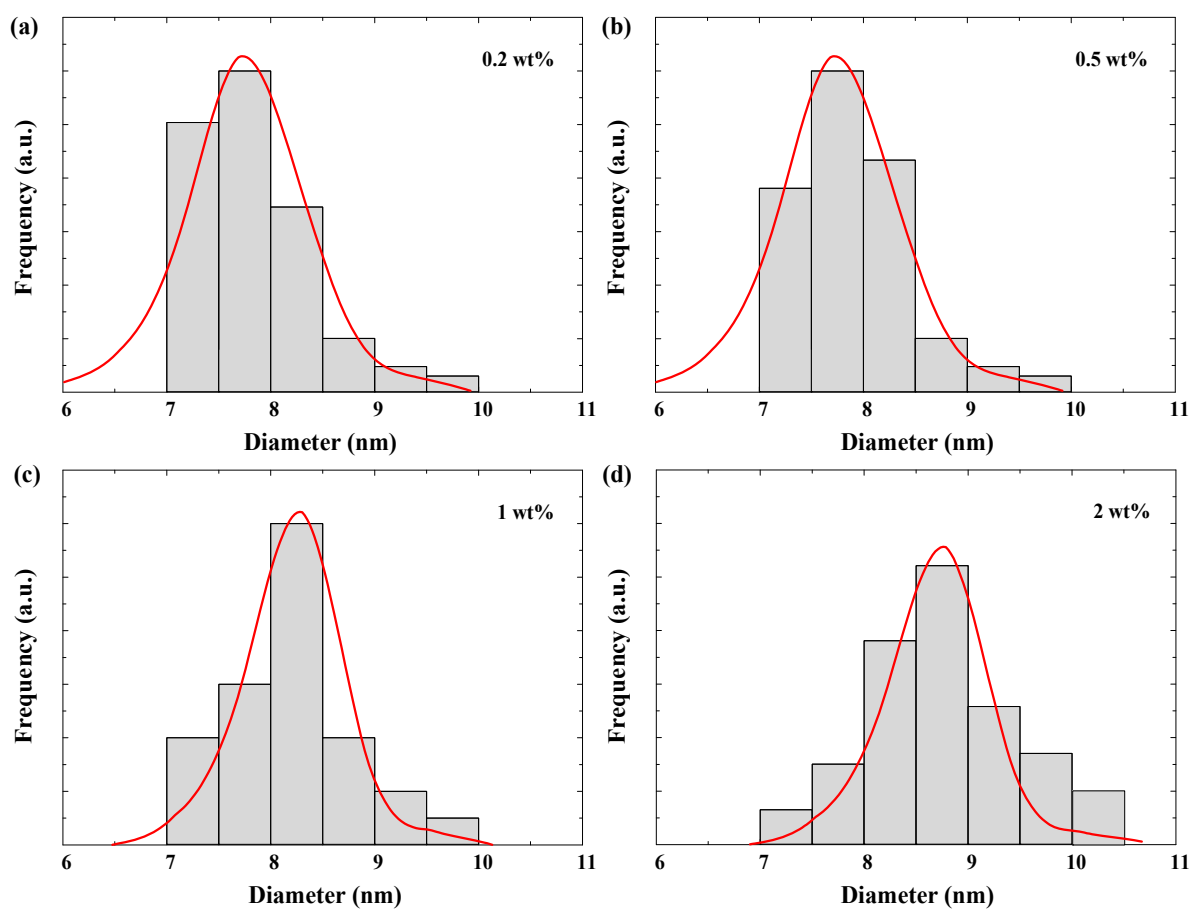


Fig. S2. Particle size distribution histogram of (a) 0.2, (b) 0.5, (c) 1 and 2 wt% ZnFe₂O₄-TiO₂ nanocomposites.

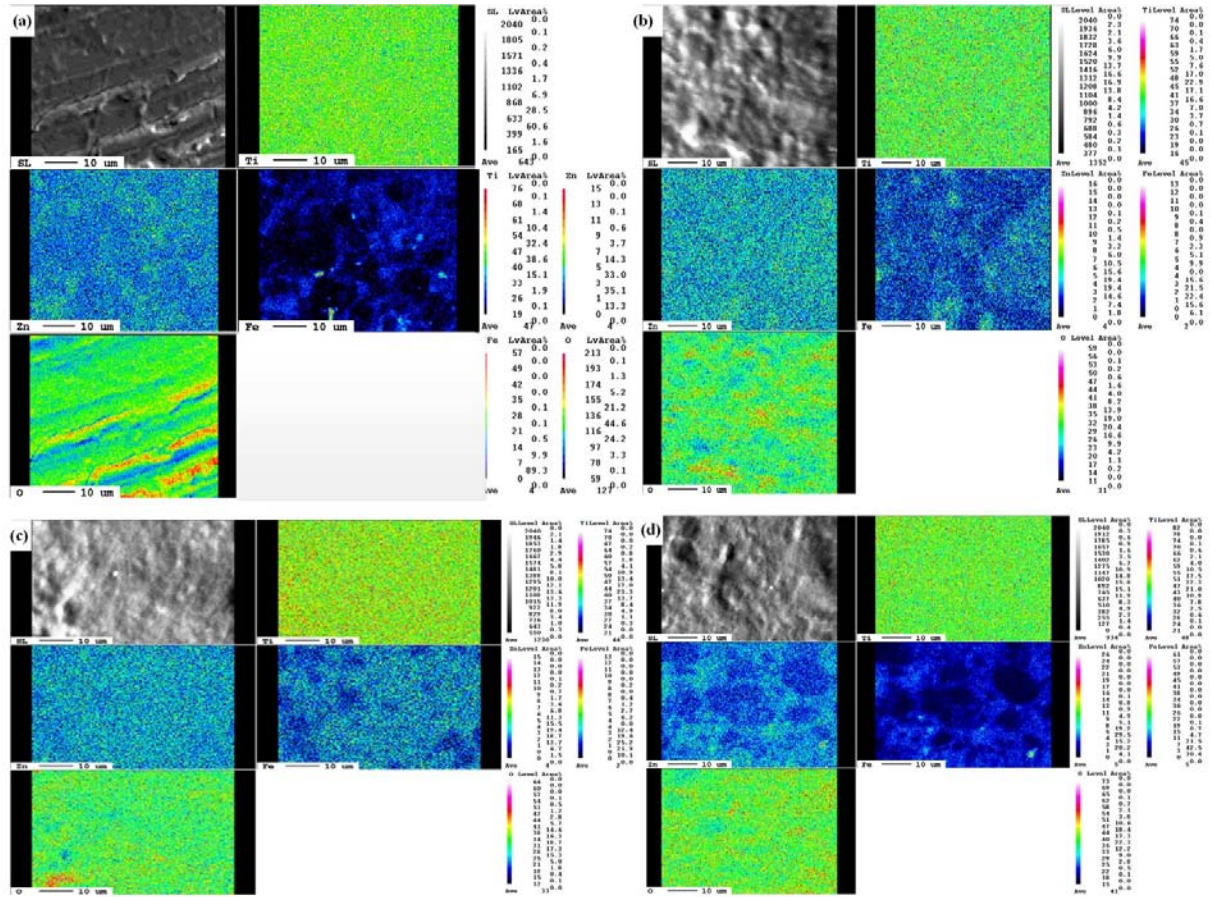


Fig. S3. EPMA images of ZnFe₂O₄-TiO₂ nanocomposites at various ZnFe₂O₄ loadings of (a) 0.2, (b) 0.5, (c) 1 and (d) 2 wt%.

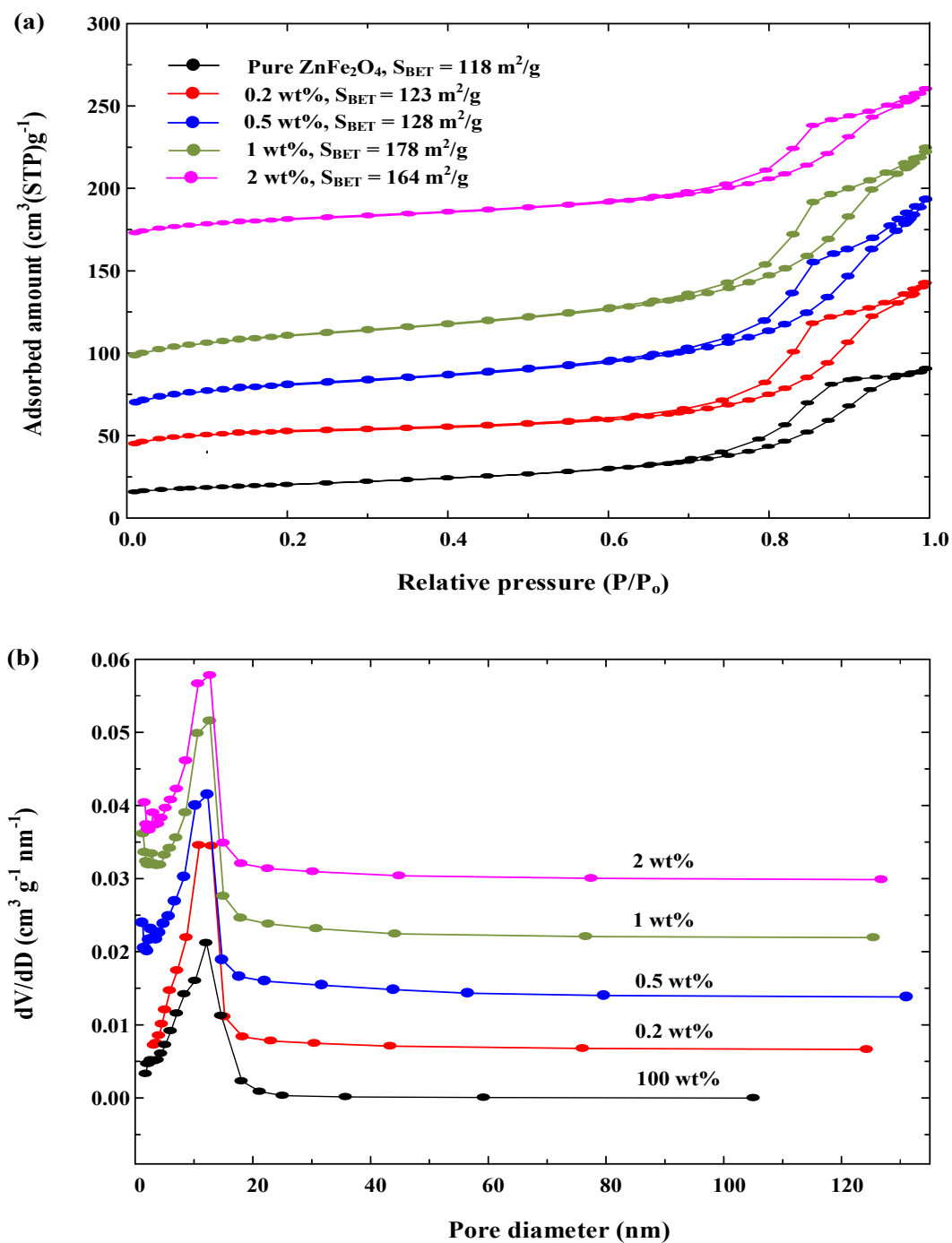


Fig. S4. (a) Nitrogen adsorption-desorption isotherms and (b) pore size distribution of ZnFe₂O₄ and ZnFe₂O₄-TiO₂ nanocomposites at various loadings of 0.2-2 wt%.

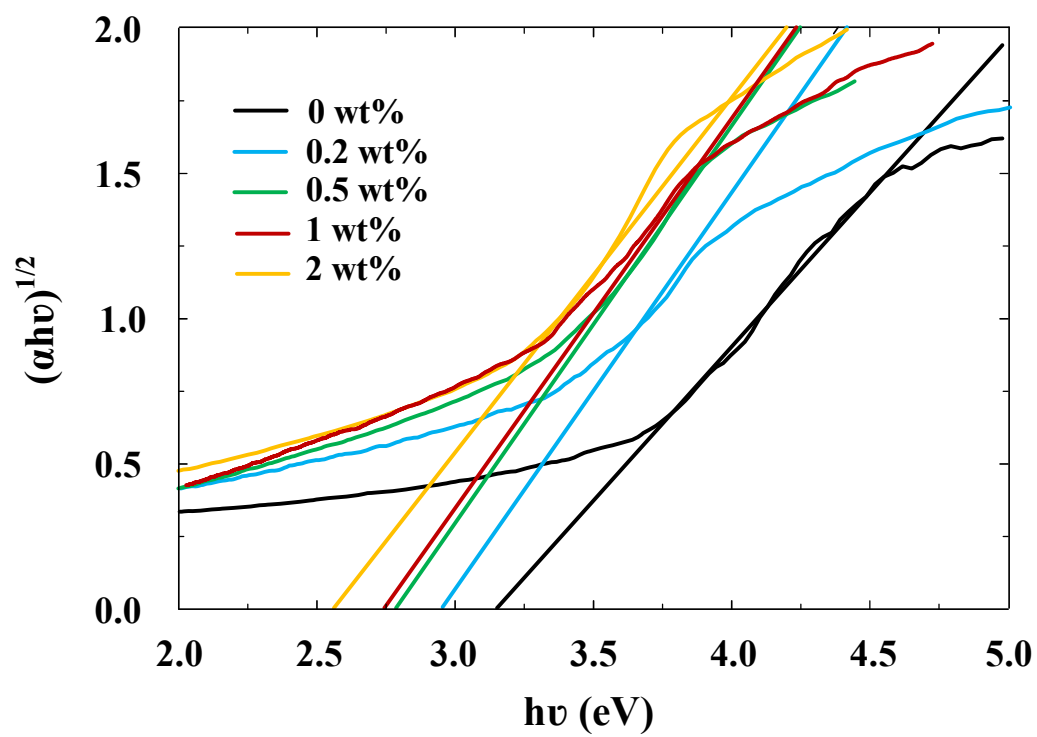


Fig. S5. Plots of $(\alpha h\nu)^{1/2}$ vs. photon energy $h\nu$.

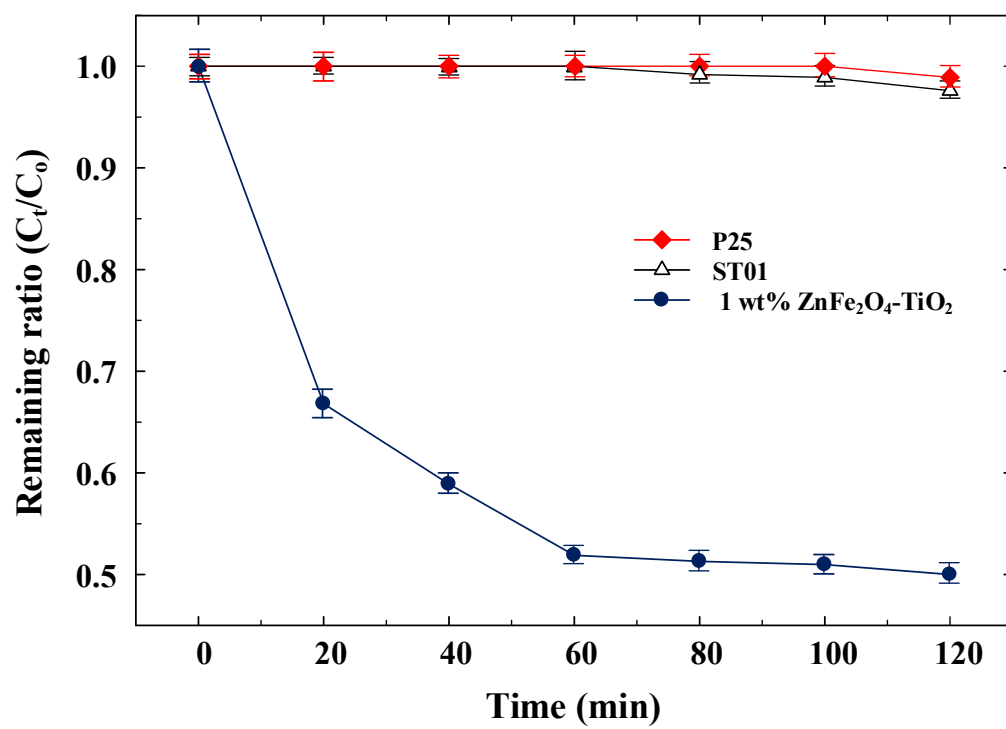


Fig. S6. The adsorption of RhB by various TiO₂-based nanomaterials.

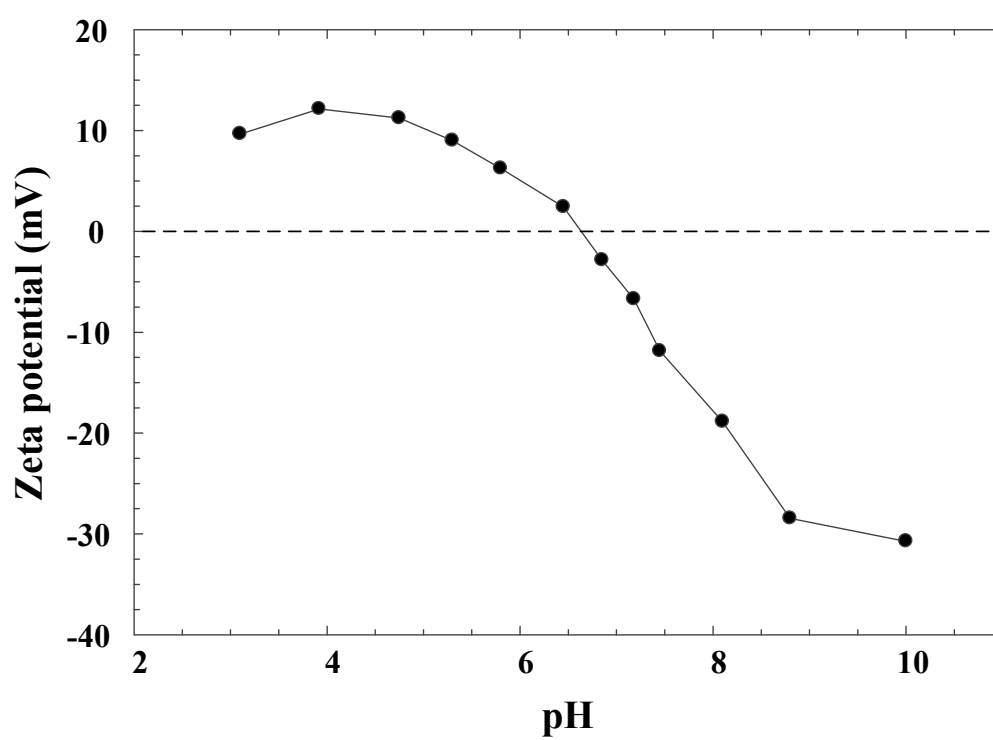


Fig. S7. Zeta potential of 1 wt% ZnFe₂O₄-TiO₂ as a function of pH value

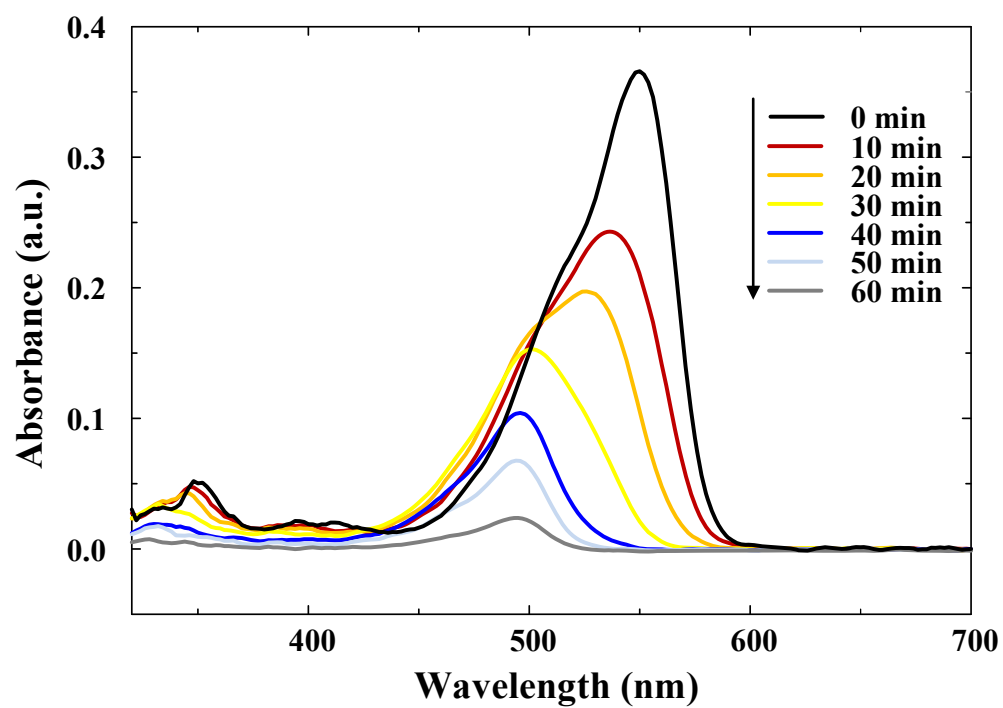


Fig. S8. UV-Vis absorption spectra of RhB in the presence of 1 wt% ZnFe₂O₄-TiO₂ at different photocatalytic degradation times.

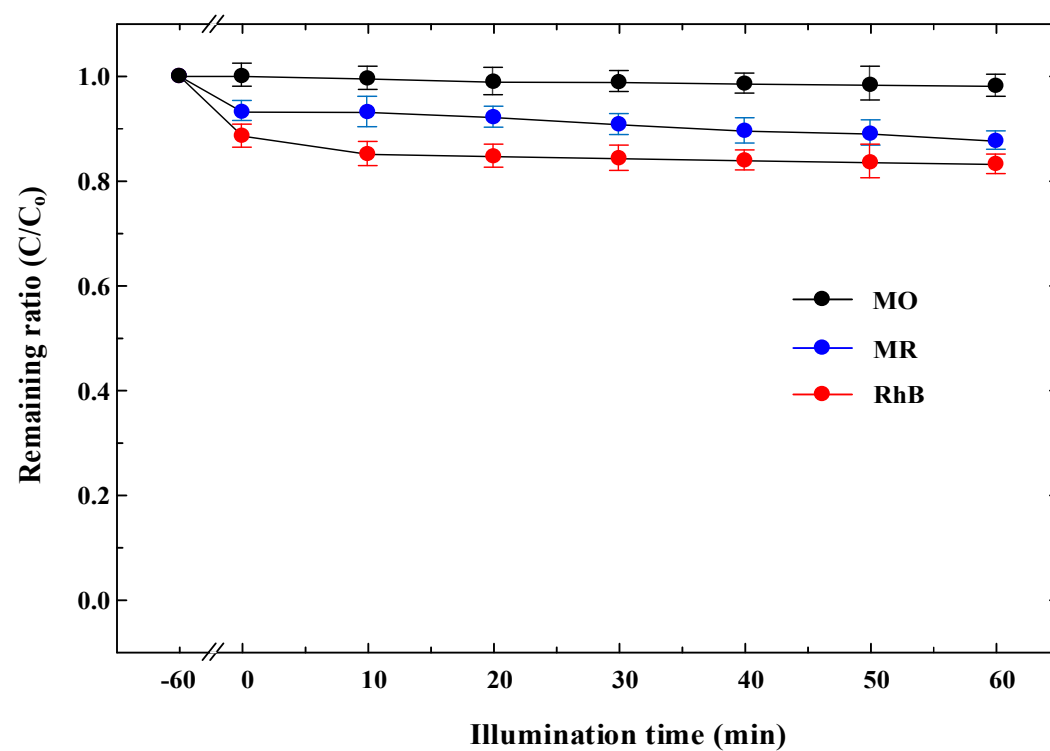


Fig. S9. Photocatalytic activity of ZnFe_2O_4 toward dyes decomposition.

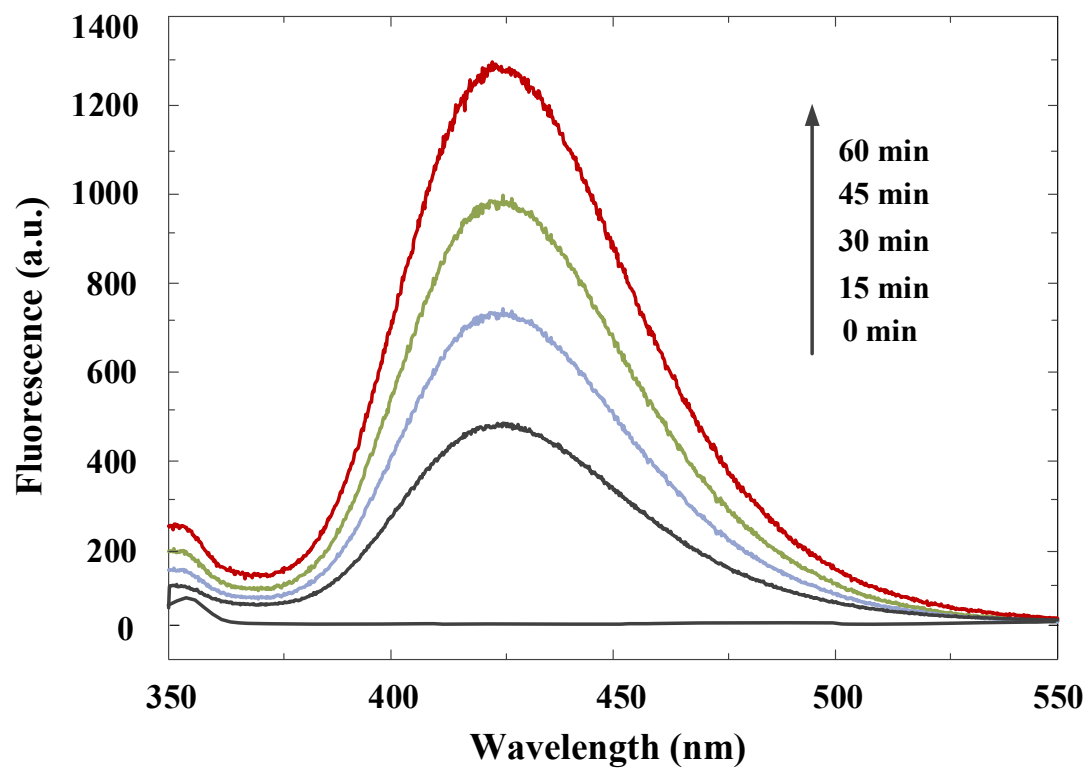


Fig. S10. Change in PL spectra with irradiation time for 1 wt% ZnFe₂O₄-TiO₂ in a 5×10^{-4} M basic solution of terephthalic acid (excitation at 315 nm) under visible light irradiation.