

Supplementary Information

Sustainable Photocatalytic Removal of Cosmetic Preservatives Parabens over a Dual S-Scheme $\text{TiO}_2/\text{ZnO}/\text{g-C}_3\text{N}_4$ Catalyst under Natural Sunlight Irradiation

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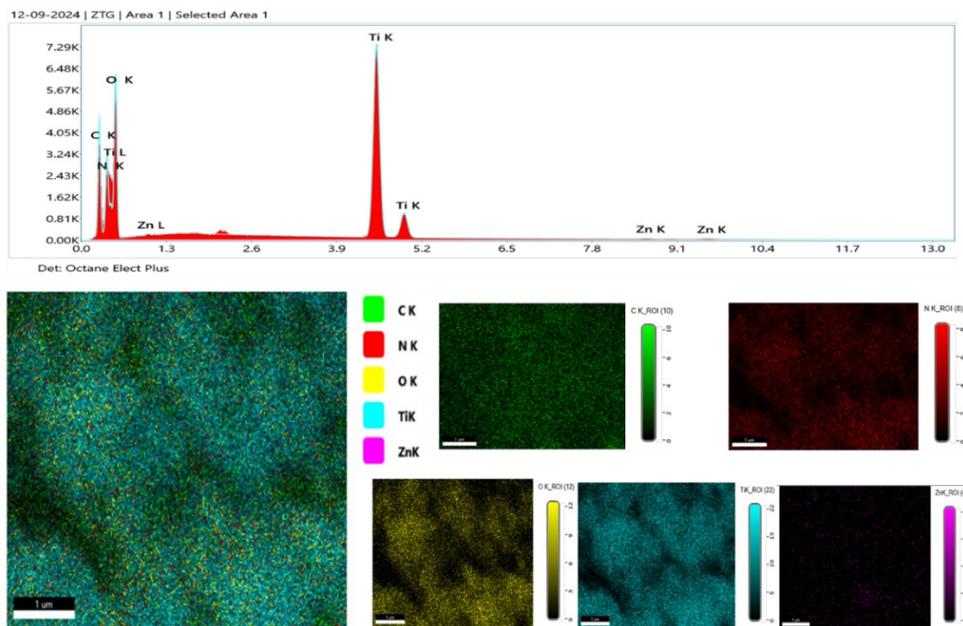


Fig. S1: EDAX pattern of TZC heterojunction

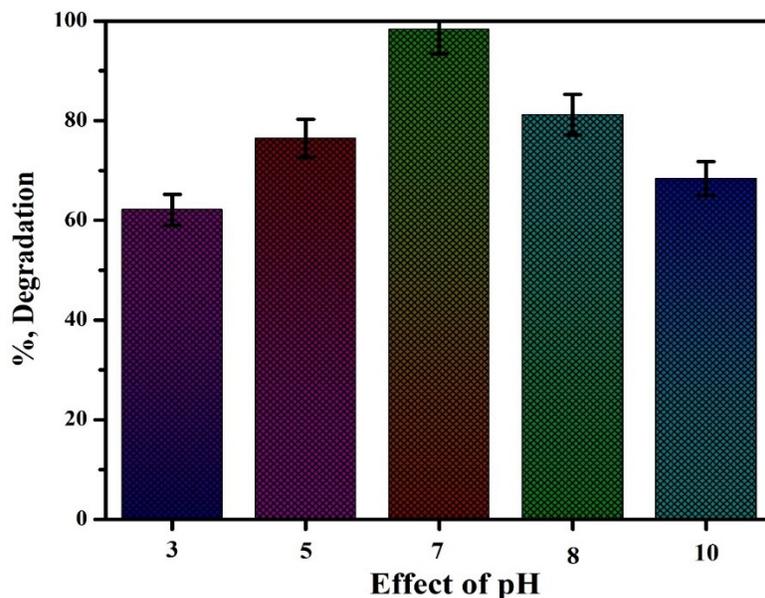


Fig. S2: Effect of pH variation on the degradation efficiency of EtP

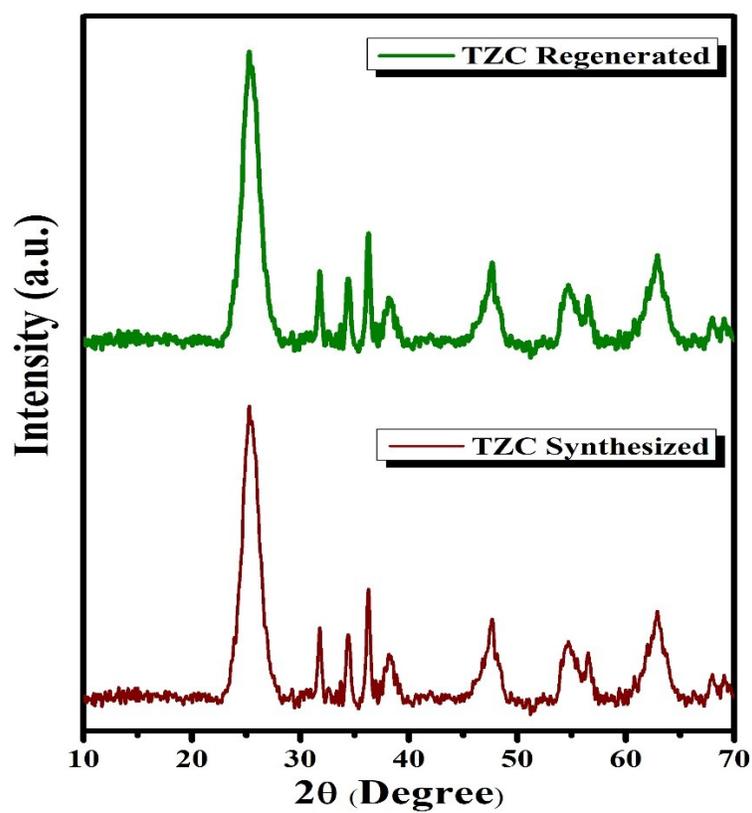


Fig. S3: XRD analysis of TZC synthesized and regenerated

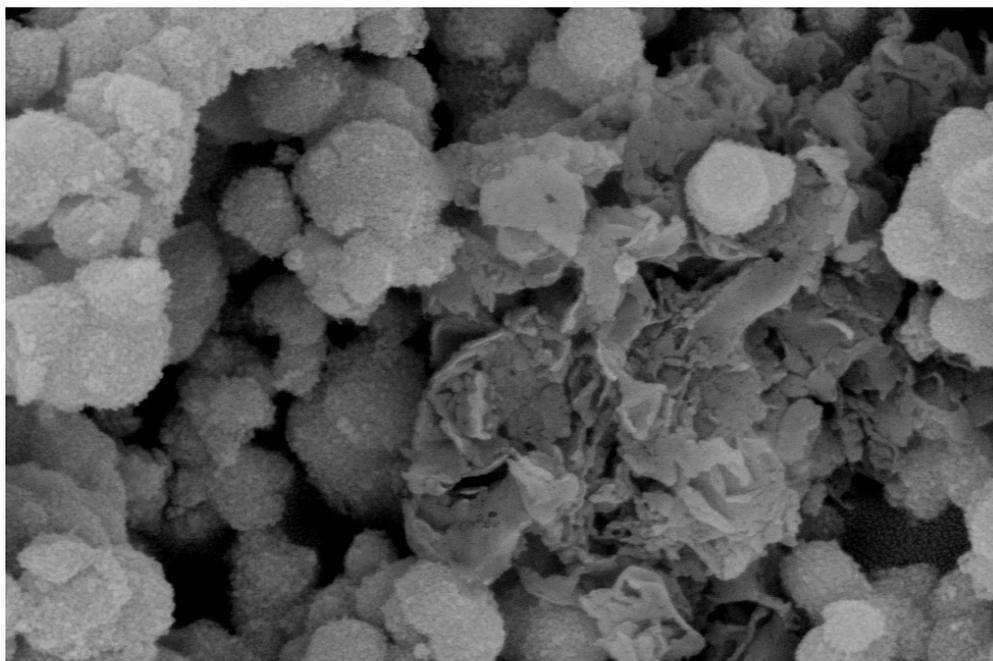


Fig. S4: FE SEM analysis of regenerated TZC

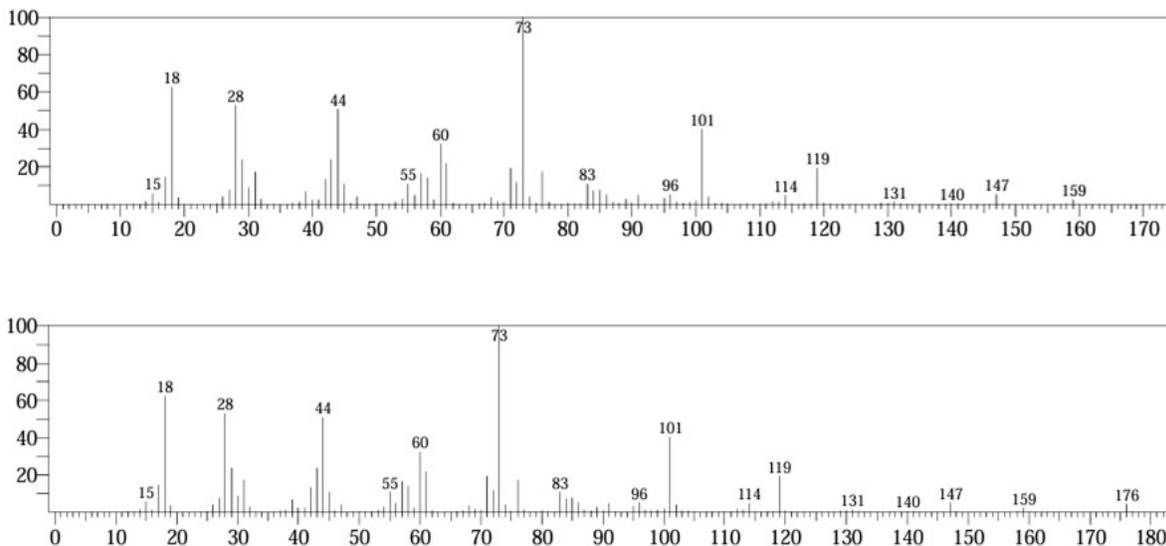


Fig. S5: LCMS spectra of EtP degradation

Mott-Schottky analysis:

The Mott-Schottky analysis of ZnO, TiO₂, and g-C₃N₄ was conducted to determine their flat-band potentials and semiconductor characteristics (**Fig. S6**). The positive slopes of the $1/C^2$ vs. potential plots for all materials confirm their n-type nature. The flat-band potential of ZnO was found to be approximately -0.75 V vs. SHE, while that of TiO₂ was around -0.69 V vs. SHE. The flat-band potential of g-C₃N₄ was -1.28 V. When converted to a common reference scale, these values indicate that the conduction band of g-C₃N₄ is more negative than ZnO and TiO₂, whereas the valence band of ZnO and TiO₂ is more positive, fulfilling the energetic requirements for S-scheme operation. Upon contact, an internal electric field (IEF) forms at the interface, directing electrons from g-C₃N₄ to ZnO and TiO₂ while the holes in the opposite direction. Under light irradiation, photogenerated electrons in the conduction band of ZnO and TiO₂ recombine with holes in the valence band of g-C₃N₄. This retains the highly reductive electrons in g-C₃N₄ along with highly oxidative holes in ZnO and TiO₂, enhancing the redox ability. Thus, the Mott-Schottky results

provide strong evidence supporting the formation of a dual S-scheme heterojunction, which facilitates efficient charge separation and promotes photocatalytic degradation of EtP.

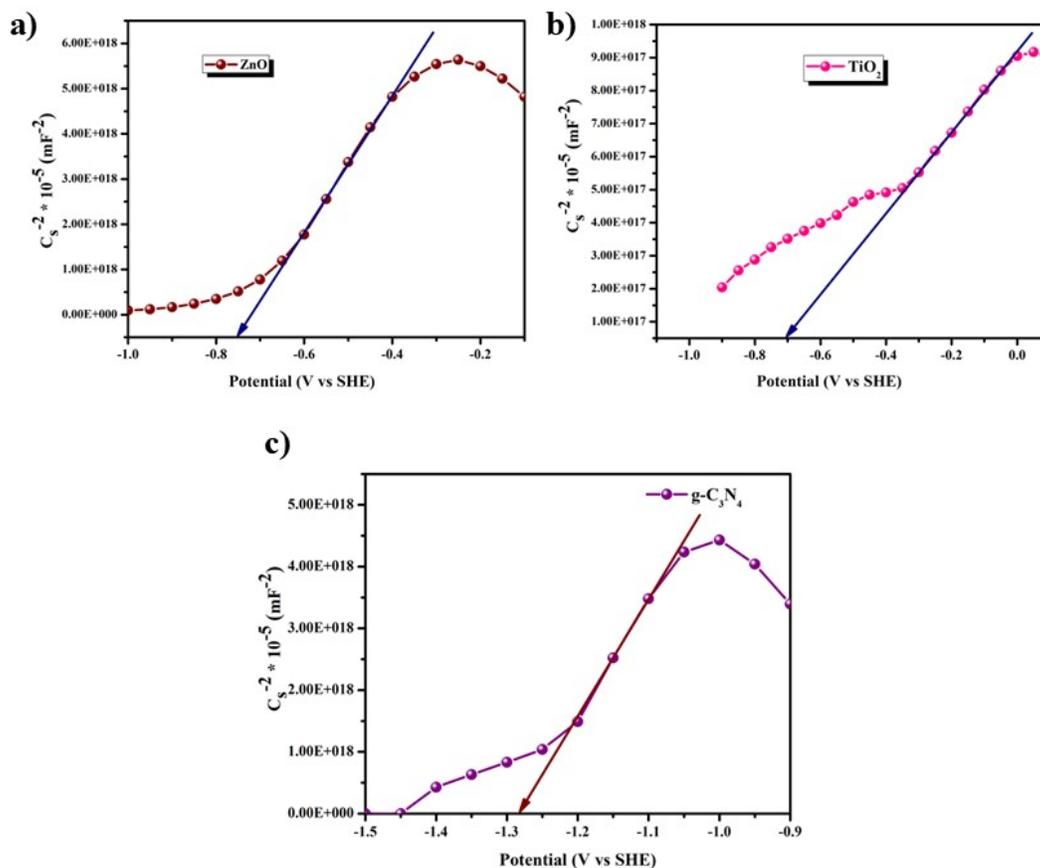


Fig. S6: Mott-Schottky analysis of ZnO, TiO₂, g-C₃N₄ individually

Entry	Catalyst	Catalyst Dosage (gL ⁻¹)	EtP (mgL ⁻¹)	Source; duration; pH	Degradation, %	Ref.
1	CuOx/BiVO ₄	0.5	0.5	Sunlight/1h	98	1
2	Ag ₃ PO ₄	0.5	1.5	Sunlight/1h	90	2
3	Bi ₂ Ti ₂ O ₇ /TiO ₂	1	10	500 W UV/30 min	91	3
4	rGO/TiO ₂	0.7 L ⁻¹	15	UV /40 min	98	4
5	MI-Ag-TiO ₂	0.3	10	500 W UV/ 2 h	93	5
6	Ag/BiVO ₄	0.5	10	40 W LED/ 4 h	94	6
7	TiO ₂ /ZnO/g-C ₃ N ₄	1 gL ⁻¹	10	Sunlight/1.5 h/7	99	Present work

Table TS1: Comparative table for the photocatalytic degradation of EtP

References

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