

Supplementary Information

In situ growth of heterojunction cocatalyst on g-C₃N₅ surface enhances charge transfer to improve photocatalytic activity

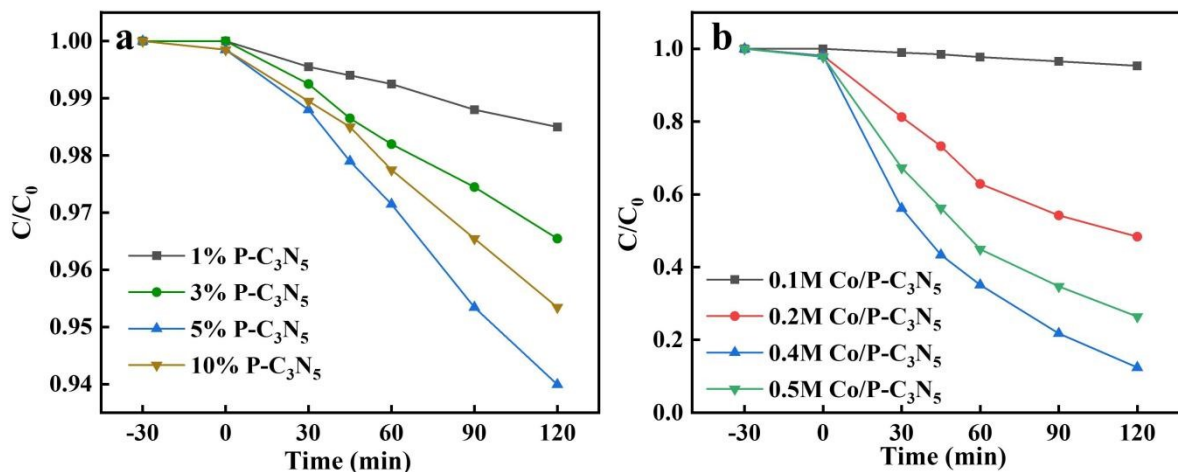


Figure.S1. (a)Photocatalytic degradation of TC under visible light (30 min in the dark) by different P contents of P-C₃N₅, and (b) by different Co contents of CoOOH·CoO_x/P-C₃N₅

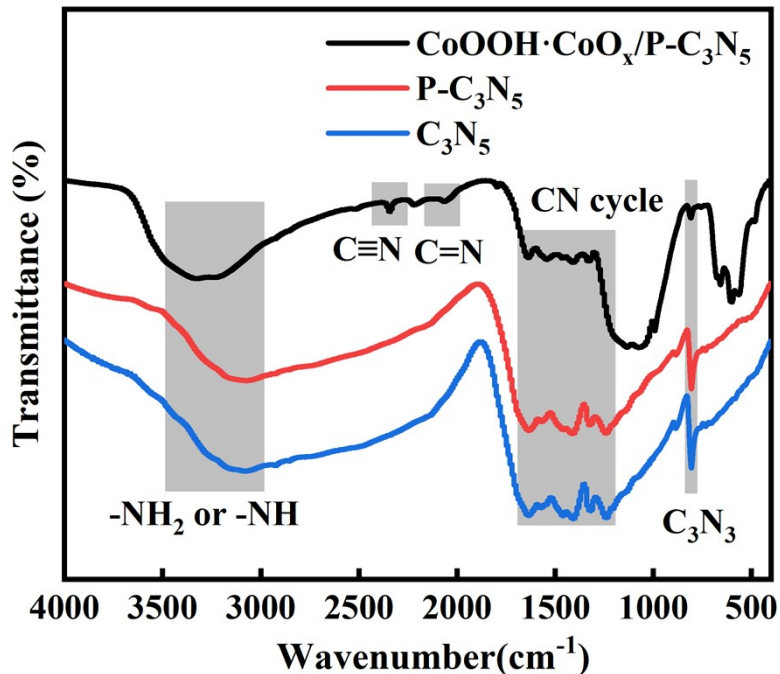


Figure.S2. FTIR of C₃N₅, P-C₃N₅ and CoOOH·CoO_x/P-C₃N₅

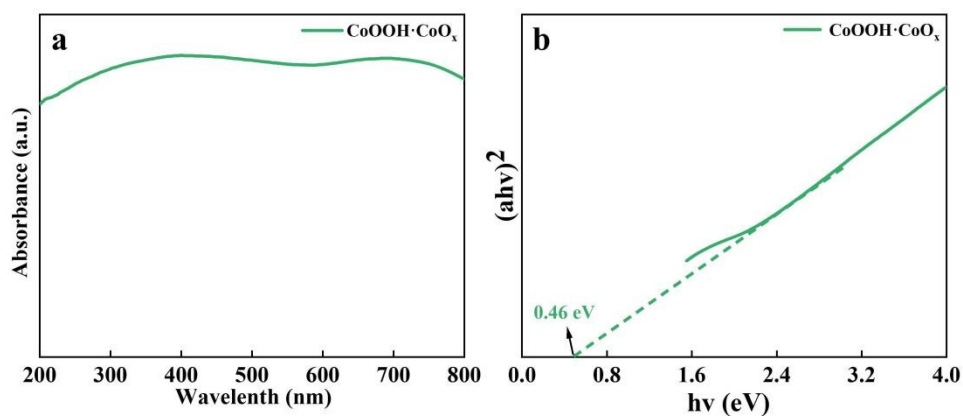


Figure. S3.(a) UV-vis absorption spectra and (b) $(ah\nu)^2$ versus $h\nu$ curves of the $\text{CoOOH}\cdot\text{CoO}_x$.

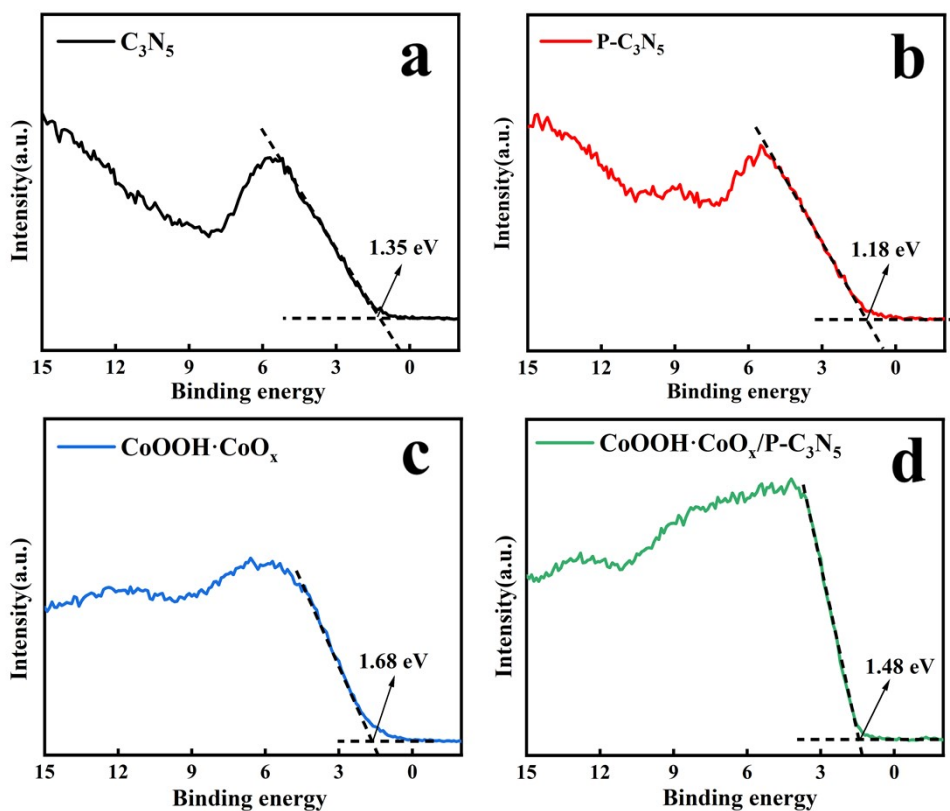


Figure. S4. XPS Vb of (a) C_3N_5 , (b) $\text{P-C}_3\text{N}_5$, (c) $\text{CoOOH}\cdot\text{CoO}_x$ and (d) $\text{CoOOH}\cdot\text{CoO}_x/\text{P-C}_3\text{N}_5$

As shown in Figure S3 and Figure S4, the $\text{CoOOH}\cdot\text{CoO}_x$ in these figures is not in-situ growth on the surface of $\text{P-C}_3\text{N}_5$, which may be different from the $\text{CoOOH}\cdot\text{CoO}_x$ in the proposed material, but it is still of certain reference value.

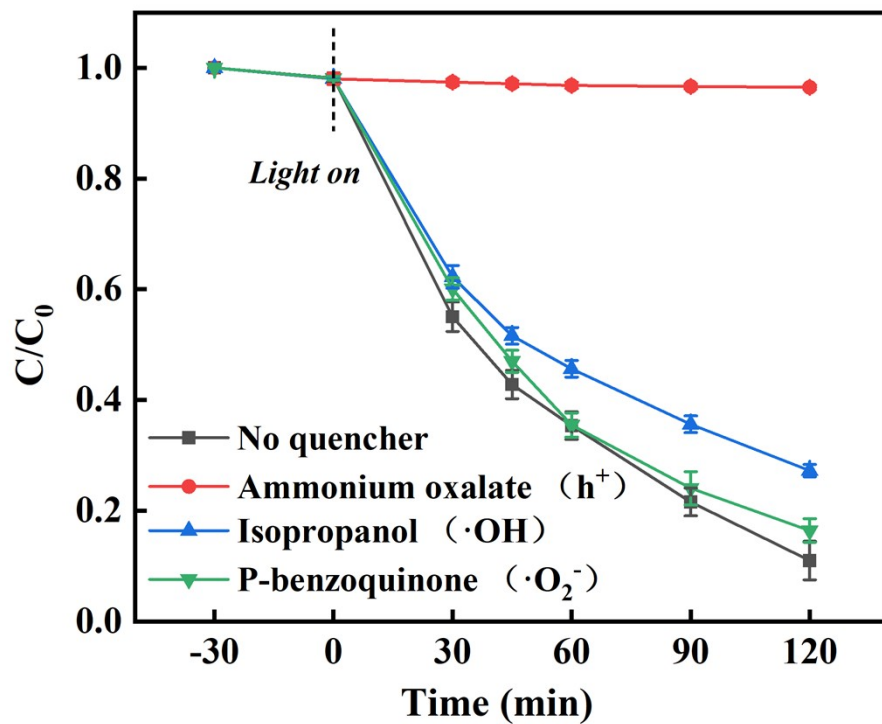


Figure.S5. Masking experiment of TC degradation by $\text{CoOOH}\cdot\text{CoO}_x/\text{P-C}_3\text{N}_5$

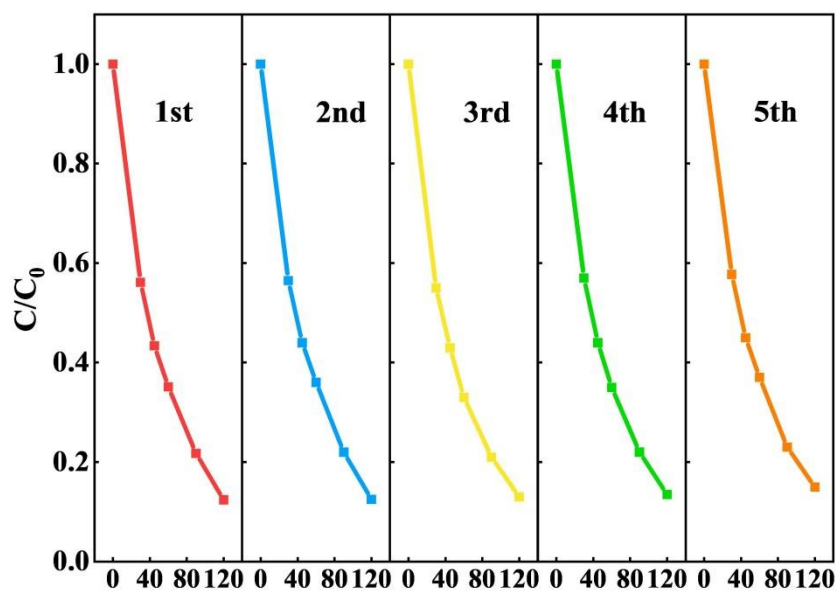


Figure.S6. Cyclic experiment of TC degradation by $\text{CoOOH}\cdot\text{CoO}_x/\text{P-C}_3\text{N}_5$

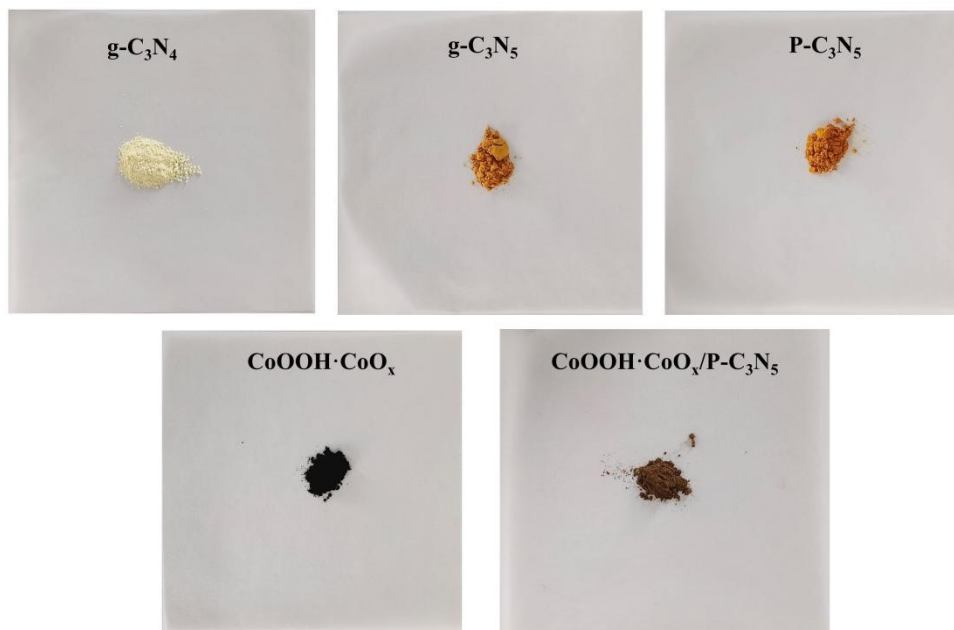


Figure.S7. Photos of $g-C_3N_4$, $g-C_3N_5$, $P-C_3N_5$, $CoOOH \cdot CoO_x$ and $CoOOH \cdot CoO_x/P-C_3N_5$

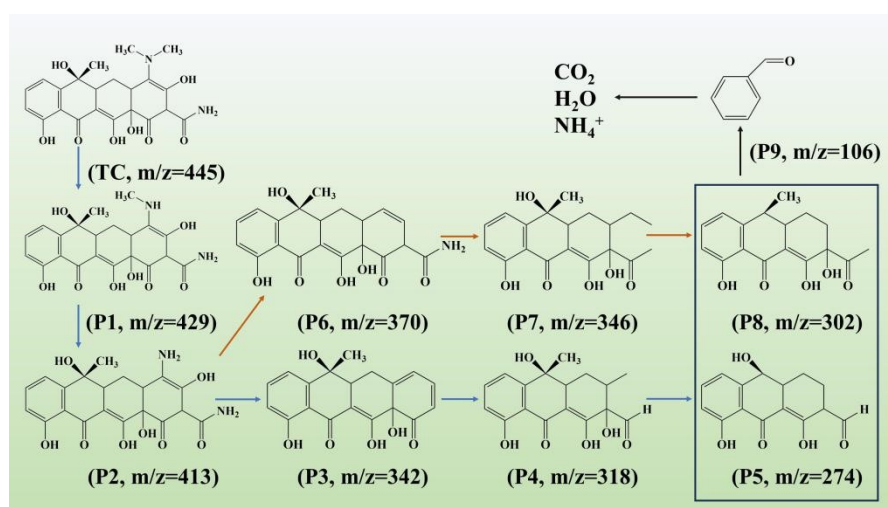


Figure.S8. The proposed photocatalytic-degradation pathways of TC

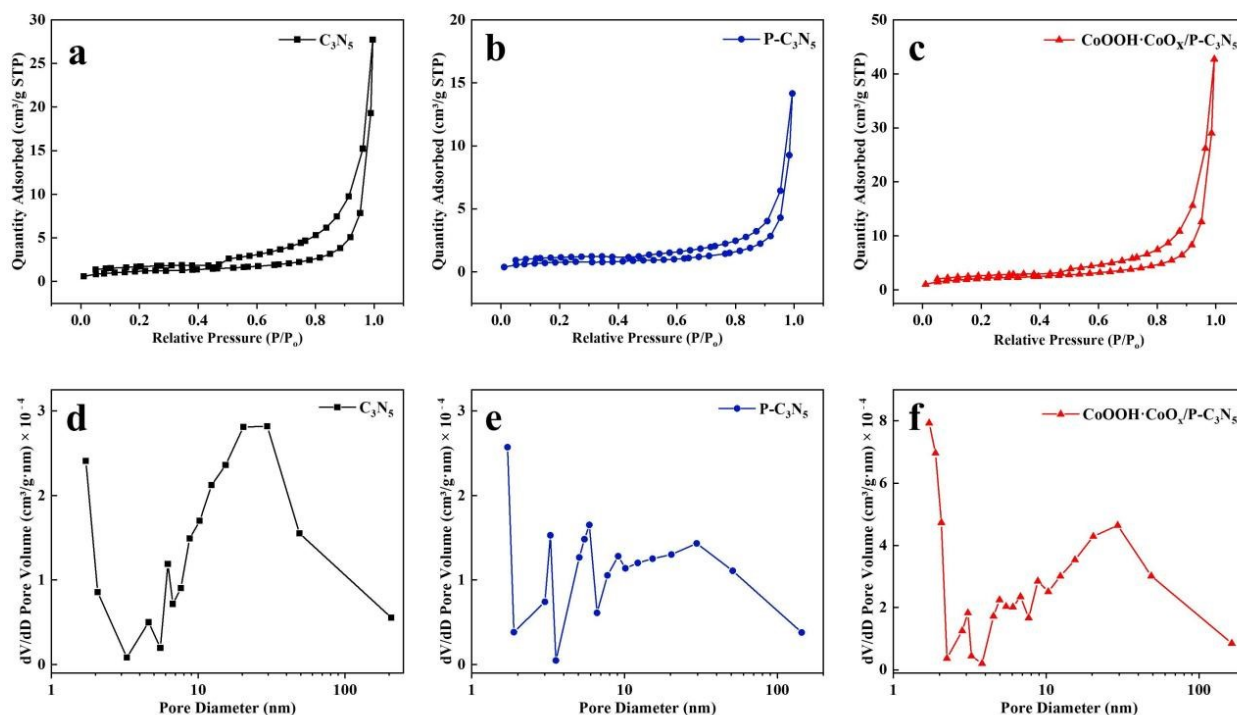


Figure.S9. N₂ adsorption–desorption isotherms of C₃N₅ (a), P-C₃N₅ (b) and CoOOH·CoO_x/P-C₃N₅ (c), the pore-size distribution curves for C₃N₅ (d), P-C₃N₅ (e) and CoOOH·CoO_x/P-C₃N₅ (f).

As shown in Fig. S9, nitrogen adsorption–desorption is used to investigate the pore structure and surface areas of C₃N₅, P-C₃N₅ and CoOOH·CoO_x/P-C₃N₅. All the samples display type IV curves, indicating the presence of mesopores [1]. The shape of the hysteresis loops for the three samples is categorized as type H3, which implies the formation of slit-like pores owing to the aggregation of the sheet-like particles [1]. The CoOOH·CoO_x/P-C₃N₅ sample contains small mesopores (3–4 nm) and large mesopores (20–50 nm). As shown in Table S1, the BET surface areas (S_{BET}) and pore volume of CoOOH·CoO_x/P-C₃N₅ are the highest, which is consistent with the results of SEM. This mesoporous hierarchical architecture could provide a transport path for the reactants and products and offers more active sites and enhance photo-energy harvesting in photocatalysis [1–4].

Table S1. Summary of the S_{BET} , pore volume, and peak pore size.

| Samples | S_{BET} (m ² /g) | Pore volume (cm ³ /g) | Peak pore size (nm) |
|---|--------------------------------------|----------------------------------|---------------------|
| C ₃ N ₅ | 4.3 | 0.043 | 6.4/23.5 |
| P-C ₃ N ₅ | 2.8 | 0.022 | 5.9/27.3 |
| CoOOH·CoO _x /P-C ₃ N ₅ | 7.6 | 0.066 | 8.6/29.6 |

Reference

- [1] J. Zhang, J. Yu, Y. Zhang, Q. Li, J.R. Gong, Visible Light Photocatalytic H₂-Production Activity of CuS/ZnS Porous Nanosheets Based on Photoinduced Interfacial Charge Transfer, *Nano Lett.* 11 (2011) 4774–4779. <https://doi.org/10.1021/nl202587b>.
- [2] G. Li, D. Zhang, J.C. Yu, M.K.H. Leung, An Efficient Bismuth Tungstate Visible-Light-Driven Photocatalyst for Breaking Down Nitric Oxide, *Environ. Sci. Technol.* 44 (2010) 4276–4281. <https://doi.org/10.1021/es100084a>.
- [3] X. Yu, J. Yu, B. Cheng, M. Jaroniec, Synthesis of Hierarchical Flower-like AlOOH and TiO₂/AlOOH Superstructures and their Enhanced Photocatalytic Properties, *J. Phys. Chem. C* 113 (2009) 17527–17535. <https://doi.org/10.1021/jp906992r>.
- [4] W. Shao, F. Gu, L. Gai, C. Li, Planar scattering from hierarchical anatase TiO₂ nanoplates with variable shells to improve light harvesting in dye-sensitized solar cells, *Chem. Commun.* 47 (2011) 5046. <https://doi.org/10.1039/c1cc10548e>.