## Hollow $In_2O_3$ (a) $ZnFe_2O_4$ heterojunctions for highly efficient photocatalytic degradation of tetracycline under visible light

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## **Supporting Information**



Fig. S1. XRD pattern of In-MIL-68 prisms.



**Fig. S2.** TGA curve of In-MIL-68 prisms in air atmosphere with a heating rate of 5 °C/min.



Fig. S3.  $N_2$  sorption isotherms and BET surface area of  $In_2O_3$ . Inset is the corresponding pore size distribution curve.



Fig. S4. EDX spectrum of In<sub>2</sub>O<sub>3</sub>.



Fig. S5. TGA curve of ZnFe-LDH in air atmosphere with a heating rate of 5 °C/min.



Fig. S6. TEM images of (a) ZnFe-LDH and (b) ZnFe<sub>2</sub>O<sub>4</sub>.



Fig. S7. EDX spectrum of  $In_2O_3$ @ZnFe<sub>2</sub>O<sub>4</sub>-500s.



Fig. S8. SEM images of (a, b) In<sub>2</sub>O<sub>3</sub>@ZnFe<sub>2</sub>O<sub>4</sub>-400s, (c, d) In<sub>2</sub>O<sub>3</sub>@ZnFe<sub>2</sub>O<sub>4</sub>-600s.



Fig. S9.  $N_2$  sorption isotherms and BET surface area of  $ZnFe_2O_4$ . Inset is the corresponding pore size distribution curve.



Fig. S10. N<sub>2</sub> sorption isotherms and BET surface area of  $In_2O_3$ @ZnFe<sub>2</sub>O<sub>4</sub>-500s. Inset is the corresponding pore size distribution curve.



Fig. S11. (a) Tauc plots and (b) Valence band XPS spectra of  $In_2O_3$  and  $ZnFe_2O_4$ .

Table S1.	The zeta potential	and the maximum	UV-vis absorption	peak (λ <sub>max</sub> )	of TC
	p			P • • • • • • • • • • • • • • • • • • •	

under	different	nH
unuor	uniterent	pm.

рН	3	5	7	9	11
zeta potential (mV)	12.1	14.7	-1.4	-9.3	-36.2
$\lambda_{max}$ (nm)	357	357	357	363	377

Photocatalysts	Pollutant	Concentration	Reaction time	Catalyst dosage	Removal rate (%)	Ref.	
		(mg/L)	(min)	(min) (g/L)			
In <sub>2</sub> O <sub>3</sub> @ZnFe <sub>2</sub> O <sub>4</sub>	TC	150	60	0.5	90	Our work	
Carbon dots/MoO <sub>3</sub> /g-C <sub>3</sub> N <sub>4</sub>	TC	20	90	0.6	88.4	1	
Ag@g-C <sub>3</sub> N <sub>4</sub> @BiVO <sub>4</sub>	TC	20	60	0.3	82.75	2	
Ag <sub>3</sub> PO <sub>4</sub> /CuBi <sub>2</sub> O <sub>4</sub>	TC	20	60	0.5	75	3	
type II AgI/CuBi <sub>2</sub> O <sub>4</sub>	TC	10	60	0.5	80	4	
Z-scheme AgBr/CuBi <sub>2</sub> O <sub>4</sub>	TC	10	60	0.5	90	4	
Ag/AgCl/Bi2MoO6	Rhb	10	60	1.0	90.9	5	

Table S2. Comparison of different contaminants degradation time and efficiency for In<sub>2</sub>O<sub>3</sub>@ZnFe<sub>2</sub>O<sub>4</sub> with previously reported catalysts.



Fig. S12. LC-MS spectra at 0 min (a), 20 min (b), 40 min (c) and 60 min (d).



Fig. S13. TC photocatalytic degradation pathway on In<sub>2</sub>O<sub>3</sub>@ZnFe<sub>2</sub>O<sub>4</sub>.



**Fig. S14.** (a). TEM image of  $In_2O_3@ZnFe_2O_4$ -500s after photocatalytic process. (b). Cycling runs for the degradation of TC (pH=11, TC=50 mg/L,  $In_2O_3@ZnFe_2O_4$ -500s = 0.5 g/L), (c). XRD patterns and XPS spectra of  $In_2O_3@ZnFe_2O_4$ -500s before and after 10 times photocatalytic reactions: (d) survey , (e) In 3d, (f) Zn 2p, (g) Fe 2p and (h) O 1s, (i). Magnetic hysteresis loop of  $In_2O_3@ZnFe_2O_4$ -500s (insert: the solution before and after magnetic separation).

## References

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