

# Supplementary Information

## A Bifunctional Electroactive Ti<sub>4</sub>O<sub>7</sub>-based Membrane System for Highly Efficient Ammonia Decontamination

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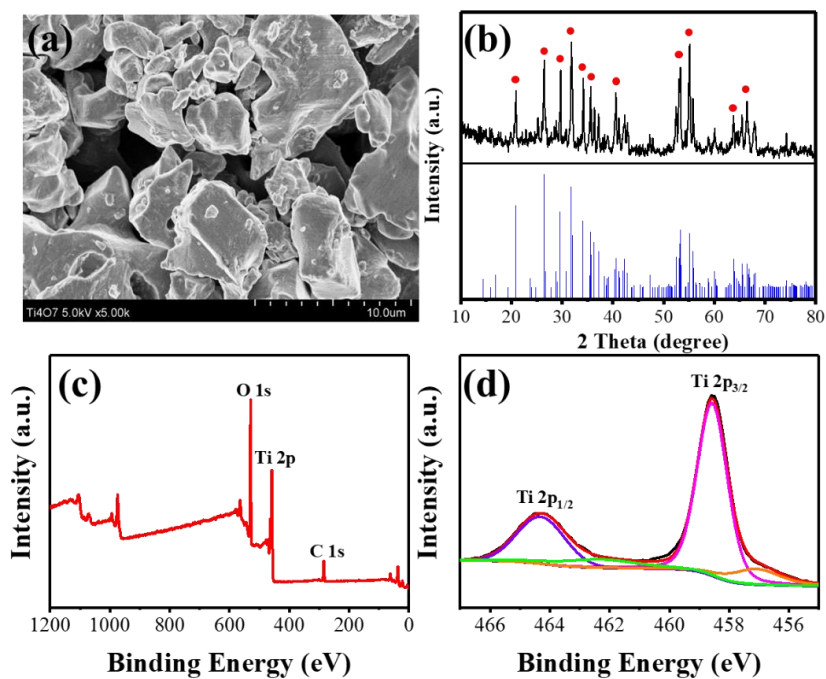
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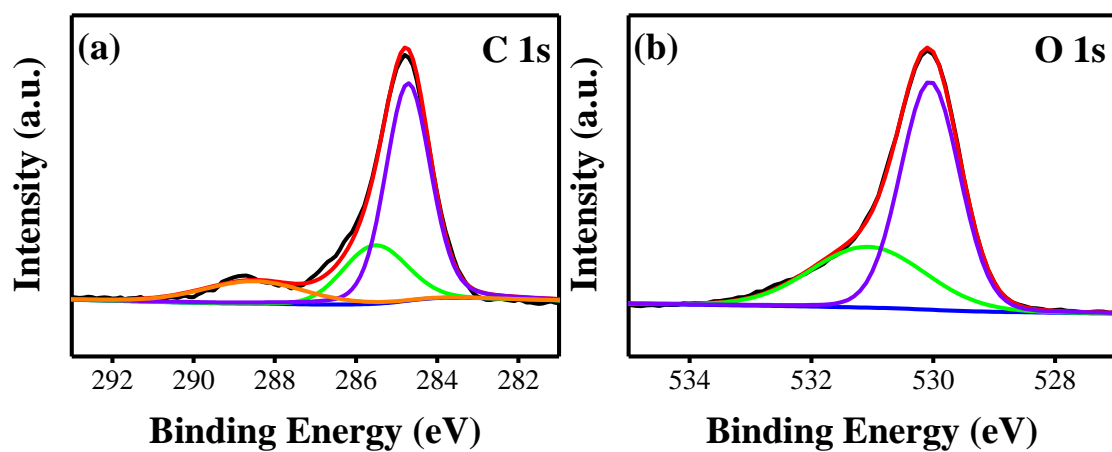
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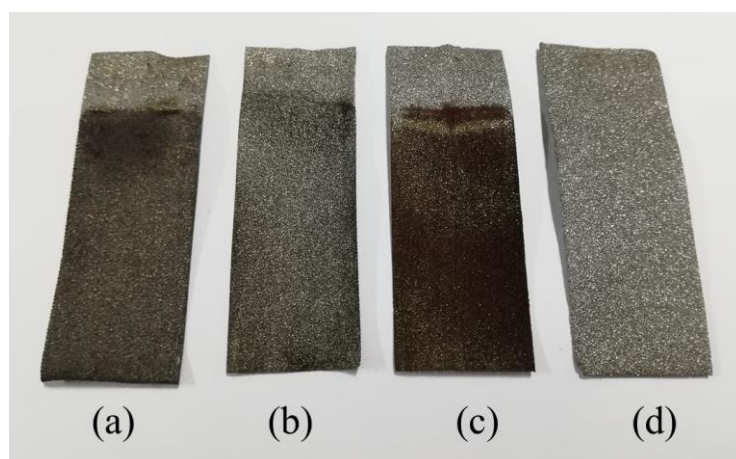
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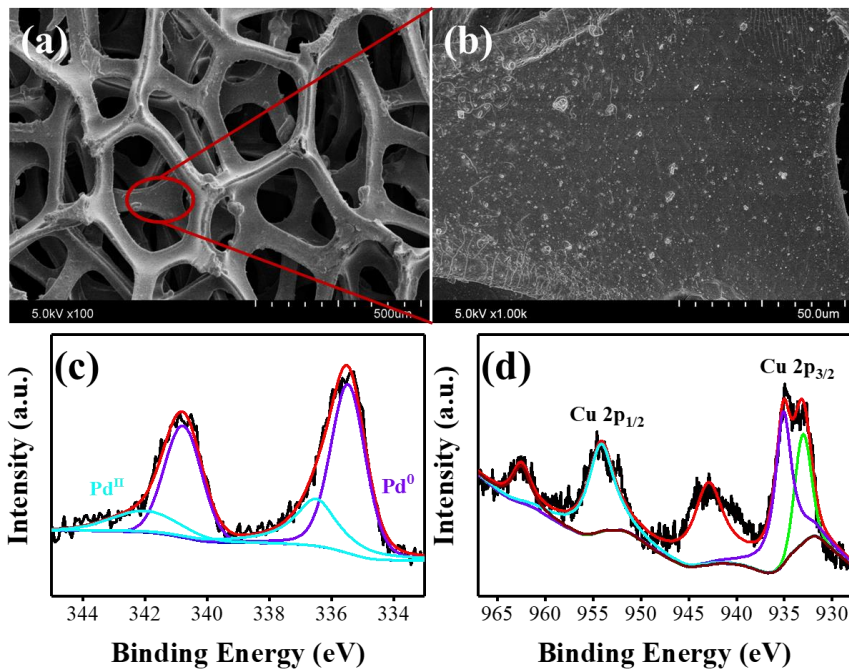
**Figure S1.** (a) FESEM characterization, (b) XRD pattern, and (c) XPS survey pattern of the  $\text{Ti}_4\text{O}_7$  anode. XPS narrow scan of (d) Ti 2p on  $\text{Ti}_4\text{O}_7$  electrode.



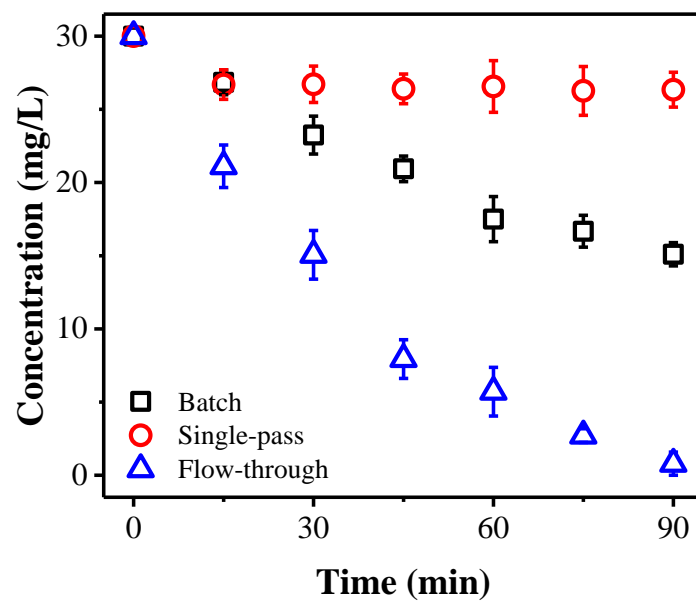
**Figure S2.** XPS spectra for the narrow scan of (a) C 1s, and (b) O 1s on Ti<sub>4</sub>O<sub>7</sub> electrode.



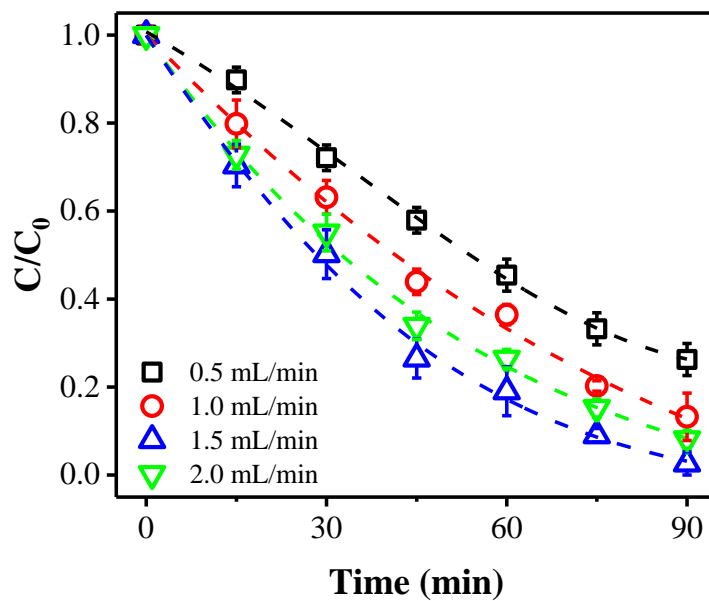
**Figure S3.** Digital pictures of (a) Pd-Cu/NF electrode, (b) Pd/NF electrode, (c) Cu/NF electrode and (d) pure Ni foam electrode.



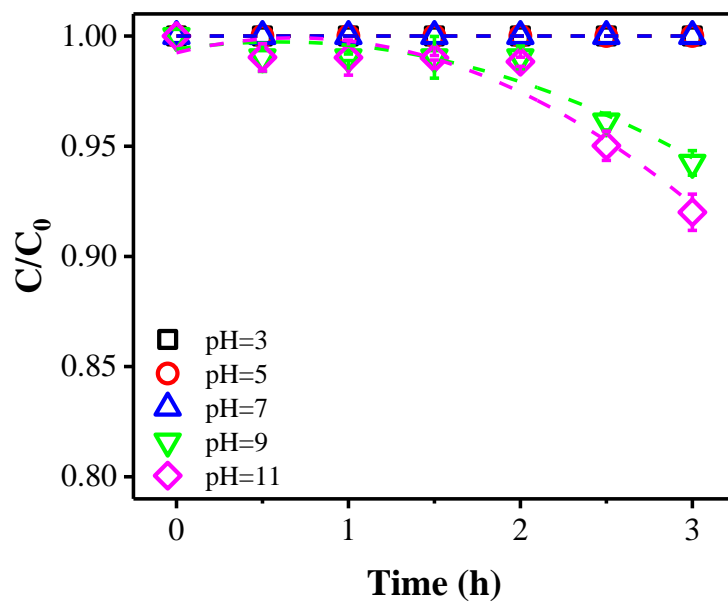
**Figure S4.** FESEM images (a, b) of the Pd/Cu-Ni foam electrode. XPS spectra for the narrow scan of (c) Pd, and (d) Cu of a fresh Pd–Cu/NF cathode.



**Figure S5.** Comparison of ammonia conversion kinetics in batch, single-pass and flow-through systems. Reaction conditions: anode potential of 2.8 V vs. Ag/AgCl, [Cl<sup>-</sup>] concentration of 0.14 M, and pH of 9.

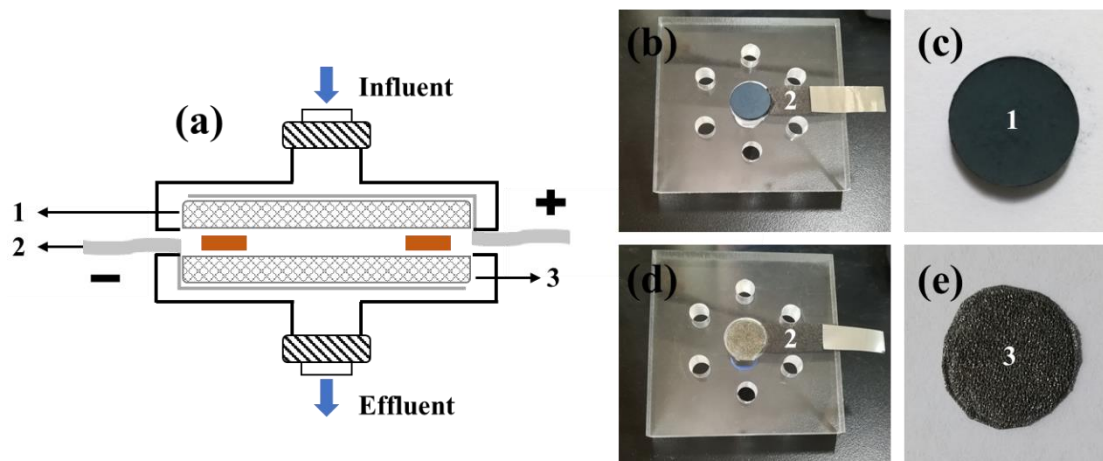


**Figure S6.** The effect of flow rate on ammonia conversion. Reaction conditions: anode potential of 2.8 V vs. Ag/AgCl,  $[Cl^-]$  concentration of 0.14 M, and pH of 9.



**Figure S7.** Conversion effect of ammonia with 0.14 M  $\text{Na}_2\text{SO}_4$  background electrolyte at different anode potentials. Experimental conditions:  $[\text{Cl}^-]$  of 0.14 M, flow rate of  $1.5 \text{ mL}\cdot\text{min}^{-1}$ , and pH of 9.





**Figure S8.** Schematic of the flow-through filtration setup. (a) Modified acrylic filter sleeve, including 1) a  $\text{Ti}_4\text{O}_7$  anode, 2) a Ti plate current collector, and 3) a Pd-Cu/NF cathode. (b, d) Images of the modified filter sleeve. (c, e) Images of the anode and cathode.

**Table S1.** Comparison of the performance of different materials on ammonia conversion

Anode	Ammonia concentration	Experimental conditions	Removal efficiency	References
WO <sub>3</sub>	20 mg L <sup>-1</sup>	Cl <sup>-</sup> /NH <sub>4</sub> <sup>+</sup> = 20, pH = 4, 1.0 V vs. SCE	99.9% (1.5 h)	[1]
SnO <sub>2</sub> -CNT	30 mg L <sup>-1</sup>	2.5 V vs. Ag/AgCl, [Cl <sup>-</sup> ] = 0.1 M, pH = 7 flow rate = 4 mL min <sup>-1</sup>	> 99.9% (1.5 h)	[2]
Ti/Pt	150 mg L <sup>-1</sup>	NaCl = 0.8% (w/v) pH = 9 I = 0.075 A cm <sup>-2</sup>	82% (1 h)	[3]
Ti/RuO <sub>2</sub> -TiO <sub>2</sub>	1060-1380 mg L <sup>-1</sup>	I = 116 mA cm <sup>-2</sup> flow rate = 2000 L h <sup>-1</sup>	49% (3 h)	[4]
Ti <sub>4</sub> O <sub>7</sub>	30 mg L <sup>-1</sup>	2.8 V vs. Ag/AgCl, [Cl <sup>-</sup> ] = 0.14 M, pH = 9 flow rate = 1.5 mL min <sup>-1</sup>	97.4% (1.5 h)	/

## References

- [1] Ji Y.Z.; Bai J.; Li J.H.; Luo T.; Qiao L.; Zeng Q.Y.; Zhou B.X. Highly selective transformation of ammonia nitrogen to N<sub>2</sub> based on a novel solar-driven photoelectrocatalytic-chlorine radical reactions system. *Water Res.* **2017**, *125*, 512-519.
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