

Supporting information

Table S1. A representative summary of the enhanced photocatalytic activity of g-C₃N₄ photocatalysts based on sodium ion treatment for water splitting.

| Precursors | Approach | Sacrificial and agentcocatalysts | Light source | Activity (unit) | ref |
|---|-------------------------------|----------------------------------|--|---|-----|
| Melamine and disodium ethylenediaminetetraacetate | General thermal condensation | Triethanolamine and Pt | 300 W Xe lamp equipped with a UV-cutoff filter ($\lambda > 420$ nm) | H ₂ ; 258.4 $\mu\text{mol h}^{-1}\text{g}^{-1}$ | 1 |
| Melamine and sodium tripolyphosphate | General thermal condensation | Methanol and Pt | 350 W Xe arc lamp | H ₂ ; 3820 $\mu\text{mol h}^{-1}\text{g}^{-1}$ | 2 |
| Melamine and sodium hydroxide | Step calcination | Triethanolamine and Pt | 350 W Xe lamp with a UV cutoff filter ($\lambda > 400$ nm) | H ₂ ; 935 $\mu\text{mol h}^{-1}\text{g}^{-1}$ | 3 |
| Melamine and sodium borohydride | Step calcination | Silver nitrate | 300 W Xe lamp with a UV cutoff filter ($\lambda > 300$ nm) | O ₂ ; 561.2 $\mu\text{mol h}^{-1}\text{g}^{-1}$ | 4 |
| Triazole ring and sodium hydroxide | Condensation reaction | Cobalt Hydroxide | 50 W halogen tungsten lamp irradiation | Degradation of methylene blue ; the maximum monolayer adsorption capacity 35mg/g | 5 |
| Dicyandiamide and sodium hydroxide | Hydrothermal treatment | hydrogen peroxide | 350 W Xe lamp with a UV cutoff filter ($\lambda > 420$ nm) | The photocatalytic oxidation (PCO) of NO ; Promote three times | 6 |
| Melamine and sodium chloride | Ground in planetary ball mill | Pt | 300 W xenon lamp with a 420 nm cutoff filter | H ₂ and O ₂ ; 31.5 $\mu\text{mol h}^{-1}\text{g}^{-1}$, 15.2 $\mu\text{mol h}^{-1}\text{g}^{-1}$ | 7 |
| Melamine and potassium chloride-sodium chloride | Molten salt method | - | 250 W high-pressure sodium lamp with main emission from 400 to 800 nm | H ₂ O ₂ ; 4.6 mmol L ⁻¹ | 8 |
| Melamine and potassium chloride-sodium chloride | Molten salt method | - | 250 W high-pressure sodium lamp with main emission in the range of 400-800 | Degradation of rhodamine B ; RhB degradation rate at 90% | 9 |

| | | | | nm | | |
|-------------------------------|------------------------------|-----------------|--|----|--|----|
| Urea and sodium hydroxide | High temperature calcination | Methanol and Pt | 250 W high-pressure sodium lamp (400 $<\lambda<800$ nm) | | H ₂ and H ₂ O ₂ ; 900 μmol h ⁻¹ g ⁻¹ ,800 μmol g ⁻¹ | 10 |
| Melamine and sodium hydroxide | Thermal polymerization | - | 300 W xenon lamp with a 420 nm cutoff filter | | Tetracycline degradation ; the photocatalytic degradation efficiency is 80.61% | 11 |

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The N_2 adsorption and desorption isotherms of the catalyst after solution regulation are shown in the figure S1. All samples show type IV adsorption desorption isotherms with H_3 hysteresis ring, indicating that the existence of mesoporous structure is the result of the accumulation of flake particles.

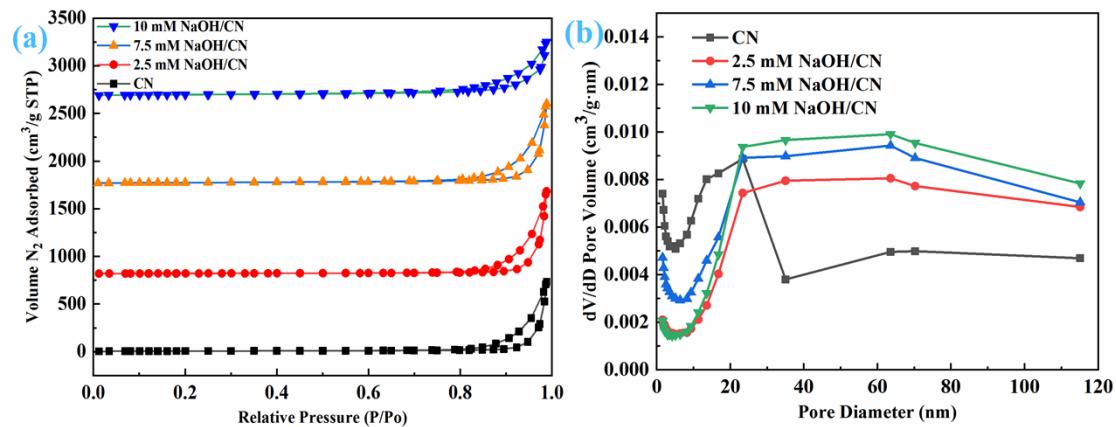
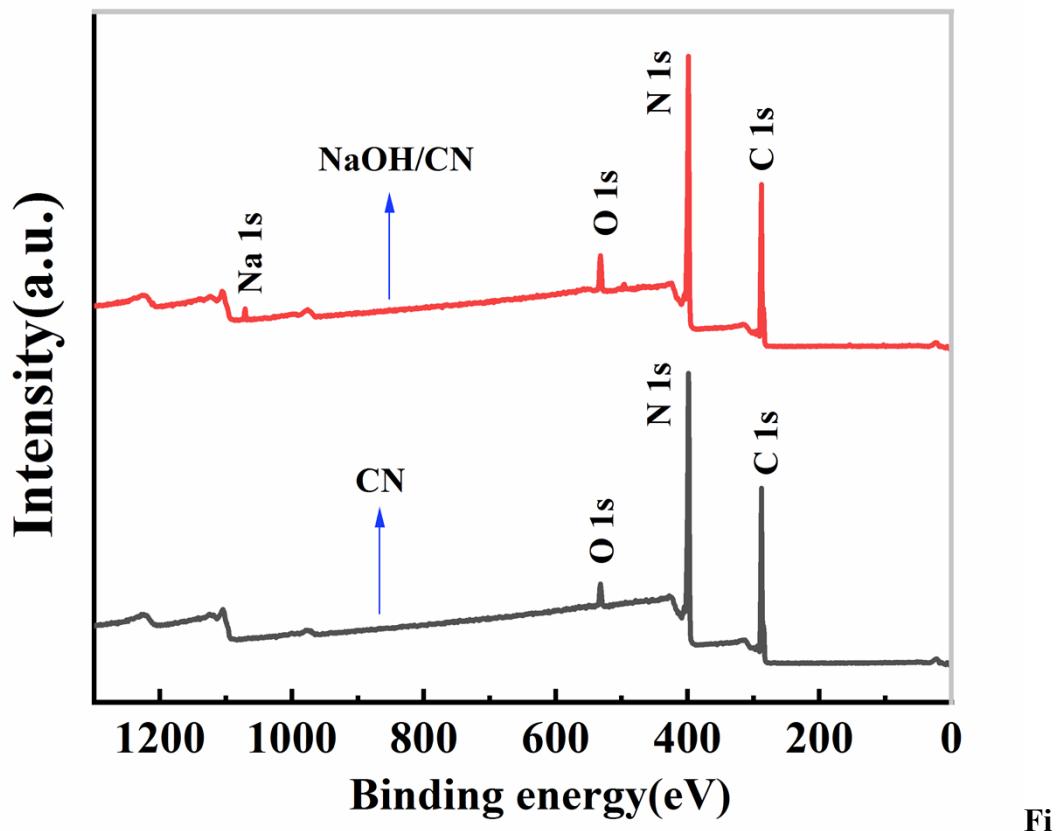


Fig. S1 (a) Nitrogen adsorption-desorption isotherms of CN, 2.5 mM NaOH/CN, 7.5 mM NaOH/CN and 10 mM NaOH/CN; (b) The pore size distribution of CN, 2.5 mM NaOH/CN, 7.5 mM NaOH/CN and 10 mM NaOH/CN.



g. S2 XPS fully scanned spectrum of CN and 7.5 mM NaOH/CN.

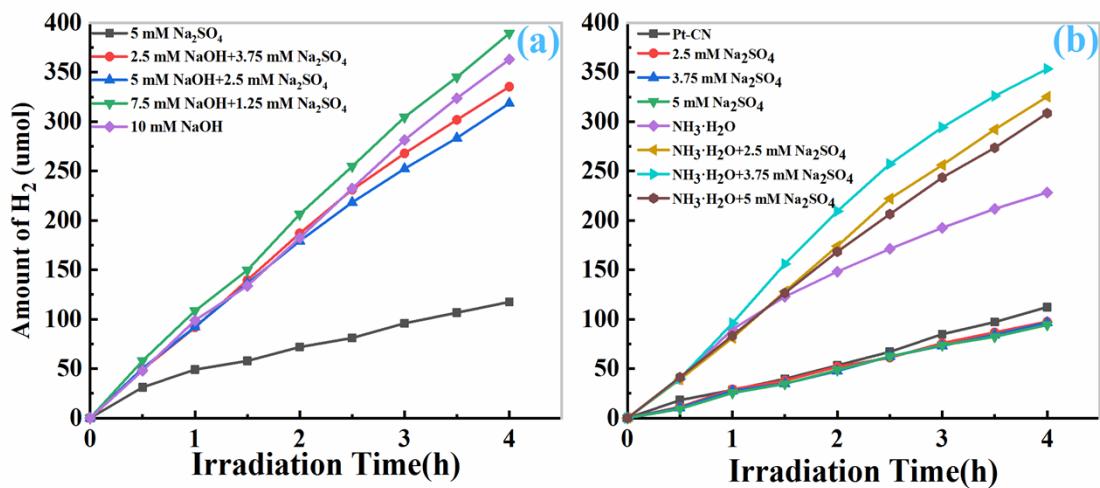


Fig. S3 (a) The time courses of H_2 evolution at different pH and (b) the H_2 production of different Na^+ concentration.

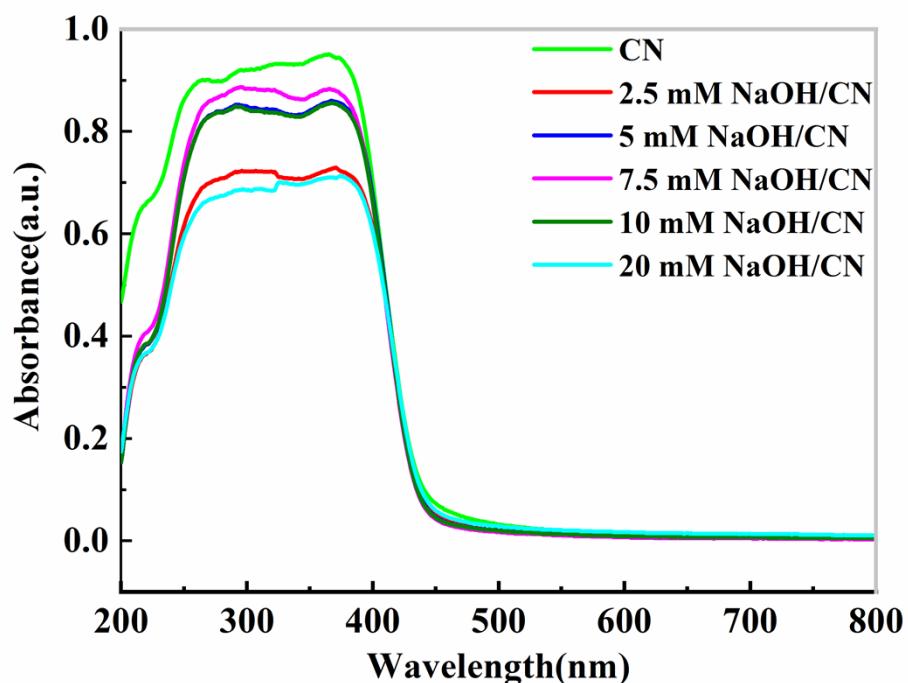


Fig. S4 UV-vis DRS patterns of CN and $(\text{NaOH})_n/\text{CN}$ ($n = 2.5, 5, 7.5, 10, 20 \text{ mM}$)