

## In-situ N-doped TiO<sub>2</sub>/mesoporous g-C<sub>3</sub>N<sub>4</sub> nanosheets S-scheme heterojunction derived from supramolecular precursor with enhanced visible-light photocatalytic performance

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### ABSTRACT

The N-doped TiO<sub>2</sub>/g-C<sub>3</sub>N<sub>4</sub> S-scheme heterojunction with holy nanosheet framework has been synthesized by a simple pyrolysis process of amorphous TiO<sub>2</sub> and melamine-cyanurate complex. During the formation of heterojunction photocatalysts, the interaction of two precursors not only provided in-situ nitrogen doping in TiO<sub>2</sub>, but also construct S-scheme charge transfer channel. Also, N-TiO<sub>2</sub>/g-C<sub>3</sub>N<sub>4</sub> exhibited a porous nanosheet structure due to the co-crystallization of TiO<sub>2</sub> and g-C<sub>3</sub>N<sub>4</sub> at 450°C. The enhancement in narrower bandgap and photogenerated charge carrier separation resulted in superior photocatalytic activities with the highest degradation rate of Methylene blue was recorded by MCN/T-50, which was 2.3 and 3.7 times that of bulk g-C<sub>3</sub>N<sub>4</sub> and pristine TiO<sub>2</sub>. The possible mechanism for the enhanced photocatalytic performance is proposed in this study. The doped S-scheme heterojunction materials prepared from the co-crystallization of precursors in calcination give a broad prospect for the future design of a highly efficient visible-light-driven photocatalyst.

**Keywords:** N-doped TiO<sub>2</sub>; Mesoporous g-C<sub>3</sub>N<sub>4</sub>; Melamine-cyanurate complex; S-scheme heterojunction; Degradation

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