

**A study on the influence of chemical oxidation on Pt containing graphitic carbon nitride photocatalysts in Photocatalytic Hydrogen production**

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In this study, we prepared Pt-containing graphitic carbon nitride (g-C<sub>3</sub>N<sub>4</sub>) catalysts modified by a simple chemical oxidation of g-C<sub>3</sub>N<sub>4</sub> and applied them for photocatalytic hydrogen evolution tests for the first time. The hydrogen production rates of the chemically oxidative Pt/g-C<sub>3</sub>N<sub>4</sub> photocatalysts were at least five times as high as those of the bulk Pt/g-C<sub>3</sub>N<sub>4</sub>. The chemical oxidation of g-C<sub>3</sub>N<sub>4</sub> introduced the oxygen-containing functional groups on the tri-s-triazine units, resulting in more negatively charged surface of g-C<sub>3</sub>N<sub>4</sub>. During the Pt photodeposition on the g-C<sub>3</sub>N<sub>4</sub> surface, the chemically oxidative g-C<sub>3</sub>N<sub>4</sub> with more negatively charged surface and the functional groups not only promoted the agglomeration of Pt nanoparticles on the g-C<sub>3</sub>N<sub>4</sub> surface but also maintained the high ratio of Pt<sup>2+</sup>/Pt<sup>0</sup> for the Pt nanoparticles, which enhanced the hydrogen evolution rate by suppressing the reversible reaction route of H<sub>2</sub> to 2H<sup>+</sup>. In addition, the presence of the oxygen-containing functional groups on the chemically oxidative g-C<sub>3</sub>N<sub>4</sub> increased the separation efficiency of photo-excited charges over Pt/g-C<sub>3</sub>N<sub>4</sub>.