

Photocatalytic H₂O₂ production over g-C₃N₄ Nanostructures Utilizing Modified Cyanuric Acid-Melamine Complex as Precursor

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The anthraquinone method and a direct production with H₂ and O₂ as the conventional methods for H₂O₂ production are suffering from several unavoidable challenges such as abundant energy and organic solvents consumption and potential explosion hazard of the mixture of oxygen and hydrogen. Photocatalytic H₂O₂ production has been recently introduced as an energy-saving, environment-friendly, facile, and safe method in this regard. Among different materials used as photocatalyst, graphitic carbon nitride (g-C₃N₄) as an organic metal-free photocatalyst has been widely investigated over the past few years. Despite the advantages, pristine g-C₃N₄ possesses poor photocatalytic activity due to the low specific surface area and easy recombination of photo-induced electrons and holes. Although, various strategies have been developed to overcome these weaknesses, scientific attempts are still made to improve the photocatalytic activity of g-C₃N₄-based photocatalysts. Starting materials used as synthesis precursors have exhibited a significant impact on the physical, optical, and electronic properties of the obtained g-C₃N₄. Commonly, some triazine derivatives such as melamine are utilized as starting substances for g-C₃N₄ preparation. However, utilization of supramolecular structures, especially cyanuric acid-melamine (CM) complex as starting materials has attracted researchers' attention in recent years. In this study, we have investigated the utilization of cyanuric acid-melamine complex (CM) modified by KH₂PO₄ as a starting material in order to improve the robustness of the starting material and provide a dopant source for synthesis of phosphorous and potassium co-doped g-C₃N₄. The as-prepared catalyst exhibited an enhancement in visible light photocatalytic H₂O₂ production than pristine melamine and CM-complex based g-C₃N₄ samples. The experimental results indicated an improved H₂O₂ production of 208 μM under 1 h visible light irradiation as compared with that of the bulk (71 μM) and the pristine nanostructured sample (137 μM).

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