

Enhancement of Broadband Solar Light Absorption and Photocurrent Increase of C₃N₄ Nanoparticles Combined with TiN and Carbon Nanoparticles

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The use of solar energy to produce hydrogen fuel from overall water splitting is a promising means of renewable energy storage. In the past years, various inorganic and organic materials have been developed as photocatalysts for water splitting driven by visible solar light.¹ Plasmonic metal nanostructures has been proposed to offer a route to improve the solar energy conversion efficiency of inorganic/semiconductors materials system.^{1,2} C₃N₄ is an Earth-abundant and low-cost semiconducting photocatalyst material capable of generating H₂ and H₂O₂ from water.³ The band gap energy of 2.7 eV and high valence band and conduction band positions [1.8 and -0.9 eV versus reversible hydrogen electrode (RHE)] makes it promising material for visible light photocatalysis. During water splitting, C₃N₄ require sacrificial reagent and also suffers from poisoning by the produced H₂O₂, which is difficult to remove from the C₃N₄ surface.⁴ Various attempt have been made to improve the catalytic activity of C₃N₄.^{3,4}

Here, we show the strategy to increase the solar light absorption by making a composite of C₃N₄ nanoparticles and plasmonic TiN nanoparticles to improve the photo-electrochemical water splitting performance under simulated solar radiation. Utilization of the broadband plasmonic resonance of the TiN particles and the incorporation of carbon dots (C-Dots) into the C₃N₄ matrix (Fig. 1a) leads to an increase in the UV-vis to NIR absorption over the entire solar spectrum range. The simple chemical synthesis route is used to grow TiN nanoparticles on C₃N₄-carbon dots composite. The hot electron injection from plasmonic nanostructure to composite and C₃N₄ plays role in photocatalysis (Fig. 1b), whereas C-dots acts as chemical catalyst for the decomposition of H₂O₂ into O₂. C-dots plays major role in avoiding the sacrificial reagent and catalytic poisoning. This two-step approach overcomes the low optical absorption, spectral utilization and charge recombination losses, and gives effective way to improve the photocatalytic activity. By incorporating TiN the catalytic performance of C₃N₄-C-dots is increased by 6-fold.

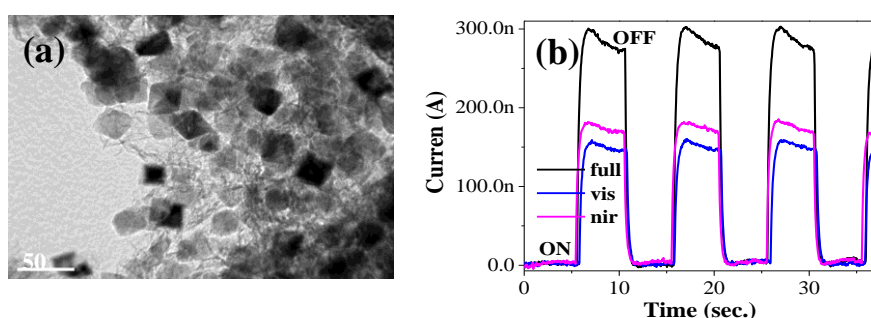


Figure 1. TEM image of TiN decorated C₃N₄ sheets and photocurrent response under simulated visible light.

References:

1. Liu, J. et. al. Science **2015**, 347, 970.
2. Chen, J. et. al. Chem. Commun. **2010**, 46, 7492.
3. J. Liu, Y. Zhang, L. Lu, G. Wu, W. Chen, Chem. Commun. **2012**, 48, 8826.
4. X. Wang et al., Nat. Mater. **2009**, 8, 76.