

Photocatalytic Reduction of CO₂ into Methanol over CuFe₂O₄/TiO₂ under Visible Light Irradiation

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ABSTRACT

The present study aims to focus the photocatalytic reduction of carbon dioxide (CO₂) into methanol on TiO₂ loaded copper ferrite (CuFe₂O₄) photocatalyst under visible light (500 W xenon lamp) irradiation. In this perspective, CuFe₂O₄ and CuFe₂O₄/TiO₂ photocatalysts were synthesized following the sol-gel method from copper(II) nitrate, Cu(NO₃)₂·3H₂O (99 %) and iron(III) nitrate, Fe(NO₃)₃·9H₂O (99 %) as precursors. The phases and crystallite size of the photocatalysts were characterized by X-ray diffraction (XRD), morphology by scanning electron microscopy (SEM), absorption spectrum by ultraviolet-visible spectroscopy (UV-Vis), electron-hole (e⁻/h⁺) recombination process by photoluminescence spectrophotometer, and elemental compositions by energy dispersive X-ray spectroscopy (EDX) instruments. The loading of TiO₂ on CuFe₂O₄ enhanced the photocatalytic activity in the visible light range. The enhanced photoactivity in CuFe₂O₄/TiO₂ semiconductor catalyst can be attributed to interfacial transfer of photogenerated charges, which led to effective charge separation and inhibited the recombination of photogenerated electron-hole (e⁻/h⁺) pairs. Methanol was observed as the main product over CuFe₂O₄/TiO₂ and the photocatalytic activity of CuFe₂O₄/TiO₂ for CO₂ reduction was found to be about three times higher (651 μmol/g_{cat} L) than that of CuFe₂O₄ photocatalyst which might be due to the modification of band gap through TiO₂ loading.

KEYWORDS: CO₂ reduction; Visible light irradiation; CuFe₂O₄/TiO₂ photocatalyst; Band gap energy; Methanol

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