

Enhanced Photocatalytic Hydrogen Production Under Visible Light over Ag Doped TiO₂

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ABSTRACT

TiO₂ is the most widely used photocatalyst for water and air purification, and for hydrogen production, due to its good properties such as chemical and photo-corrosion resistance and low cost. One disadvantage of this material, resides in its bandgap energy (3.2eV), which lies in the UV spectrum. For this reason, studies have been conducted to modify TiO₂ bandgap into the visible light range. Doping elements used for this purpose are noble metals such as Au, Pt and Ag. However, Au and Pt are expensive and scarce materials, leaving Ag as a preferred candidate. TiO₂ and doped TiO₂ were synthesized via Sol-Gel/hydrothermal (SGH) named TiO₂-F and TiO₂Ag-F, respectively, while under the Sol-Gel/hydrothermal/thermal (SGHT) technique was named as TiO₂Ag-C, using titanium butoxide as a precursor and ethanol as solvent. XRD characterization resulted in the presence of the anatase phase in all three synthesized samples as well as the characteristic signals for Ag in TiO₂Ag-F. Samples crystal sizes were determined by the Scherrer equation, and were ~ 10 nm. Light absorption exhibited a shift in the Eg value from 3 eV for TiO₂-F to 1.98 eV for TiO₂Ag-F. TGA indicates the presence of residual organic material for samples synthesized through the SGH technique. BET surface area for the SGH and SGHT photocatalysts were of 140 and 90 m²/g, respectively. SEM images presented particle agglomerates of irregular morphology. Photocatalytic evaluation for hydrogen production was performed using a 250 W mercury light lamp, filtering the UV spectrum. TiO₂Ag-F was the only sample that showed activity, producing 175 μmol of H₂/g catalyst over an 8-h irradiation period. This activity can be mainly attributed to the ability of this material to be activated under the visible light spectrum.

Keywords: TiO₂ doped Ag;, Water splitting; Hydrothermal method.

