

PHOTOCATALYTIC DEGRADATION OF TNT FROM WATER IN UV-VIS/Fe-TiO₂ SYSTEM

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Abstract

2,4,6-Trinitrotoluene (TNT) is one of most common toxic pollutant identified in wastewater generated from ammunitions plants. Due to its potential carcinogenic characteristics, TNT presence in water bodies represents a risk for human health and aquatic life. Among modern treatment methods, TiO₂ photocatalysis was successfully applied in order to remove toxic pollutants. Fe-TiO₂ assisted photocatalytic degradation of TNT in aqueous media, under UV-VIS irradiation was studied. The effects of operating parameters on photocatalytic process performances, kinetic and mechanism of pollutant degradation were investigated. Solutions with $(0.27-2.72) \times 10^{-4}$ M TNT content were photo-oxidized using a medium pressure Hg lamp as UV-VIS light source ($\lambda = 320 - 550$ nm), in the following working conditions: pH = 7; photocatalyst dose = 50 – 500 mg/L; irradiation time = 30 - 240 min. Prior to irradiation, the photocatalyst was added to samples, and resulted suspension was bubbled with air (50 L/h). In order to evaluate the effect of the main active species involved in Fe-TiO₂ assisted photocatalytic degradation of TNT we suppressed the free $\cdot\text{OH}$ radicals mediated process by addition of 16×10^{-3} M iso-propanol (i-PrOH) scavenger. Lock of $\cdot\text{OH}_{\text{ads}}$ radicals' production on the catalyst surface was assured by addition of 16×10^{-3} M sodium iodide (NaI). The initial and irradiated samples were analysed for TNT, NO₃⁻, NO₂⁻ and NH₄⁺ concentrations by Gas Chromatography (GC), and Ion Chromatography respectively. In the tested experimental conditions, at 2.72×10^{-4} M pollutant concentration, the increase of catalyst load up to 200 mg/L leads to the enhancement of initial TNT degradation rate up to 0.64×10^{-7} Ms⁻¹. Since, ten times increase of initial TNT content has a negative effect on pollutant degradation rate constant, in similar experimental condition, prolonged irradiation time from 60 to 240 min was needed in order to assure pollutant advanced degradation efficiencies ($\geq 99.9\%$). The TNT degradation and its inorganic by-products formation obeyed a pseudo-first-order kinetics. The experimental results of the reactive species quenching showed that $\cdot\text{OH}$ radicals was the predominant oxidant species participated in reaction, and the pollutant degradation occurred mainly on the surface of catalyst.

Keywords: 2,4,6-Trinitrotoluene, Fe-TiO₂ photocatalyst, AOPs