

PHOTOCATALYTIC ACTIVITY OF METAL OXIDE NANOPARTICLES FOR REMOVAL OF THE HERBICIDE FLUROXYPYR FROM WATER**Vesna Despotović¹, Paula Sfirloaga², Daniela Šojić Merkulov¹, Biljana Abramović¹**¹*University of Novi Sad Faculty of Sciences, Department of Chemistry, Biochemistry and Environmental Protection, Trg Dositeja Obradovića 3, 21000 Novi Sad, Serbia*²*National Institute for Research and Development in Electrochemistry and Condensed Matter, Department of Condensed Matter, 1 Plautius Andronescu St., 300224 Timisoara, Romania**e-mail: vesna.despotovic@dh.uns.ac.rs***Abstract**

The photocatalytic materials absorb the photons having energy equals to or more than the band gap energy between the valence and conduction bands of the photocatalyst. The positive holes in the valence band either oxidize the water to produce hydroxyl radical or pollutant, whereas excited electrons reduce the adsorbed oxygen on the photocatalyst in the conduction band [1]. Till date a wide range of metal oxide/semiconductor-based nanomaterials have been explored for the photocatalytic degradation of harmful and toxic organic pollutants into the non-toxic products [2,3]. The aim of this work was to investigate removal of the herbicide fluroxypyr from double distilled water in the presence of novel TiO₂, MgO and ZnO nanoparticles under UV and simulated solar irradiation. It was found that TiO₂ photocatalyst showed higher photocatalytic activity for removal of fluroxypyr from water compared with MgO and ZnO. However, a major drawback of photocatalyst is that it has higher rate of recombination of photogenerated electron-hole pairs, which suppressed its catalytic potential [1,4]. Because of that it was investigated the photodegradation behaviour of fluroxypyr in aquatic systems using UV and simulated solar irradiation in the presence of TiO₂/(NH₄)₂S₂O₈. The efficiency of elimination the herbicide from double distilled water was monitored by UFLC–DAD technique.

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References

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