

# Enhanced photoactivity of cerium tungstate-modified graphitic carbon nitride heterojunction photocatalyst for the photodegradation of moxifloxacin

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## Abstract

Design and optimization of visible-light-driven photocatalysts for degradation of organic pollutants is an important step towards environmental decontamination. In this study, wolframite cerium tungstate ( $\text{Ce}_2(\text{WO}_4)_3$ , (CW)) hybridized with g- $\text{C}_3\text{N}_4$  (CN) nanosheets was synthesized via a simple hydrothermal route followed by an ultrasound-assisted synthesis method. The prepared  $\text{Ce}_2(\text{WO}_4)_3@ \text{g-C}_3\text{N}_4$  (CW@CN) heterojunction was investigated for photocatalytic degradation of the antibiotic moxifloxacin (MXF) under visible light irradiation. Structural, morphological, and optical properties as well as chemical composition of the as-synthesized heterojunction were investigated by transmission electron microscopy (TEM), X-ray photoelectron spectroscopy (XPS), X-ray diffraction (XRD), UV-Vis diffuse reflectance spectroscopy (UV-Vis DRS) and photoluminescence (PL). MXF photocatalytic degradation by the binary nanostructure ( $\text{Ce}_2(\text{WO}_4)_3@ \text{g-C}_3\text{N}_4$ ) (94.1%) was the highest compared to g- $\text{C}_3\text{N}_4$  (53.6%) and  $\text{Ce}_2(\text{WO}_4)_3$  (46.4%). Such enhanced activity could be ascribed to efficient suppression of the charge carriers' recombination, leading to adequate formation of the reactive species responsible for MXF degradation. Furthermore, the  $\text{Ce}_2(\text{WO}_4)_3@ \text{g-C}_3\text{N}_4$  heterojunction showed remarkable stability over five consecutive cycles, with only 11.5% reduction after the 5th cycle. This work established the potential applicability of  $\text{Ce}_2(\text{WO}_4)_3@ \text{g-C}_3\text{N}_4$  nanostructures towards photocatalytic removal of MXF.