

Adsorption and photocatalytic characteristics for degradation of Persistent Organic Pollutants on Carbon-coated TiO₂

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Carbon-coated TiO₂ samples were prepared from the mixture of poly (vinyl alcohol) and commercial TiO₂ (P-25) with different mass ratios at a heating temperatures of 500–900°C for 1h in high purity Argon gas flow. The samples were characterized by XRD, SEM, TEM, TGI and BET analysis. Adsorption properties and photocatalytic activities of commercial and carbon-coated TiO₂ catalysts were compared for the oxidation of methylene blue (MB). The photocatalytic degradation of MB in a batch photocatalytic reactor, using a UV lamp as a light source and TiO₂ as a photocatalyst, was investigated. The effects of various parameters such as the initial MB concentration, and the initial pH on the MB degradation rate of TiO₂ photocatalysis were examined. In the presence of both UV light illumination and TiO₂ catalyst, MB was more effectively degraded than with either UV or TiO₂ alone. The reaction rate was found to obey pseudo first-order kinetics represented by the Langmuir-Hinshelwood model. It was also observed that carbon-coated samples gave faster kinetics for the oxidation of organic molecules.