

Photocatalytic activity of TiO₂ treated by EtOH, H₂O and plasma

Myoung Joo Kim, Yuan Luo, Kwang-Dae Kim and Young Dok Kim*

Department of Chemistry, Sungkyunkwan University, Suwon, 440-746, Korea

Due to the high catalytic activity and good thermal stability, TiO₂ is widely used in various photocatalyst applications. In the present work, the photocatalytic activities of pristine and variously treated Degussa P-25 catalysts were compared. The surface structure and chemical properties were characterized by Brunauer, Emmett & Teller (BET), X-ray diffraction (XRD) UV-vis spectroscopy, and Fourier transform infrared spectroscopy (FT-IR). The photocatalytic activity was evaluated by the photocatalytic degradation of methylene blue (MB) under UV irradiation (254 nm) at room temperature. The samples treated by N₂ and O₂ plasma were more effective than commercial TiO₂ in photocatalytic degradation of MB. The photocatalytic activity became even higher when the samples were immersed into water or EtOH and dried prior to the photocatalysis-experiment. The FT-IR spectra indicated the adsorption of various water species chemisorbed in TiO₂. The peak at 3400 cm⁻¹ and 3200 cm⁻¹ were attributed to water molecules on TiO₂ with random (liquid-like) and tetrahedral (ice-like) arrangements, respectively. When the relative amount of liquid-like water increased, photocatalytic activity was enhanced.