

SYNTHESIS AND PHOTO-CATALYTIC ACTIVITIES OF N-DOPED TITANIA NANO-CHAINS

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Photocatalysis has emerged to be one of the most promising pollution remediation technologies in recent decades. Of all the photocatalysts, TiO2 is the most widely studied one. It is very stable, cheap to produce and is capable of degrading a wide range of organic pollutants. However, the semiconductor TiO2 has its shortcomings. Due to its wide bandgap, it can only be activated by near-UV radiation. In order to overcome these problem, some metal ions have been doped into the lattice of the TiO2 catalysts. This modifies the electronic structure of TiO₂ by narrowing the bandgap, rendering the doped TiO₂ more sensitive to the visible light. In this study, the nitrogen-doped TiO₂ anatase powders were synthesized by hydrothermal digestion of the hydrolysis product of ammonium titanyl oxalate with ammonia. The TEM images of N-doped TiO2 show that chain-like structure was composed of 20nm sized single crystalline. The BET surface area of this powder was 87.75m²/g. We also evaluated the XRD patterns for this powder and the crystal structure was pure anatase. In the UV-Vis absorption spectrum, nitrogen doping causes the absorption edge of TiO2 to shift to a lower energy region. It is considered that nitrogen atoms in doped TiO2 anatase powder are responsible for the significant enhancement in the doped TiO₂ photoactivity under visible light irradiation. We found the photocatalytic activity of the N-doped TiO2 nano-chain was higher than that of the commercial TiO2 anatase for methyl orange decomposition under visible light and UV light irradiation.