

### Photocatalytic activity of $Ti_{1-x}M_xO_2$ (M = Co and Cr) crystals

김현규<sup>1,\*</sup>, 정의덕<sup>1</sup>, 김해진<sup>2</sup>, 홍경수<sup>1</sup>, 박혁규<sup>1,3</sup>, 홍석준<sup>4</sup>,  
배상원<sup>4</sup>, 지상민<sup>4</sup>, 장점석<sup>4</sup>, 이재성<sup>4</sup>

<sup>1</sup>한국기초과학지원연구원 부산센터;

<sup>2</sup>한국기초과학지원연구원 미래융합연구실;

<sup>3</sup>부산대학교 물리학과; <sup>4</sup>포항공과대학교 화학공학과

(hhgkim@kbsi.re.kr\*)

Metal-doped  $TiO_2$  nanoparticles were prepared by the sol-gel and hydrothermal synthesis methods. The estimated quantum yields (QYs) of Pt/Cr-doped  $TiO_2$  nanoparticles was ca. 0.3 %. Cr-, Co- and N-doped  $TiO_2$  showed the photocatalytic activity for IPA degradation to  $CO_2$ , but only Cr-doped  $TiO_2$  produced  $H_2$  photocatalytically in the presence of methanol-water aqueous solution under visible light ( $\lambda > 420nm$ ). Thus, in the case of Pt/Co-doped  $TiO_2$ , the electron excited to the conduction band has a sufficient reduction potential to reduce  $H^+$  ion, but hole in the valence band has lower oxidation potential than required for  $CH_3OH$  degradation to  $CO_2$ . Therefore, Cr-, Co- and N-doped  $TiO_2$  show the different activity for the photocatalytic reaction of gases and solution phases. The results of the calculated electronic structure and experimental optical properties are correlated to schematically to describe the possible mechanism of the photocatalytic behavior of the system under study.