

Supporting Information

Ti(IV) nanocluster as a promoter on semiconductor photocatalysts for oxidation of organic compounds

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Table S1. Performances of the photocatalysts under 1mW/cm² blue light irradiation

Sample	Cu(II)-Ti(IV)-WO ₃	Cu(II)-WO ₃
Rp ^a (quanta/sec)	3.91×10 ¹⁵	3.91×10 ¹⁵
Rco ₂ (μmol/h)	0.83	0.47
QE(%)	21.2	12.0

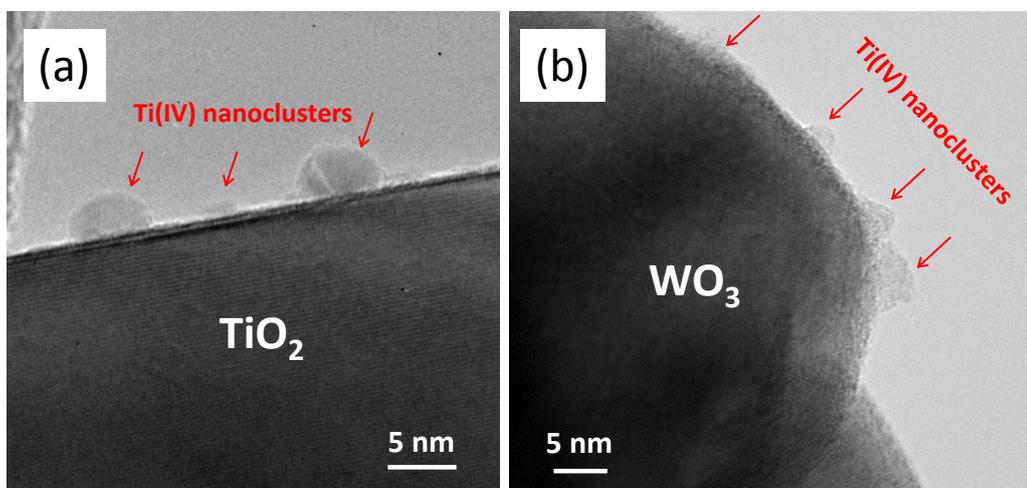


Fig. S1 High Resolution TEM images for Ti(IV)-TiO₂ (a) and Ti(IV)-WO₃ (b).

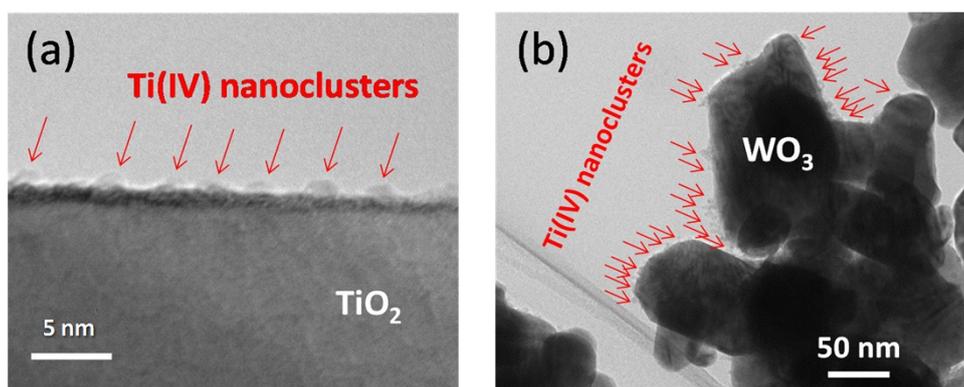


Fig. S2 TEM images for Ti(IV)-TiO₂ (a) and Ti(IV)-WO₃ (b).

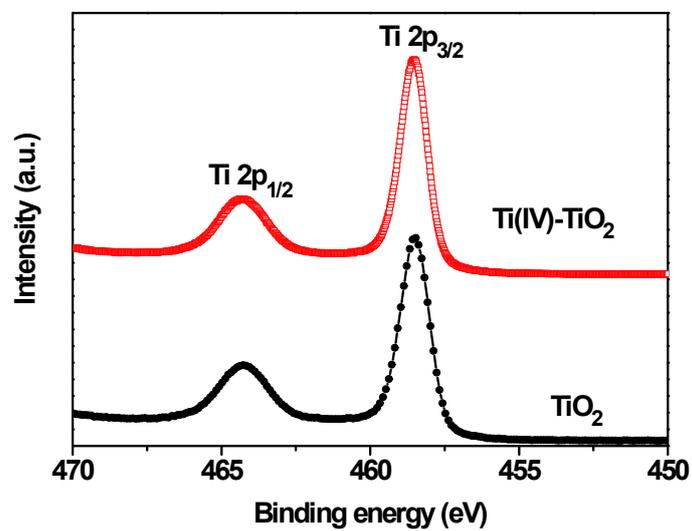


Fig. S3 XPS on Ti 2p core-level of TiO_2 and Ti(IV)-TiO_2 , respectively.

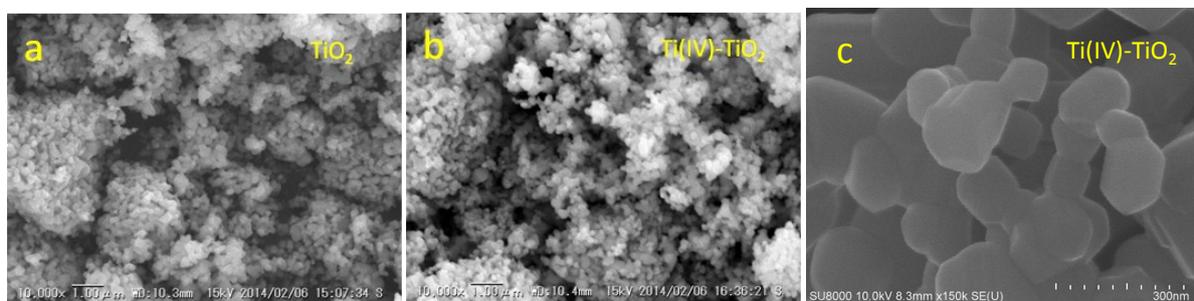


Fig. S4 SEM images of rutile TiO_2 (a) and Ti(IV)-TiO_2 (b). High resolution FE-SEM image of Ti(IV)-TiO_2 (c).

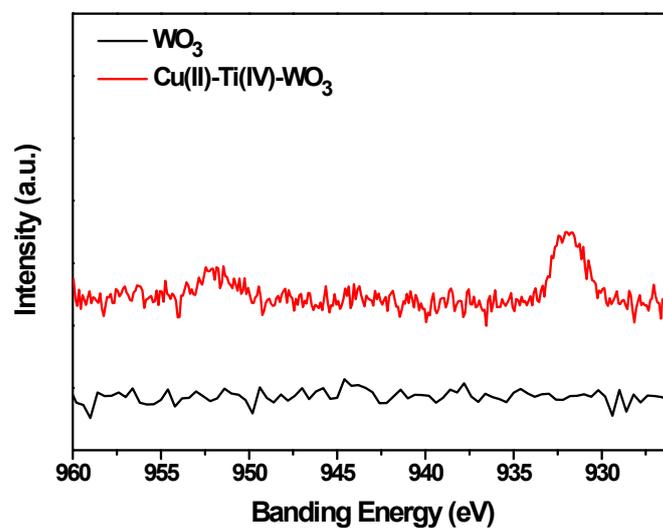


Fig. S5 XPS on Cu 2p core-level of WO_3 and $\text{Cu(II)-Ti(IV)-WO}_3$.

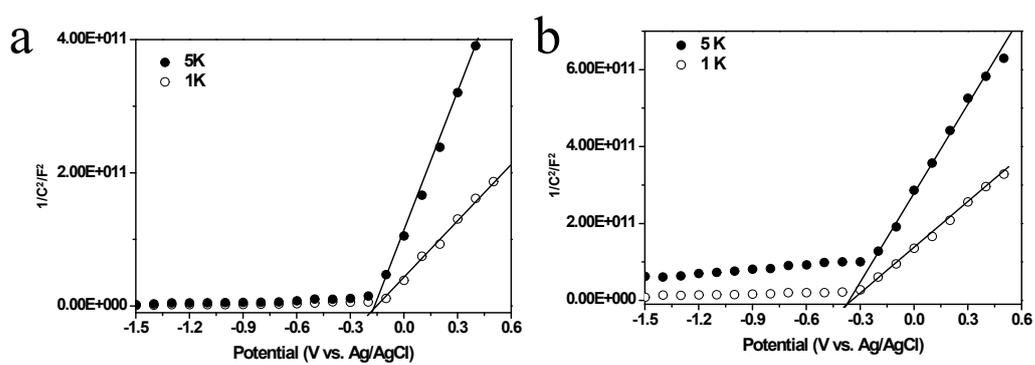


Fig. S6 Mott-Schottky plots of WO_3 (a) and Ti(IV)-WO_3 (b) at frequencies of 1 k Hz and 5 k Hz, respectively.

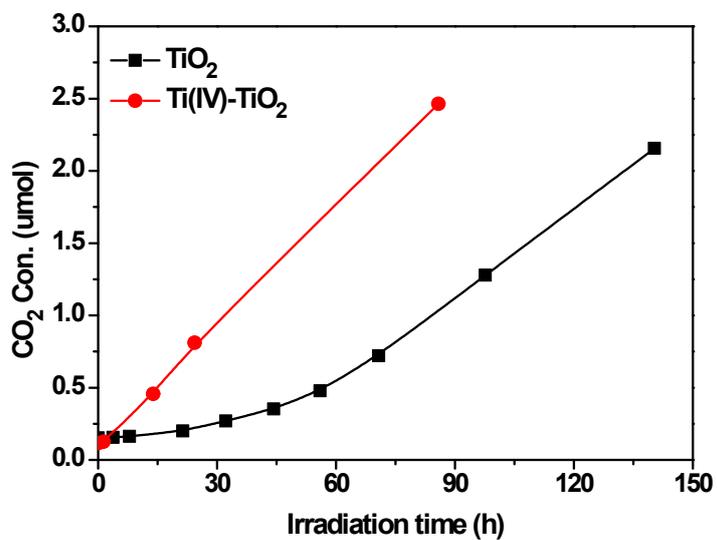


Fig. S7 Photocatalytic oxidation of 2-propanol on TiO₂ and Ti(IV)-TiO₂ under UV light irradiation.

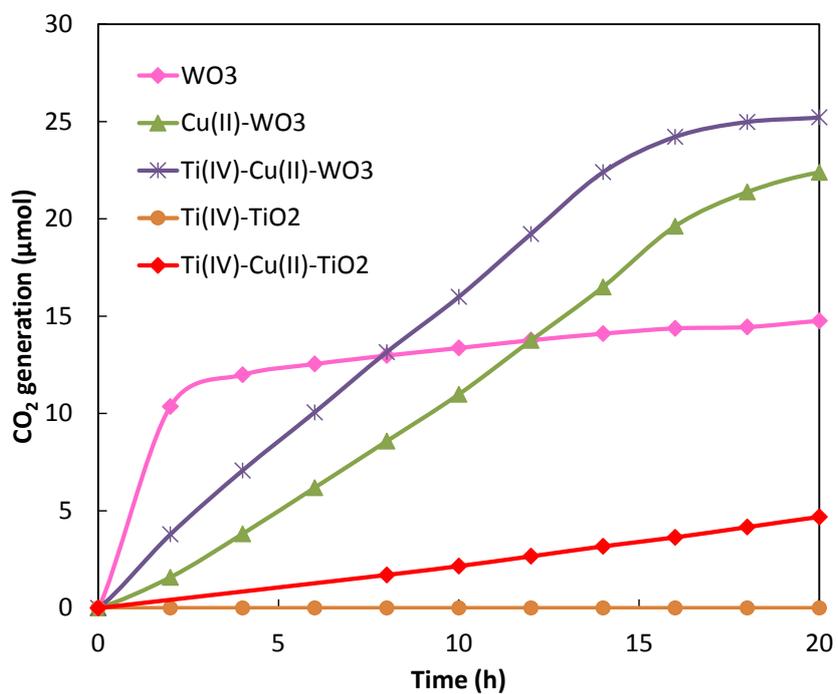


Fig. S8 Photocatalytic oxidation of acetaldehyde under a commercial white LED irradiation.

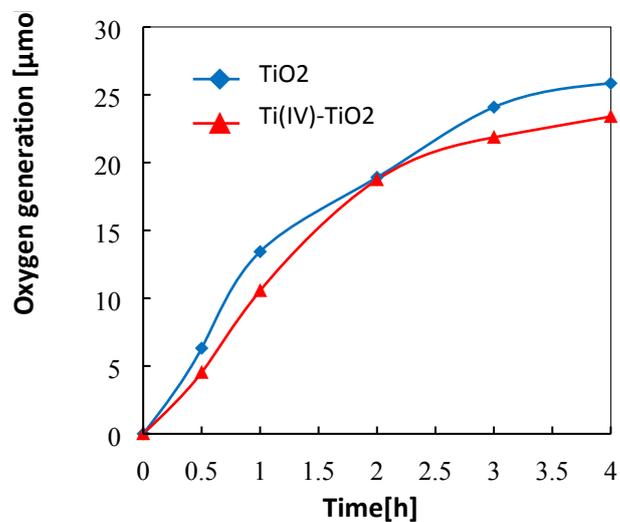


Fig. S9 Photocatalytic activities of TiO₂ and Ti(IV)-TiO₂ for water oxidation in aqueous solution containing sacrificial AgNO₃ agent (50 mmol/L). Light source was Hg-Xe lamp.

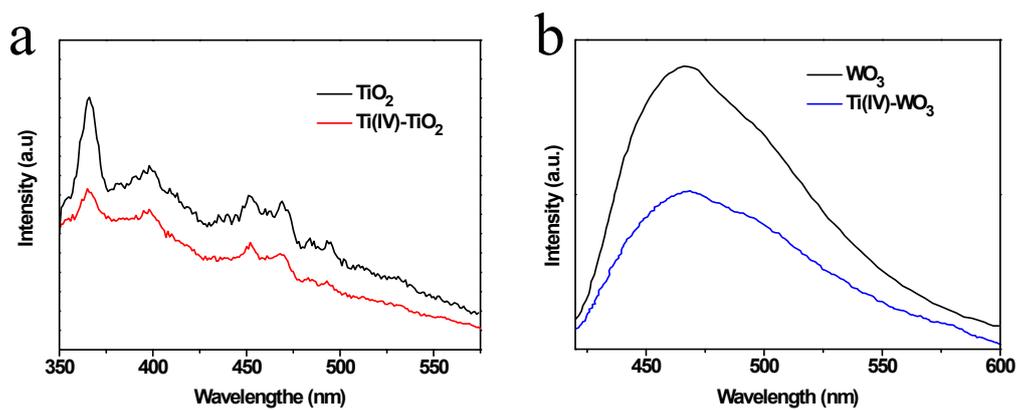


Fig. S10 Photoluminescence spectra for TiO₂ and WO₃ samples

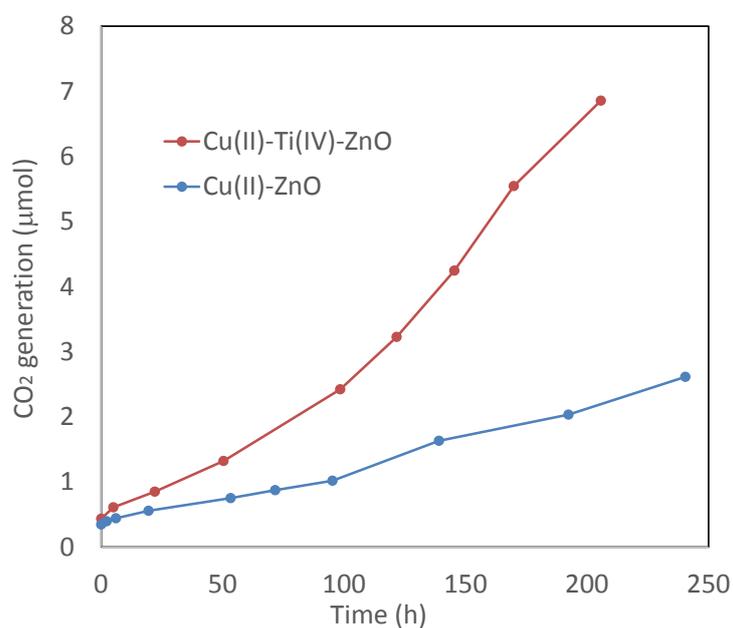


Fig. S11 Photocatalytic activities for oxidation of 2-propanol for Cu(II)-Ti(IV)-ZnO and Cu(II)-ZnO under blue light irradiation.

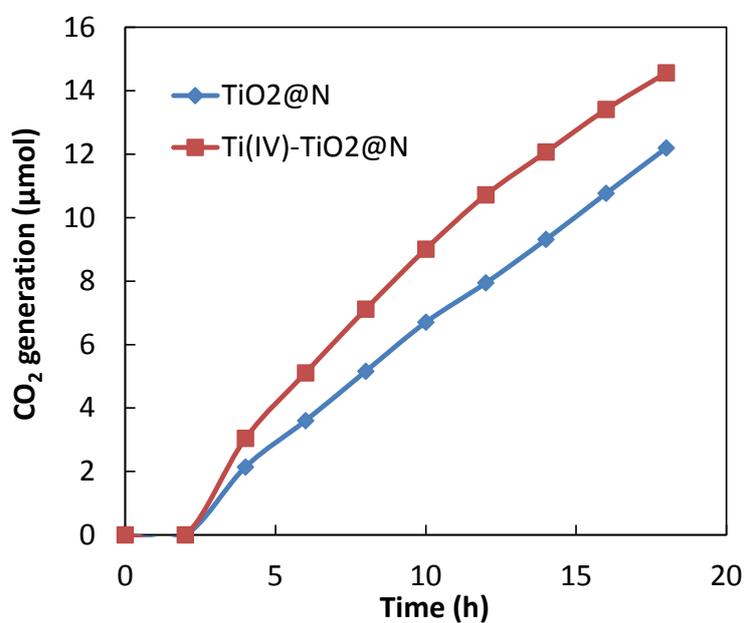


Fig. S12 Photocatalytic activities for oxidation of acetaldehyde for nitrogen doped TiO₂ (TiO₂@N) and Ti(IV) nanoclusters grafted nitrogen doped TiO₂ (Ti(IV)-TiO₂@N) under white LED irradiation.