



Title	Surface modification of TiO ₂ with metal oxide nanoclusters: a route to composite photocatalytic materials
Author(s)	Nolan, Michael
Publication date	2011-06-30
Original citation	Nolan, M. (2011) 'Surface modification of TiO ₂ with metal oxide nanoclusters: a route to composite photocatalytic materials', <i>Chemical Communications</i> , 47(30), pp. 8617-8619. doi: 10.1039/C1CC13243A
Type of publication	Article (peer-reviewed)
Link to publisher's version	http://dx.doi.org/10.1039/C1CC13243A Access to the full text of the published version may require a subscription.
Rights	© The Royal Society of Chemistry 2011. This is the accepted manuscript version of an article published in <i>Chemical Communications</i> . The version of record is available at http://dx.doi.org/10.1039/C1CC13243A
Item downloaded from	http://hdl.handle.net/10468/5187

Downloaded on 2018-08-23T19:23:04Z

Cite this: DOI: 10.1039/c0xx00000x

www.rsc.org/xxxxxx

ARTICLE TYPE

Surface Modification of TiO₂ with Metal Oxide Nanoclusters: a Route to Composite Photocatalytic Materials

Michael Nolan^{*a}*Received (in XXX, XXX) Xth XXXXXXXXX 20XX, Accepted Xth XXXXXXXXX 20XX*

DOI: 10.1039/b000000x

Density functional theory simulations show that modifying rutile TiO₂ with metal oxide nanoclusters produces composite materials with potential visible light photocatalytic activity.

Developing visible light active photocatalysts for hydrogen production is a major activity. TiO₂ has desirable qualities of being cheap, readily available, stable and non-toxic, but pure TiO₂ is not suitable, since it primarily absorbs in the ultra violet. Modifying TiO₂ to allow visible light absorption is well studied, reviewed in refs. 1 and 2.

Many efforts in enabling visible light absorption have focused on substitutional cation or anion doping at the Ti or O sites³⁻¹¹, or cation-anion co-doping¹²⁻¹⁴. However, there are issues with substitutional doping of TiO₂, such as dopant incorporation/solubility, stability and charge recombination. Strategies for surface modification of TiO₂ shifting the band gap to the visible while improving photocatalytic efficiency are being researched.

The dye sensitised solar cell (DSSC), developed by Gratzel and co-workers^{15,16} is a paradigm in this regard, in which a dye molecule is adsorbed at the surface of TiO₂ nanoparticles and plays the role of a visible light absorber. Metal sulphide quantum dots adsorbed on TiO₂ have been studied¹⁷. Recent developments have shown that oxide-oxide heterostructures¹⁸⁻²³ can display visible light absorption and improved photocatalytic activity over the pure oxides. Examples include Bi₃Ti₄O₁₂-TiO₂¹⁸, BiVO₄-WO₃¹⁹ and BiOBr-ZrFe₂O₄²⁰. The valence and conduction band alignments of the oxides in the heterostructure are postulated to allow visible light absorption and charge separation¹⁸⁻²³.

One can conceive of using metal oxide nanoclusters adsorbed at a TiO₂ surface as a composite photocatalytic material, with the oxide nanocluster playing a role similar to the dye in a DSSC or the sulphide quantum dot. A visible light active heterostructure of iron oxide supported on TiO₂ has been reported²¹⁻²³, in which *dispersed, molecular sized FeO_x species* are supported on TiO₂. Atomic layer deposition²¹ and a chemisorption-calcination cycle^{22,23} have been used to fabricate the heterostructure, which demonstrates band gap narrowing and visible light absorption, as well as reduced charge recombination, as characterised by photoluminescence spectroscopy²¹⁻²³. Band gap narrowing is suggested to arise from the presence of iron oxide states in the band gap, shifting the valence band edge upwards. The heterostructure approach has advantages over substitutional doping; alleviating many issues and allows fine tuning of properties through composition.

We have used first principles density functional theory (DFT), with Hubbard correction for on-site Coulomb interactions (DFT+U) to study TiO₂ nanoclusters supported on TiO₂ and we find that the heterostructures have reduced band gaps, and will facilitate electron-hole separation²⁴. To investigate other heterostructures composed of metal oxide nanoclusters supported on TiO₂, this paper presents DFT+U studies of metal oxide nanoclusters adsorbed on the rutile (110) surface. The rutile TiO₂ (110) surface is chosen as rutile is the most stable TiO₂ phase, and the (110) surface, as well as being well studied²⁵⁻²⁸, is energetically the most stable surface among the low index surfaces. Rutile (110) is a suitable substrate for deposition of other species^{29,30}.

In the calculations, for which the computational details are presented in the ESI (sec. S1), we adsorb and relax the metal oxide nanoclusters on the rutile (110) surface. The oxide nanoclusters are Sn₂O₄, Cr₂O₄/Cr₂O₃, Mo₂O₄ and Zr₂O₄. This size of nanocluster is studied since refs. 22 and 23 indicate that the FeO_x species supported on TiO₂ are of this size. The supported oxide nanoclusters are in a size regime realisable experimentally. The minimum energy structures for free M₂O₄ and M₂O₃ clusters are shown in figure 1(f) (see ESI) and are consistent with the global minima for these stoichiometries^{31,32}. Two oxidation states are considered for CrO_x since the energy required for the transformation of Cr₂O₄ ↔ Cr₂O₃ is small, see ESI, sec. S2.

The relaxed adsorption structures for the clusters supported on TiO₂ are shown in figure 1. The adsorption energies, computed relative to a bare (110) rutile surface and a free oxide nanocluster lie in the range of -3.43 eV (Cr₂O₄) to -7 eV (Zr₂O₄), indicating strong binding between the oxide cluster and the TiO₂ surface.

For all supported oxides, the cations of the cluster bind to bridging and in-plane oxygen atoms of TiO₂. Sn₂O₄, Zr₂O₄ and Cr₂O₃ adsorb at the TiO₂ surface using three cluster oxygen atoms to bond to surface 5-fold coordinated Ti. In Cr₂O₄ and Mo₂O₄, two cluster oxygen atoms bind with 5fold coordinated surface Ti. In the Sn₂O₄ and Zr₂O₄ clusters, the fourth oxygen bridges the cluster cations, while in Cr₂O₄ and Mo₂O₄, two oxygen atoms of the cluster do not interact with surface Ti.

In examining the geometry of the cluster at the surface, we see multiple new metal-O bonds between the cluster and the surface, which make the cations in the cluster 6 coordinate, which is a favourable coordination environment. Compared to the gas phase clusters, adsorption tends to break the symmetry of the cluster. Metal-oxygen distances in the adsorbed cluster are lengthened

compare to the free oxide clusters. The distances for the cluster cation to surface oxygen are longer than in the corresponding oxide of the cation, but are nonetheless consistent with formation of a bond between the cation and surface oxygen. Finally, Ti-O distances to oxygen of the supported cluster are generally typical of TiO₂. We note that supported Cr₂O₄ has the least cluster-

surface bonds (see complete geometry data in sec. S3, ESI) and the smallest adsorption energy, while Zr₂O₄ has the most cluster-surface bonds and this is consistent with the corresponding adsorption energies, so that maximising the number of new cluster-surface bonds enhances adsorption of the oxide cluster²⁴.

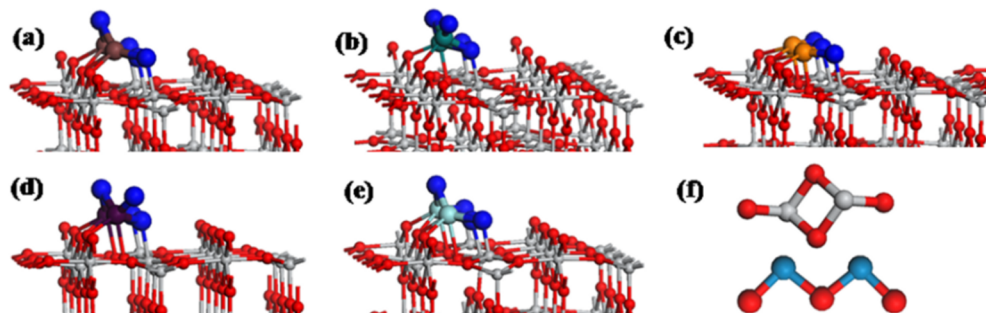


Fig. 1 Relaxed atomic structure of metal oxide clusters supported on the rutile TiO₂ (110) surface. (a) Sn₂O₄, (b) Cr₂O₄, (c) Cr₂O₃ (d) Mo₂O₄ and (e) Zr₂O₄. Part (f) shows the most stable structure for an M₂O₄ and an M₂O₃ nanocluster. In (a)-(e), Ti of the surface is grey, O from the surface is red, O from the cluster is large blue spheres and the cations in the clusters are coloured differently for each element. In (f) the cluster oxygen are red.

The alignments of the electronic states of the two oxides determine the photocatalytic properties. In experiments, the absorption spectra display a shift to longer wavelength, indicative of band gap narrowing. Photoluminescence spectra can be used to study charge recombination. XPS has been used to determine cation oxidation states. From these analyses the relative positions of the valence and conduction bands have been discussed.

In the present calculations, we examine the electronic density of states (EDOS), projected onto cation and oxygen states from the support and the cluster (PEDOS) to determine the impact of surface modification on the electronic structure of the composite; the PEDOS of the bare rutile (110) surface is shown in the ESI (sec. S4). Bader charge analysis³³ allows identification of cation oxidation states. Formation of an interface between the support and the cluster allows the alignments of the surface and cluster electronic states to be determined. It is this alignment of the electronic states upon modification of TiO₂ by the cluster that is of interest; ref. 20 highlights the importance of the interface in the heterostructure.

Figure 2 shows the PEDOS for Sn 5s, Zr 5d states and oxygen 2p states from the surface and the cluster for the composites. Analysing the electronic structure shows that the Zr₂O₄-TiO₂ composite shows no change in the band gap from pure TiO₂. The ZrO₂ derived electronic states lie below the VB and above the CB of TiO₂; this arises from the very large band gap of ZrO₂ structures. Analysis of a Hf₂O₄-TiO₂ heterostructure gives the same conclusion – oxides with band gaps much larger than that of TiO₂ will not be suitable for reducing the band gap of TiO₂; see sec. S5 in the ESI.

The top of the valence band of Sn₂O₄-modified TiO₂ is derived from the TiO₂ surface, while the bottom of the conduction band is derived from empty Sn 5s states of the cluster, shifting the conduction band down in energy by 0.2 eV, and narrowing the band gap. Although band gap changes determined with DFT+U (for which the band gap remains underestimated) will be qualitative, the shifts in the valence or conduction band edges should be useful for comparing trends across a series of materials.

The band gap reduction with Sn₂O₄ modified TiO₂ is small and may not be sufficient to shift absorption to the visible.

For supported Cr₂O₄ and Mo₂O₄ nanoclusters (see ESI, sec. S6 for Cr₂O₃), the PEDOS are shown in figure 3. In this case, the top of the valence band is pushed upwards, by between 0.4 eV and 0.6 eV, due to the presence of electronic states from the adsorbed cluster. This shift is large enough to narrow the band gap into the visible, so that if the TiO₂ surface can be modified by adsorption of these metal oxides, visible light absorption will take place.

Bader charges have been computed for the cations in the adsorbed cluster and in the surface (see ESI, table S2). For Ti, the Bader charges take values typical of Ti⁴⁺, while the various cluster cations also take Bader charges consistent with their expected oxidation states. Thus, there is no charge transfer between the supported cluster and the TiO₂ surface.

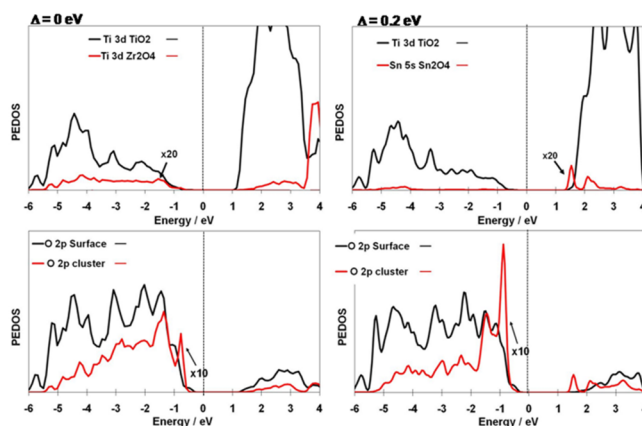


Fig. 2 PEDOS projected onto Ti 3d, Sn 5s, Zn 5d and O 2p states for Sn₂O₄ and Zr₂O₄ supported on the rutile (110) surface. **Left panel:** Sn₂O₄, **Right panel:** Zr₂O₄. The shift in the valence-conduction band energy gap is given at the top of each panel in eV. The Fermi level is indicated by vertical line at 0 eV. The cluster PEDOS are enhanced by the factors given in the figure, since there are 2 cations and 4 anions, compared with 96 surface cations and 192 cluster anions.

The alignments of the surface and cluster electronic states in CrO_x and Mo₂O₄-modified TiO₂ have the consequence that upon excitation of an electron by visible light from the nanocluster states to the CB, the excited electron will migrate to the TiO₂ surface, while the resulting hole will be found on the cluster. Even if the cluster has a short charge diffusion length, the dimensions of the cluster mitigate against this, as discussed for iron oxide modified TiO₂²⁰⁻²³. For Sn₂O₄, excitation of electrons will result in holes migrating to the TiO₂ surface and electrons to the cluster, specifically in the Sn 5s states, which are good conducting states. Thus, the heterostructure approach facilitates separation of electrons and holes after excitation, which is necessary to reduce charge recombination and is the second factor in enabling improved photocatalytic activity. The modifications of TiO₂ with metal oxide clusters thus shares some similarities with other approaches to surface modified TiO₂ for visible light activity, but provides another route to tune properties.

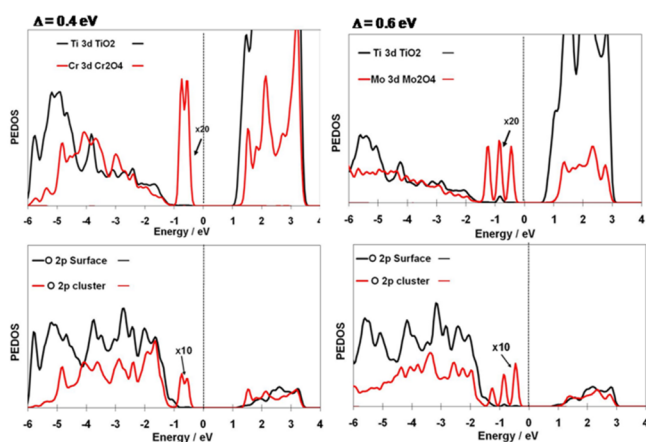


Fig. 3 PEDOS projected onto Ti 3d, Cr 3d, Mo 4d and O 2p states for Sn₂O₄ and Zr₂O₄ supported on the rutile (110) surface. **Left panel:** Cr₂O₄, **Right panel:** Mo₂O₄. The shift in the valence-conduction band energy gap is given at the top of each panel in eV. The Fermi level is indicated by the vertical line at 0 eV. The cluster PEDOS are enhanced by the factors given in the figure, since there are 2 cations and 4 anions, compared with 96 surface cations and 192 cluster anions.

In conclusion, we have shown that TiO₂ modified with metal oxide nanoclusters, in particular CrO_x and Mo₂O₄, is a heterostructured material system with the necessary properties to be a visible light active photocatalytic material. With advances in synthesis of heterostructured composites, this is another approach to developing visible light active photocatalytic materials.

Support from Science Foundation Ireland through a Starting Investigator Research Grant, (“EMOIN”: SFI 09/SIRG/I1620) is acknowledged, as are computing resources funded by SFI and provided by the SFI/Higher Education Authority Funded Irish Centre for High End Computing.

Notes and references

^a Tyndall National Institute, University College Cork, Lee Maltings, Prospect Row, Cork, Ireland. Email: michael.nolan@tyndall.ie

† Electronic Supplementary Information (ESI) available: details of the computational methods employed, geometry data, TiO₂ (110) surface EDOS, data on CrO_x and Hf₂O₄ supported on TiO₂. See DOI: 10.1039//

- 1 A. Fujishima, X. Zhang and D. A. Tryk, *Surf. Sci. Rep.* 2008, **63**, 515.
- 2 X. L. Nie, S. P. Zhou, G. Maeng and K. Sohlberg, *Int. J. Photoenergy* 2009, 294042
- 3 Y. Cui, H. Du and L. S. Wen, *J. Mat. Sci. and Tech.* 2008, **24**, 675
- 4 H. W. Peng, J. B. Li, S. S. Li and J. B. Xia, *J. Phys. Cond. Matt.* 2008, **20**, 125207
- 5 C. Di Valentin, G. Pacchioni, H. Onishi and A. Kudo, *Chem. Phys. Lett.* 2009, **469**, 166
- 6 J. G. Yu, Q. J. Xiang and M. H. Zhou, *Appl. Cat. B Environmental* 2009, **90**, 595
- 7 L. Bian, M. X. Song, T. L. Zhou, X. Y. Zhao and Q. Q. Dai, *J. Rare Earths* 2009, **27**, 461
- 8 C. Di Valentin, E. Finazzi, G. Pacchioni, A. Selloni, S. Livraghi, M. C. Paganini and E. Giamello, *Chem. Phys.* 2007, **339**, 44
- 9 A. M. Czoska, S. Livraghi, M. Chiesa, E. Giamello, S. Agnoli, G. Granozzi, E. Finazzi, C. Di Valentin and G. Pacchioni, *J. Phys. Chem C* 2008, **112**, 8951
- 10 R. Long and N. J. English, *J. Phys. Chem. C* 2010, **114**, 11984
- 11 J. W. Zheng, A. Bhattacharyya, P. Wu, Z. Chen, J. Highfield, Z. L. Dong and R. Xu, *J. Phys. Chem. C* 2010, **114**, 7063
- 12 Y. Q. Gai, J. B. Li, S. S. Li, J. B. Xia and S. H. Wei, *Phys. Rev. Lett.* 2009, **102**, 036402
- 13 W. G. Zhu, X. F. Qiu, V. Iancu, X. Q. Chen, H. Pan, W. Wang, N. M. Dimitrijevic, T. Rajh, H. M. Meyer, M. P. Paranthaman, G. M. Stocks, H. H. Weiering, B. H. Gu, G. Eres and Z. Y. Zhang, *Phys. Rev. Lett.* 2009, **103**, 2264101
- 14 J. Zhang, C. X. Pan, P. F. Fang, J. H. Wie and R. Xiong, *ACS Applied Materials and Interfaces* 2010, **2**, 1173
- 15 M. Gratzel, *MRS Bull.*, 2005, **30**, 23
- 16 D. Wei, *Int. J. Mol. Sci.*, 2010, **11**, 1103
- 17 G. Ai, W. Sun, X. Gao, Y. Zhang, and L-M. Peng, *J. Mat. Chem.* 2011, **21**, 8749
- 18 T. Cao, Y. Li, C. Wang, Z. Zhang, M. Zhang, C. Shao and Y. Liu, *J. Mat. Chem.* 2011, **21**, 6922
- 19 S. J. Hong, S. Lee, J. S. Jang and J. S. Lee, *Energy Envir. Science*, 2011, **4**, 1781
- 20 L. Kong, Z. Jiang, T. Xiao, L. Lu, M. O. Jones and P. P. Edwards, *Chem. Comm.*, 2011, **47**, 5512
- 21 J. A. Libera, J. W. Elam, N. F. Sather, T. Rajh and N. M. Dimitrijevic, *Chem. Mat.*, 2010, **22**, 409
- 22 H. Tada, Q. Jin, H. Nishijima, H. Yamamoto, M. Fujishima, S-i. Okuoka, T. Hattori, Y. Sumida and H. Kobayashi, *Angew Chem.* 2011, **50**, 3501
- 23 Q. Jin, N. Fujishima and H. Tada, *J. Phys. Chem. C* 2011, **115**, 6478
- 24 A. Iwaszuk and M. Nolan, *Phys. Chem. Chem. Phys.* 2011, **13**, 4963
- 25 U. Diebold, *Surf. Sci. Rep.* 2003, **48**, 53
- 26 Z. Dohnalek, I. Lyubinetsky and R. Rousseau, *Prog. In Surf. Sci.*, 2010, **85**, 161
- 27 E. Wahlstrom, E. K. Vestergaard, R. Schaub, A. Ronnau, M. Vestergaard, E. Laegsgaard, I. Stensgaard and F. Besenbacher, *Science* 2004, **303**, 511
- 28 R. Schaub, E. Wahlstrom, A. Ronnau, E. Laegsgaard, I. Stensgaard and F. Besenbacher, *Science* 2003, **299**, 377
- 29 C. L. Pang, R. Lindsay and G. Thornton, *Chem. Soc. Rev.*, 2008, **37**, 2328
- 30 G. D. Sheng, J. X. Li, S. W. Wang and X. K. Wang, *Prog. In Chem.*, 2009, **21**, 2492
- 31 S. A. Shevlin and S. M. Woodley, *J. Phys. Chem. C* 2010, **114**, 17333
- 32 A. Walsh and S. M. Woodley, *Phys. Chem. Chem. Phys.*, 2010, **12**, 8446
- 33 G. Henkelman, A. Arnaldsson and H. Jónsson, *Comput. Mater. Sci.*, 2006, **36**, 254