



Activation of Water on MnOx-Nanocluster-Modified Rutile (110) and Anatase (101) TiO₂ and the Role of Cation Reduction

Rhatigan, S., & Nolan, M. (2019). Activation of Water on MnOx-Nanocluster-Modified Rutile (110) and Anatase (101) TiO₂ and the Role of Cation Reduction. *FRONTIERS IN CHEMISTRY*, 7, Article 67.
<https://doi.org/10.3389/fchem.2019.00067>

[Link to publication record in Ulster University Research Portal](#)

Published in:
FRONTIERS IN CHEMISTRY

Publication Status:
Published (in print/issue): 12/02/2019

DOI:
[10.3389/fchem.2019.00067](https://doi.org/10.3389/fchem.2019.00067)

Document Version
Publisher's PDF, also known as Version of record

Document Licence:
CC BY

General rights

The copyright and moral rights to the output are retained by the output author(s), unless otherwise stated by the document licence.

Unless otherwise stated, users are permitted to download a copy of the output for personal study or non-commercial research and are permitted to freely distribute the URL of the output. They are not permitted to alter, reproduce, distribute or make any commercial use of the output without obtaining the permission of the author(s).

If the document is licenced under Creative Commons, the rights of users of the documents can be found at <https://creativecommons.org/share-your-work/ccllicenses/>.

Take down policy

The Research Portal is Ulster University's institutional repository that provides access to Ulster's research outputs. Every effort has been made to ensure that content in the Research Portal does not infringe any person's rights, or applicable UK laws. If you discover content in the Research Portal that you believe breaches copyright or violates any law, please contact pure-support@ulster.ac.uk



Activation of Water on MnO_x-Nanocluster-Modified Rutile (110) and Anatase (101) TiO₂ and the Role of Cation Reduction

Stephen Rhatigan and Michael Nolan*

Tyndall National Institute, University College Cork, Cork, Ireland

OPEN ACCESS

Edited by:

Javier Carrasco,
CIC energigune, Spain

Reviewed by:

Jose J. Plata,
Universidad de Sevilla, Spain
Li-Min Liu,
Beihang University, China

*Correspondence:

Michael Nolan
michael.nolan@tyndall.ie

Specialty section:

This article was submitted to
Physical Chemistry and Chemical
Physics,
a section of the journal
Frontiers in Chemistry

Received: 12 October 2018

Accepted: 24 January 2019

Published: 12 February 2019

Citation:

Rhatigan S and Nolan M (2019)
Activation of Water on
MnO_x-Nanocluster-Modified Rutile
(110) and Anatase (101) TiO₂ and the
Role of Cation Reduction.
Front. Chem. 7:67.
doi: 10.3389/fchem.2019.00067

Surface modification of titania surfaces with dispersed metal oxide nanoclusters has the potential to enhance photocatalytic activity. These modifications can induce visible light absorption and suppress charge carrier recombination which are vital in improving the efficiency. We have studied heterostructures of Mn₄O₆ nanoclusters modifying the TiO₂ rutile (110) and anatase (101) surfaces using density functional theory (DFT) corrected for on-site Coulomb interactions (DFT + U). Such studies typically focus on the pristine surface, free of the point defects and surface hydroxyls present in real surfaces. In our study we have considered partial hydroxylation of the rutile and anatase surfaces and the role of cation reduction, via oxygen vacancy formation, and how this impacts on a variety of properties governing the photocatalytic performance such as nanocluster adsorption, light absorption, charge separation, and reducibility. Our results indicate that the modifiers adsorb strongly at the surface and that modification extends light absorption into the visible range. MnO_x-modified titania can show an off-stoichiometric ground state, through oxygen vacancy formation and cation reduction spontaneously, and both modified rutile and anatase are highly reducible with moderate energy costs. Manganese ions are therefore present in a mixture of oxidation states. Photoexcited electrons and holes localize at cluster metal and oxygen sites, respectively. The interaction of water at the modified surfaces depends on the stoichiometry and spontaneous dissociation to surface bound hydroxyls is favored in the presence of oxygen vacancies and reduced metal cations. Comparisons with bare TiO₂ and other TiO₂-based photocatalyst materials are presented throughout.

Keywords: TiO₂, water activation, reduction, oxygen vacancies, photocatalysis

INTRODUCTION

Photocatalysts are semiconductor materials which absorb photons of energies in excess of the bandgap to produce electron-hole pairs. These charge carriers separate and migrate to the surface of the catalyst where they drive chemical reactions *via* reduction and oxidation of adsorbed species. Photocatalysis has a variety of applications, including, but not limited to, the solar production of hydrogen from water splitting (Ni et al., 2007; Fujishima et al., 2008; Maeda and Domen, 2010; Jiang et al., 2017).

A practical photocatalyst must meet a number of criteria, such as visible light absorption, efficient charge carrier separation, stability, and active surface sites for the adsorption of feedstock species. Reducible metal cations in the catalyst can be important for enhancing the activity of the catalyst toward the difficult step of water dissociation. The development of metal oxide photocatalysts is of interest as these materials are cheap, earth abundant and, in many instances, non-toxic. Indeed, the most widely studied photocatalyst is titanium dioxide (TiO₂) (Ni et al., 2007; Fujishima et al., 2008; Dimitrijevic et al., 2011; Pelaez et al., 2012; Habisreutinger et al., 2013; Tada et al., 2014; Etacheri et al., 2015) which was first demonstrated as a photoanode for water splitting by Fujishima and Honda (1972). The large bandgap (>3 eV) means that photoactivity is restricted to the UV and has limited the real-world application of TiO₂-based photocatalyst technologies. As a result, significant scientific effort has focused on extending the light absorption edge of TiO₂ to longer wavelengths.

Substitutional doping of TiO₂ with cations and/or anions is a widely studied approach to inducing visible light absorption through the emergence of impurity-derived energy levels in the TiO₂ bandgap (Di Valentin et al., 2007; Czoska et al., 2008; Haowei et al., 2008; Ikeda et al., 2008; Gai et al., 2009; Nie et al., 2009; Valentin et al., 2009; Xu et al., 2009; Yang et al., 2009; Yu et al., 2009; Zhu et al., 2009; Long and English, 2010a,b; Zhang et al., 2010; Zheng et al., 2010; Herrmann, 2012; Etacheri et al., 2015; Li, 2015; Na Phattalung et al., 2017). First principles studies of such doped systems typically focus on bandgap reduction (Cui et al., 2008; Yang et al., 2009; Zhu et al., 2009; Long and English, 2010a,b; Zhang et al., 2010, 2014; Chand et al., 2011; Guo and Du, 2012; Na Phattalung et al., 2017) and questions of charge localization and surface reactivity are often overlooked. These are important considerations as dopant-derived defect states have been shown to act as recombination centers (Herrmann, 2012; Etacheri et al., 2015; Li, 2015) and photocatalysis is generally a surface mediated phenomenon.

Studies of the chemistry and electronic properties of surfaces and interfaces are key to understanding and screening materials for photocatalysis. The enhanced performance of the benchmark material, P25, which consists of chemically interfaced rutile and anatase phases, has been attributed to the favorable alignment of the conduction and valence bands at the interface which facilitates charge transfer between phases and the suppression of charge carrier recombination (Scanlon et al., 2013). In addition, the interface can promote the formation of active catalytic sites. This effect can be tuned by considering heterostructures with metal oxides of different compositions. Such heterostructures have been realized experimentally and shown to exhibit enhanced photocatalytic activity (Boppana and Lobo, 2011; Boppana et al., 2013; Chae et al., 2017; Sotelo-Vazquez et al., 2017; Wang et al., 2017). Nanostructuring of metal oxides has been investigated as an approach to enhancing charge transfer kinetics and increasing surface area while providing low-coordinated metal and oxygen sites for the adsorption of feedstock species (Gordon et al., 2012; Bhatia and Verma, 2017; Zhang et al., 2017a; Ong et al., 2018).

Further, nanostructuring can also facilitate the reduction of metal cations.

Surface modification of metal oxide surfaces with dispersed metal oxide nanoclusters combines the properties of hetero- and nano-structuring. Sub-nm nanoclusters of iron oxide were deposited on TiO₂ surfaces *via* chemisorption-calcination cycle (CCC) (Jin et al., 2011; Tada et al., 2014) and atomic layer deposition (ALD) (Libera et al., 2010). FeO_x-modified TiO₂ exhibited bandgap reduction and enhanced visible light photocatalytic activity. The modification was shown to suppress carrier recombination as indicated by photoluminescence spectroscopy (Jin et al., 2011). The red-shift in light absorption was attributed to cluster-derived states above the valence band maximum (VBM) which were identified by X-ray photoelectron spectroscopy (XPS) and density functional theory (DFT) simulations (Jin et al., 2011; Nolan, 2011a; Tada et al., 2014).

These studies, and the subsequent development of similar systems, (Jin et al., 2012; Boppana et al., 2013; Iwaszuk et al., 2013; Bhachu et al., 2014; Fronzi et al., 2016a) mean that a multitude of nanocluster-surface composites can be investigated. Considerations for tuning these systems for optimal performance include composition, surface termination, nanocluster size, and stoichiometry; all of which contribute to the light absorption properties, charge carrier mobility and surface reactivity. DFT simulations can be used to illuminate the properties underpinning experimental observations (Nolan, 2011a; Jin et al., 2012; Nolan et al., 2012; Iwaszuk et al., 2013) and to screen candidate materials worthy of further investigation (Park et al., 2009; Graciani et al., 2010; Nolan, 2011b, 2012, 2018; Iwaszuk and Nolan, 2013; Lucid et al., 2014; Nolan et al., 2014; Fronzi et al., 2016b; Rhatigan and Nolan, 2018a,b).

In the present study we use first principles DFT calculations to examine the photocatalytic properties of manganese oxide modified TiO₂, using model systems of Mn₄O₆-nanoclusters modifying the rutile (110) and anatase (101) surfaces and consider the role of partial surface hydroxylation in the interfacial chemistry. Our analysis includes an assessment of the stability of the composite surfaces, their ground state stoichiometry and reducibility *via* oxygen vacancy formation. Point defects, such as oxygen vacancies, are active sites at metal oxide surfaces and can be produced thermally. A more reducible surface will lose oxygen more readily and be more active in solar thermal (Muhich et al., 2016) or Mars and van Krevelen processes (1954, (Ganduglia-Pirovano et al., 2007)). Computed density of states plots elucidate the impact of modification on the light absorption properties and a model for the photoexcited state (Di Valentin and Selloni, 2011) is used to examine charge separation and localization. Finally, we study the interaction of water with the modified surfaces and focus particularly on the role of oxygen vacancies and reduced cations on water adsorption. We identify the characteristics of activation, such as dissociation, geometry distortions and charge transfer to the adsorbed species. The importance of oxygen vacancies as active sites for water dissociation at the rutile (110) surface (Schaub et al., 2001; Henderson et al., 2003) and ceria surfaces (Mullins et al., 2012) has been widely discussed and reduced Ti³⁺ ions have been shown to be active in the chemistry at titania surfaces

(Lira et al., 2011; Xiong et al., 2012). For anatase TiO₂, oxygen vacancies have been shown to be more stable at subsurface and bulk sites than on the surface (He et al., 2009; Scheiber et al., 2012). However the surface can be reduced by electron bombardment (Scheiber et al., 2012; Setvin et al., 2016) and the reaction of these vacancy sites with water and O₂ results in water dissociation. These studies highlight the necessity of engineering photocatalytic surfaces for which vacancies can be produced with moderate energy costs.

MnO_x is an interesting modifier as manganese is a multi-valent, reducible element which crystallizes in oxides with a variety of oxidation states; (Franchini et al., 2007) this will have implications for the light absorption properties and reducibility of sub-nm nanoclusters of MnO_x dispersed at the titania surfaces. We have previously studied similar systems of MnO_x-modified TiO₂, in collaboration with experiment, to interrogate their activity for CO₂ capture and reduction (Schwartzberg et al., 2017). In the present study, we focus on the potential for these catalysts to be active toward water activation. Furthermore, we investigate the impact of surface hydroxylation on the reduction of the heterostructures *via* oxygen vacancy formation and apply a model for photoexcitation to examine the associated energetics and charge localization. In reference Schwartzberg et al. (2017), the Mn₄O₆-TiO₂ composites were found to be stoichiometric in the ground state for both modified rutile and anatase, albeit with moderate costs to produce reducing oxygen vacancies (+0.59 eV for rutile and +1.1 eV for anatase). However, the impact of surface hydroxyls on the formation of oxygen vacancies was not investigated; in this paper we show that vacancy formation is in fact promoted with hydroxyls already present at the TiO₂ surfaces. The photoexcited state model, which examines localization of electrons and holes at nanocluster metal and oxygen sites, sheds light on experimental observations which suggest that the MnO_x-modifiers may facilitate recombination (Schwartzberg et al., 2017). In addition, active oxygen vacancy sites play a crucial role in the subsequent interaction of water molecules and their adsorption modes. In particular, dissociation is favored for the reduced systems; this is an important step in the water oxidation reaction.

METHODOLOGY

Periodic plane wave DFT calculations are performed using the VASP5.4 code (Kresse and Hafner, 1994; Furthmüller et al., 1996) with an energy cut-off of 400 eV. The core-valence interaction is described with projector augmented wave (PAW) potentials, (Blöchl, 1994; Kresse and Joubert, 1999) with 4 valence electrons for Ti, 6 for O, 13 for Mn and 1 for H species. The Perdew-Wang (PW91) approximation to the exchange-correlation functional is used (Perdew et al., 1996).

The TiO₂ rutile (110) and anatase (101) substrates are modeled as 18 and 12 atomic layer slabs, respectively. The bulk lattice parameters for rutile were computed as $a = 4.639$ Å and $c = 2.974$ Å, and the rutile (110) surface was modeled with a (2 × 4) surface expansion. For anatase the bulk lattice parameters are $a = 3.814$ Å and $c = 9.581$ Å and a (1 × 4) expansion was

used for the anatase (101) surface. These parameters correspond to surface areas per supercell of 13.120×11.896 Å and 10.312×15.255 Å for rutile (110) and anatase (101), respectively. The surfaces are separated from their periodic images by vacuum gaps of 20 Å, as used in our previous studies (Fronzi and Nolan, 2017; Schwartzberg et al., 2017; Nolan, 2018; Rhatigan and Nolan, 2018a,b). Γ -point sampling is used and the convergence criteria for the energy and forces are 10^{-4} eV and 0.02 eVÅ⁻², respectively. All calculations are spin polarized.

A Hubbard U correction is implemented to consistently describe the partially filled Mn 3d states and reduced Ti³⁺ states (Anisimov et al., 1991; Dudarev et al., 1998). The values of U used are U(Ti) = 4.5 eV and U(Mn) = 4 eV and these have been chosen based on previous work on TiO₂ (Morgan and Watson, 2007; Nolan et al., 2008; Iwaszuk and Nolan, 2011; Nolan, 2011a; Fronzi et al., 2016a; Fronzi and Nolan, 2017; Rhatigan and Nolan, 2018a) and manganese oxides (Franchini et al., 2007; Kitchaev et al., 2016).

To model surface hydroxylation (before the nanoclusters are adsorbed) and the impact on the heterostructure chemistry, four water molecules are dissociatively adsorbed at the clean rutile (110) and anatase (101) surfaces which gives a partial coverage of 50%. The computed energy gain when the TiO₂ surfaces are hydroxylated at half coverage is -1.03 eV per water molecule for rutile (110) and -0.8 eV for anatase (101), referenced to the total energy of four gas phase water molecules. These indicate favorable water adsorption and surface hydroxylation and these models have been used in our previous studies (Fronzi et al., 2016a; Fronzi and Nolan, 2017; Schwartzberg et al., 2017; Rhatigan and Nolan, 2018a). The nature of water molecules adsorbed at metal oxide surfaces, and in particular TiO₂ surfaces, is widely investigated both experimentally and computationally (Valdés et al., 2008; Fronzi and Nolan, 2017; Rhatigan and Nolan, 2018a) and readers are referred to reference (Mu et al., 2017) for a review of the state of the art. These models are representative of hydroxylated rutile and anatase surfaces, while we are not attempting to describe the most stable solutions for water or dissociative water adsorption at these titania surfaces (Fronzi et al., 2016a; Fronzi and Nolan, 2017; Rhatigan and Nolan, 2018a). The hydroxylated surfaces are denoted by OH-r110 and OH-a101. For the O²⁻ ions of the pristine titania surfaces, computed Bader charges are in the range of 7.3–7.4 electrons and this is our reference. After hydroxylation, Bader charges for those oxygen atoms of the surface to which H atoms are adsorbed increase to values in the range 7.6–7.7 electrons, with similar values for oxygen ions of the water-derived hydroxyls.

The Mn₄O₆ nanocluster (see **Supporting Information**) was adsorbed in different configurations at the hydroxylated rutile (110) and anatase (101) surfaces and the adsorption energies are computed using:

$$E_{ads} = E_{surf+A} - E_{surf} - E_A \quad (1)$$

where E_{surf+A} , E_{surf} and E_A are the energies of the adsorbate-surface composite system, the hydroxylated titania surface and the gas phase nanocluster, respectively.

For the reduction of the composite surface, each of the six O sites of the supported nanocluster is considered for the formation of an oxygen vacancy, O_V . One oxygen ion is removed from the Mn_4O_x cluster and the vacancy formation energy is calculated as:

$$E_{vac} = E(Mn_4O_{x-1}) + 1/2E(O_2) - E(Mn_4O_x) \quad (2)$$

where the first and third terms on the right hand side are the total energy of the cluster-surface composite with and without an oxygen vacancy and the energy is referenced to half the total energy for molecular O_2 . Having identified the most stable structure with a single O_V , the calculation is repeated for each of the five remaining O sites to determine the most stable structure with two O_V . Oxidation states are investigated with Bader charge analysis and computed spin magnetizations.

We apply a model for photoexcitation to the ground state configuration of each modified surface and to the unmodified OH-r110 and OH-a101 surfaces for comparison. This model involves imposing a triplet electronic state on the system (Di Valentin and Selloni, 2011) to promote an electron to the conduction band, with a corresponding hole in the valence band. The analysis of the energies and charge localization is discussed in more detail in the **Supporting Information**.

For the interaction of water with the modified surfaces, H_2O molecules are adsorbed in various configurations at the oxygen deficient systems and the adsorption energies are calculated as:

$$E_{ads} = E_{surf+H_2O} - E_{surf} - E_{H_2O} \quad (3)$$

where E_{surf+H_2O} , E_{surf} and E_{H_2O} refer to the energies of the H_2O molecule and modified surface in interaction, the modified surface, and the gas phase H_2O , respectively.

Oxygen atoms of the surface, cluster and surface-bound hydroxyls are denoted O_S , O_C and O_{OH} , respectively, and similar notation is adopted for OH groups. For the interaction of water with the modified surfaces, water-derived oxygen and hydroxyls are denoted O_W and OH_W .

RESULTS

Stoichiometric Mn_4O_6 -Modified TiO_2 OH-Rutile (110) and OH-Anatase (101)

Figures 1A,B show the adsorption energies and relaxed atomic structures of the stoichiometric Mn_4O_6 -nanocluster modifying the OH-r110 and OH-a101 surfaces. The large, negative adsorption energies indicate that the nanocluster-surface interaction is favorable and that the nanoclusters will be stable against desorption and aggregation (Fronzi et al., 2016b; Nolan et al., 2016; Fronzi and Nolan, 2017; Nolan, 2018; Rhatigan and Nolan, 2018a,b). For Mn_4O_6 -OH-r110 (**Figure 1A**), three Mn ions are 4-fold coordinated and to each of these is bound a terminal OH. Of these OH groups, one has migrated from a Ti site in the rutile surface to an Mn ion of the cluster (OH_{OH}) and two OH groups result from the migration of hydrogen from surface hydroxyls to O_C atoms (OH_C). The fourth Mn ion is 5-fold coordinated and is bound to three O_C and two O_S ions (one bridging O_S and one in-plane O_S). Five O ions of the OH-r110 surface bind with Mn of the nanocluster (three O_S and two

O_{OH}) and two O_C ions bind to Ti of the surface. Mn-O distances are in the range 1.8–2.1 Å; the shorter distances involve 2-fold coordinated O ions and for Mn bound to the in-plane O_S ion the Mn-O distance is 2.2 Å. Ti ions which bind to the nanocluster migrate out from the surface by 0.1 Å, however, distortions to the geometry of the rutile (110) surface are minimal.

For Mn_4O_6 -OH-a101 (**Figure 1B**), three Mn ions are 4-fold coordinated and one Mn is 5-fold coordinated. Five O_C sites are 2-fold coordinated with one O_C ion binding to three Mn ions and a H atom which has migrated from a bridging O_S site. Of the six interfacial bonds between the Mn_4O_6 nanocluster and OH-a101, three involve Mn and OH_{OH} groups; two involve Mn and bridging O_S sites and the sixth is a Ti- O_C bond. Mn-O distances are in the range 1.7–2.1 Å.

For Mn_4O_6 adsorbed at OH-r110 and OH-a101, the computed Bader charge for each of the Mn ions is 11.3 electrons, which are typical of Mn^{3+} ions (see **Table 1**). (Schwartzenberg et al., 2017) The spin magnetizations for these sites are each 3.9 μ_B , which reflects the $3d^4$ configuration of the Mn^{3+} ion.

For the Mn_4O_6 nanocluster adsorbed at OH-a101, there is an accumulation of positive charge at those O_C sites which are doubly-coordinated to Mn ions of the nanocluster. Computed Bader charges of 7.0 electrons for these O_C sites compare with 7.3–7.7 electrons computed for O^{2-} anions of the OH-a101 surface. The nanocluster-surface interaction is not as strong at the OH-a101 surface as indicated by the smaller adsorption energy. The consequence of this is that the supported nanocluster retains characteristics of the gas phase, for which the O_C ions have computed Bader charges in the range 7.0–7.1 electrons.

Reduction of Mn_4O_6 -Modified TiO_2 OH-Rutile (110) and OH-Anatase (101) Via Oxygen Vacancy Formation

The most stable modified surfaces with a single O_V are shown in **Figure 1C** for Mn_4O_5 -OH-r110 and **Figure 1D** for Mn_4O_5 -OH-a101. For the modified OH-r110 surface the formation energy of a single O_V is -0.26 eV and this formation energy indicates that O_V will form spontaneously. The next three most stable vacancy sites have formation energies in the range 0.60–0.82 eV. After formation of the most stable vacancy, two Mn ions are 3-fold coordinated and the third and fourth Mn cations are 4- and 5-fold coordinated. Two bridging and one in-plane surface oxygen are bound to Mn ions of the nanocluster. Two O_C ions bind to surface Ti sites while three O_C ions are bound only to Mn and H ions.

The formation of the neutral oxygen vacancy releases two electrons. Bader charge analysis reveals that the electrons localize at the 3-fold coordinated Mn sites of the nanocluster. The computed Bader charges on these sites increase from 11.3 to 11.5 electrons; see (**Table 1**) for computed Bader charges of reduced Ti and all Mn sites. The computed spin magnetizations are 4.6 μ_B for these Mn sites; this is typical of the formation of reduced Mn^{2+} ions which has an electronic configuration of $3d^5$.

The most favorable structure with one O_V is more stable than the second most favorable by 0.9 eV. However, the relaxed atomic structures of these configurations are very similar (compare

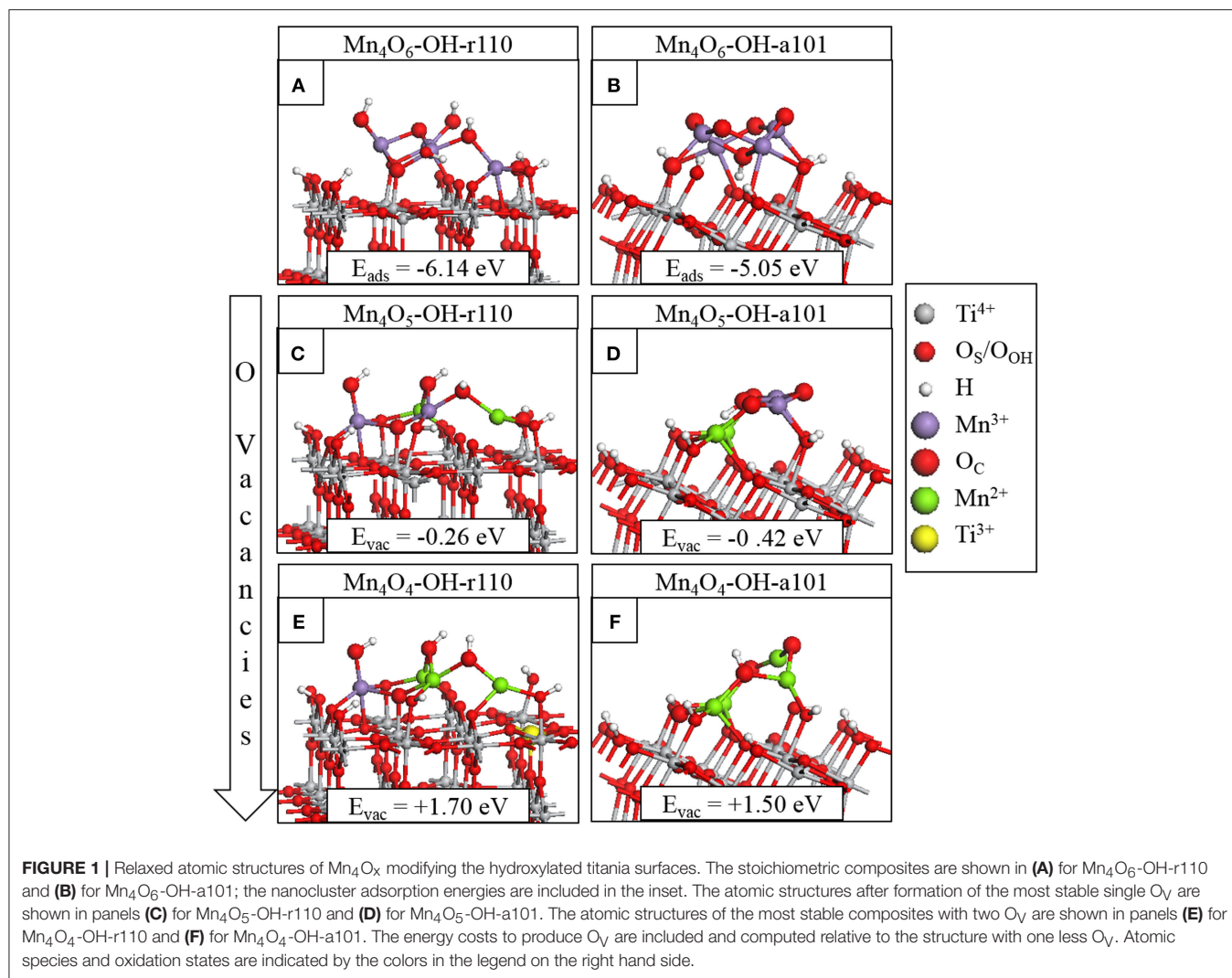


TABLE 1 | Computed Bader charges for the manganese ions in the supported nanoclusters before and after formation of one or more O_V .

Surface	OH-r110			OH-a101		
	Mn_4O_6	Mn_4O_5	Mn_4O_4	Mn_4O_6	Mn_4O_5	Mn_4O_4
Mn_I	11.3	11.5	11.5	Mn_I	11.3	11.5
Mn_{II}	11.3	11.2	11.2	Mn_{II}	11.3	11.3
Mn_{III}	11.3	11.2	11.5	Mn_{III}	11.3	11.3
Mn_{IV}	11.3	11.5	11.5	Mn_{IV}	11.3	11.5
Ti_I	1.3	1.3	1.7			

Also included are Bader charges for titanium ions of the support which are reduced after vacancy formation. Reduced Mn^{2+} and Ti^{3+} are highlighted in bold.

Figure 1C with **Figure S3A**). The difference in energy arises from the distribution of excess charge. For the O_V structure shown in **Figure S3A**, one excess charge localizes at a 5-fold coordinated surface Ti site for which the Bader charge increases from 1.3 to 1.7 electrons. A computed spin magnetization of $1.0 \mu_B$ reflects the $3d^1$ configuration of reduced Ti^{3+} .

For the modified OH-a101 surface, the most stable O_V has a formation energy of -0.42 eV which indicates that it will spontaneously form, so that the ground state is off-stoichiometric (vacancy formation energies for other sites of the nanocluster were in the range 0.5–1.3 eV). This compares with Mn_4O_6 modifying bare anatase (101) which was found to be stoichiometric in the ground state (Schwartzberg et al., 2017). After the formation of this O_V , two Mn ions relax toward the titania surface and bind with bridging O_S sites so that, in this configuration, each Mn ion is 4-fold coordinated. Bader charge analysis and computed spin densities indicate that two Mn ions are reduced to Mn^{2+} , having computed Bader charges of 11.5 electrons and computed spin magnetisations of $4.6 \mu_B$. The next most stable structure with one O_V is shown in **Figure S3B**; in this configuration three Mn ions are reduced to Mn^{2+} and this is accompanied by an accumulation of positive charge on 2-fold coordinated O_C ions for which the Bader charges were computed as 7.0 electrons.

The formation of the second O_V has a moderate energy cost for both MnO_x -modified TiO_2 surfaces, however the

modified anatase surface is reducible at a lower energy cost. Given that the anatase surface is more easily hydroxylated, (Mu et al., 2017) which these results indicate promotes vacancy formation, one would expect more O_V present on modified anatase. That O_V formation is more facile for modified anatase corroborates previous experimental work on MnO_x - TiO_2 (Schwartzenberg et al., 2017). The most stable configurations of the heterostructures with two O_V are shown in **Figure 1E** for Mn_4O_4 -OH-r110 and **Figure 1F** for Mn_4O_4 -OH-a101. For the Mn_4O_4 -OH-r110 surface, the two most stable O_C sites for formation of a second O_V had similar formation energies. One such configuration is described here and the other is included in the **Supporting Information**. For the structure shown in **Figure 1E**, the removed O_C ion was 2-fold coordinated to a cluster Mn and surface Ti ion. After vacancy formation the Mn ion binds to a bridging O_S ion and remains 3-fold coordinated. In this configuration three Mn ions are reduced; the Bader charges and spin magnetizations for these sites are 11.5 electrons and 4.6 μ_B , respectively. Similarly, for the Ti site to which the removed O_C was bound, the Bader charge and spin magnetization are 1.7 electrons and 1.0 μ_B . Hence, the Mn_4O_4 -OH-r110 heterostructure with two oxygen vacancies has one Ti^{3+} and three Mn^{2+} ions.

For the modified OH-a101 surface, a 3-fold coordinated O_C site, which forms a hydroxyl group bridging two Mn ions, has the lowest cost to produce a second O_V . One Mn ion that was bound to the removed O_C atom is 2-fold coordinated, having been originally coordinated to three O_C ions and one O_{OH} ion. The second Mn ion is 3-fold coordinated, having been 4-fold coordinated prior to vacancy formation. The H ion which was bound to the removed O_C migrates to another O_C ion. In this Mn_4O_4 -OH-a101 configuration, there are four Mn^{2+} ions, with computed Bader charges of 11.5–11.6 electrons and spin magnetizations of 4.6 μ_B .

Additional structures with two O_V are presented in **Figure S3C** for Mn_4O_4 -OH-r110 and **Figure S3D** for Mn_4O_4 -OH-a101; these are close in energy to the configurations described above, and differ in the distribution of excess charge over Mn and Ti sites. Hence, Mn and Ti sites should be present at the surface in a variety of oxidation states.

The localization of electrons at Ti and Mn sites is also accompanied by localized geometry distortions. The cation-O distances increase by ~ 0.1 Å after reduction, reflecting the larger ionic radii of Mn^{2+} and Ti^{3+} compared to Mn^{3+} and Ti^{4+} (Shannon and Prewitt, 1969).

Electronic Properties of Mn_4O_x -Modified TiO_2 OH-Rutile (110) and OH-Anatase (101)

The projected electronic density of states (PEDOS) for the heterostructures are presented in **Figure 2**. Since the heterostructures are off-stoichiometric in the ground state, the PEDOS plot for Mn_4O_6 -OH-r110 and Mn_4O_6 -OH-a101 have been omitted from this figure and are included in the **Supporting Information** for completeness. The top panels of **Figure 2** show the PEDOS of modified OH-r110 for (A) the ground state with one O_V and (B) the reduced state with two

O_V . The PEDOS plots show that occupied nanocluster-derived states (Mn 3d and O_C 2p) extend to 0.3 and 0.8 eV above the VBM of the rutile support for Mn_4O_5 - and Mn_4O_4 -OH-r110, respectively. Unoccupied Mn 3d-derived states also emerge in the titania band gap at 0.1 and 0.3 eV below the conduction band minimum (CBM) for the ground state with one O_V and the reduced state with two O_V . Additional states emerge in the band gap due to occupied Ti^{3+} states (see inset of **Figure 2B**), for the heterostructure with two O_V .

The bottom panels of **Figure 2** display the PEDOS of the modified OH-a101 surface for (C) the ground state, with one O_V , and (D) reduced state with two O_V . The PEDOS plot for the ground state, with one O_V , shows that occupied Mn 3d- and O_C 2p-derived states extend to 1.3 eV above the titania derived VBM, while unoccupied Mn 3d states emerge at 1 eV below the CBM, leading to a significant reduction in the computed energy gap relative to TiO_2 . For the reduced structure, with two O_V , each of the Mn ions is reduced to Mn^{2+} , and the highest occupied of these states is 1 eV above the VBM. The lowest energy, unoccupied state is Mn-derived and is 1 eV below the CBM. For Mn_4O_4 -OH-a101 the energy gap is 0.6 eV, with our DFT+U set-up showing a reduction over unmodified anatase.

These features of the PEDOS for Mn_4O_x - TiO_2 can be attributed to formation of interfacial bonds, the presence of low-coordinated Mn and O_C sites and the facile formation of O_V in the supported metal oxide nanocluster. Modification pushes the VBM to higher energy and results in the emergence of empty states below the CBM; these effects, and the consequent red shift, are greater for modified anatase, consistent with previous reports (Schwartzenberg et al., 2017). These metal oxide nanocluster-modified surfaces are of interest for the oxygen evolution half reaction (OER) of the water splitting process and in this context raising the VBM from that of TiO_2 and toward the water oxidation potential is a desirable effect. Lowering of the titania CBM from its favorable position straddling the water reduction potential is detrimental to the hydrogen evolution reaction (HER) activity. However, as H adsorbs too strongly at metal oxide surfaces, such heterostructures will in any case not be suitable photocathodes for water splitting.

Photoexcitation Model

We apply the model for the photoexcited state to the ground state systems, Mn_4O_5 -OH-r110 and Mn_4O_5 -OH-a101. **Table 2** presents the computed vertical, singlet-triplet and electron-hole trapping energies, as discussed in the **Supporting Information**. As can be seen from the values listed in **Table 2**, the underestimation of the bandgap inherent in approximate DFT is present in our DFT+U computational set-up. Our goal in choosing +U corrections is to consistently describe the localization of electrons and holes rather than reproduce the bandgap of bulk TiO_2 , which is not advised. Comparison of these computed energies across different structures nonetheless yields useful qualitative information about the effect of surface modification and results for the unmodified OH-r110 and OH-a101 surfaces are included for reference. In particular, $E^{vertical}$ is analogous to the optical band gap, and a reduction in this value

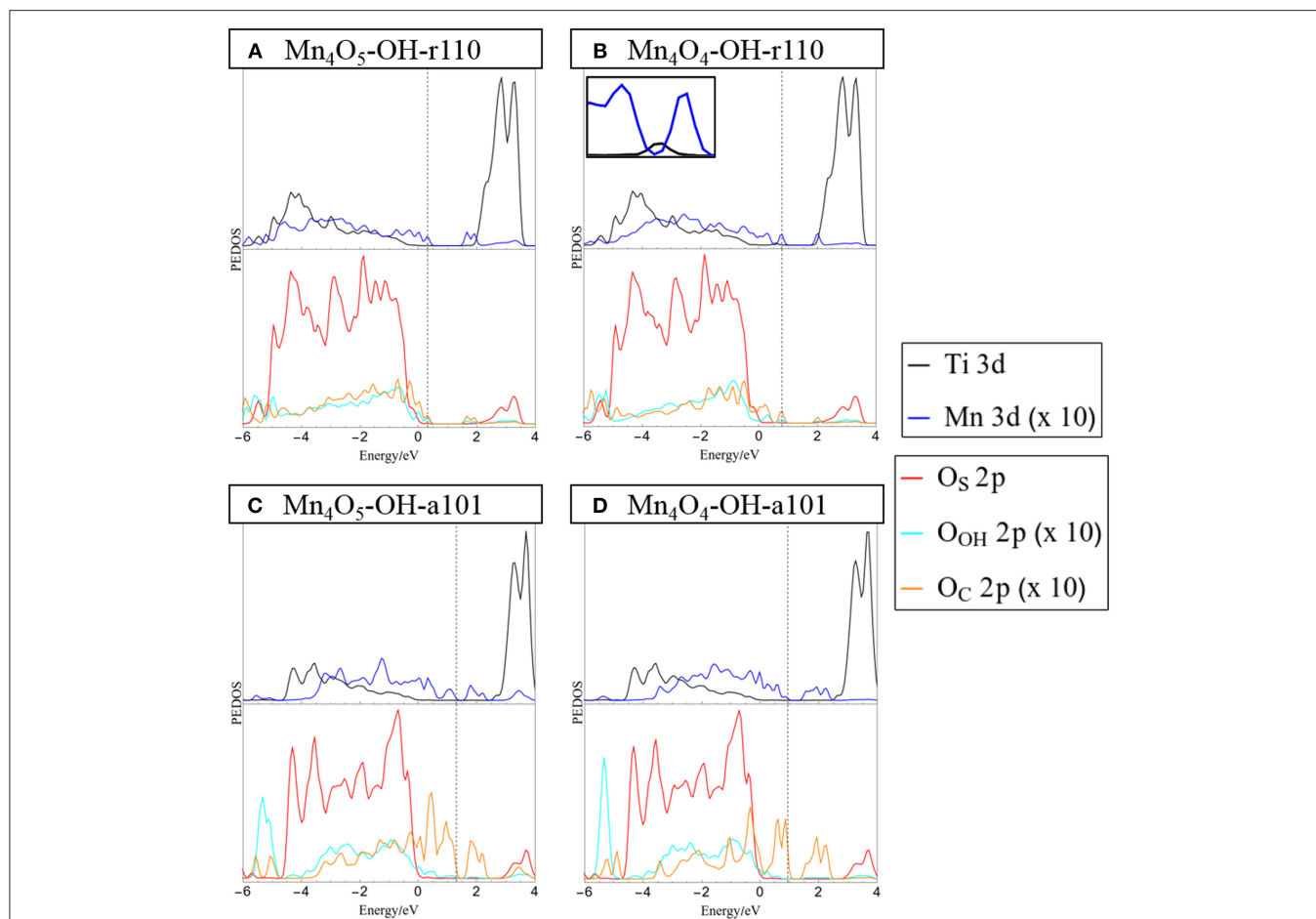


FIGURE 2 | Projected electronic density of states (PEDOS) plots for (A) Mn_4O_5 -, and (B) Mn_4O_4 -OH-r110 and (C) Mn_4O_5 -, and (D) Mn_4O_4 -OH-a101. The computed valence band max is set to 0 eV and the Fermi energy is indicated with a dashed line. The top half of each panel displays Ti and Mn 3d derived states. The bottom halves of the panels show contributions to the DOS from oxygen 2p states of the surface (O_C), surface bound hydroxyls (O_{OH}) and nanocluster (O_C). Inset in panel (B) shows the mid-gap occupied Ti 3d states in the range [0 eV, 1 eV].

TABLE 2 | Vertical singlet-triplet energy difference (E^{vertical}), the relaxed singlet-triplet energy difference (E^{excite}) and the relaxation energy (E^{relax}) for Mn_4O_5 -OH-r110 and Mn_4O_5 -OH-a101.

Composite structure	E^{vertical} (eV)	E^{excite} (eV)	E^{relax} (eV)
OH-rutile (110)	2.08	1.61	0.46
Mn_4O_5 -OH-rutile (110)	2.00	0.68	1.31
OH-anatase (101)	2.71	1.52	1.19
Mn_4O_5 -OH-anatase (101)	2.37	0.95	1.43

Values for hydroxylated rutile (110) and anatase (101) surfaces have been included for reference.

for a heterostructure relative to unmodified titania implies that modification leads to a red shift in light absorption.

When comparing Mn_4O_5 -OH-r110 with unmodified OH-r110 we can see that the values for E^{vertical} are similar, however E^{excite} is reduced by 0.93 eV for the modified surface. Comparing Mn_4O_5 -OH-a101 with unmodified OH-a101, decreases in E^{vertical} and E^{excite} by 0.34 and 0.57 eV, respectively, indicate that

modification leads to a significant red shift in light absorption. These results corroborate the analysis of the PEDOS. E^{relax} is the energy gained by the system after structural relaxation in response to the triplet electronic state and is related to the stability of the trapped electron and hole. The relaxation energy is larger for Mn_4O_5 -OH-r110 than that computed for unmodified OH-r110 (1.31 vs. 0.46 eV) and reflects the greater flexibility of the modified system in accommodating the triplet electronic state. The relaxation energies for Mn_4O_5 -OH-a101 and unmodified OH-a101 are comparable. The mixture of Mn oxidation states and the proximity of the Mn ions to each other at the anatase surface (neighboring Mn-Mn distances are in the range 2.9–3.2 Å for Mn_4O_5 -OH-a101 and 3.0–3.9 Å for Mn_4O_5 -OH-r110) restricts the degree to which the nanocluster can respond structurally to the localization of photoexcited charges.

Through analysis of Bader charges and spin magnetizations we can determine the electron and hole localization sites and the results of this analysis are represented graphically in Figure 3. For Mn_4O_5 -OH-r110, in Figures 3A,B, the electron localizes at

an Mn site; the Bader charge and spin magnetization for this site are 11.5 electrons and $4.6 \mu_B$ after electron localization, which are typical of Mn^{2+} formation. The hole localizes at an O_C site which is 2-fold coordinated to the Mn^{2+} ion and a surface Ti. In this instance the Bader charge is 6.8 electrons and the spin magnetization is $0.8 \mu_B$, which are consistent with formation of O^- . The $Mn^{2+}-O^-$ distance increases by 0.2 \AA , relative to the ground state. The $Ti-O^-$ distance decreases by 0.1 \AA .

For $Mn_4O_5-OH-a101$ (Figures 3C,D), the photoexcited electron localizes at an Mn site of the nanocluster, as confirmed by a computed Bader charge of 11.5 electrons and spin magnetization of $4.5 \mu_B$. The hole state localizes predominantly at an O_C site which bridges Mn^{2+} and Mn^{3+} ions. After hole localization the Bader charge for the O^- ion is 6.7 electrons and the spin magnetization is $0.8 \mu_B$. The $Mn^{2+}-O$ distances increase by $0.2-0.3 \text{ \AA}$.

These results show that the electron localizes at an Mn site of the supported nanocluster and the hole state localizes at a neighboring O_C site. Based on this model for the photoexcited state, we can conclude that modification does not necessarily promote the spatial separation of photoexcited charges. However, both electrons and holes will be available at the modified surface for transfer to adsorbed species.

H₂O Adsorption at Mn₄O_x-Modified OH-Rutile (110) and OH-Anatase (101)

For the interaction of water at the modified surfaces, only those composites with O_V present were considered, as such vacancies are known to be active sites at metal oxide surfaces (Schaub et al., 2001; Wang et al., 2016; Ruiz Puigdollers et al., 2017; Zhang et al., 2017b). Water adsorption is favorable at multiple sites of both modified surfaces and the geometries of the most stable adsorption configurations are displayed in Figure 4, while the Supporting Information shows other, less stable, water adsorption structures.

We adsorb water in molecular form at the heterostructures and relax the geometry. For $Mn_4O_5-OH-r110$, shown in Figure 4A, water is adsorbed exothermically in molecular form with a computed adsorption energy of -0.75 eV . In this instance, the water molecule binds to a 3-fold coordinated Mn^{2+} site with a $Mn-O_W$ distance of 2.3 \AA . Figure S5A shows dissociative water adsorption at the $Mn_4O_5-OH-r110$ surface, which has an adsorption energy of -0.31 eV . Upon dissociation, an H atom migrates to a bridging O_S site and the water-derived hydroxyl (OH_W) is singly coordinated to an Mn site with an $Mn-O_W$ distance of 1.9 \AA . The dissociation is accompanied by a transfer of charge from O_W to the nanocluster modifier, indicated by a decrease of 0.4 electrons in the computed Bader charge for the O_W ion. The Bader charges and spin magnetizations of cation sites are unchanged by the adsorption and dissociation.

Water adsorbs molecularly at $Mn_4O_5-OH-a101$, as shown in Figure 4B, with an adsorption energy of -0.74 eV . The H_2O binds to a 4-fold coordinated Mn^{3+} ion with a $Mn-O_W$ distance of 2.2 \AA . Since $Mn_4O_5-OH-r110$ and $Mn_4O_5-OH-a101$ are the ground states of the systems, the single O_V having formed spontaneously, these composites favor non-stoichiometry so that the strength of interaction with the water molecule is not

sufficient to promote spontaneous dissociation and adsorption in molecular form is favored.

The surfaces with two O_V show higher reactivity to water, as indicated by the larger adsorption energies in Figures 4C,D. Water adsorbs and spontaneously dissociates at both $Mn_4O_4-OH-r110$ (Figure 4C) and $Mn_4O_4-OH-a101$ (Figure 4D). For $Mn_4O_4-OH-r110$, the water molecule adsorbs at an O_V site. An H atom migrates to a bridging O_S site and the OH_W group is doubly coordinated to an Mn and a surface Ti site. The $Mn-O_W$ and $Ti-O_W$ distances are 2.2 \AA . Bader charge analysis reveals that 0.3 electrons are transferred from the O_W to the surface. Despite this charge transfer, the Ti ion which binds to OH_W and which was reduced to Ti^{3+} due to O_V formation prior to water adsorption, remains in the Ti^{3+} state. This agrees with work by Henderson et al. in which no charge transfer was observed between Ti^{3+} and bridging hydroxyls bound at oxygen vacancy sites at the TiO_2 rutile (110) surface (Henderson et al., 2003). The reduced Ti site was only oxidized after interaction of O_2 with the $Ti^{3+}-OH$ group. After water adsorption and dissociation the distribution of cation oxidation states is unchanged so that there are three Mn^{2+} ions and one Ti^{3+} . The Bader charge for the bridging O_S site to which the H ion binds increases from 7.3 to 7.7 electrons which, as discussed in the methodology, is typical of hydroxyl formation.

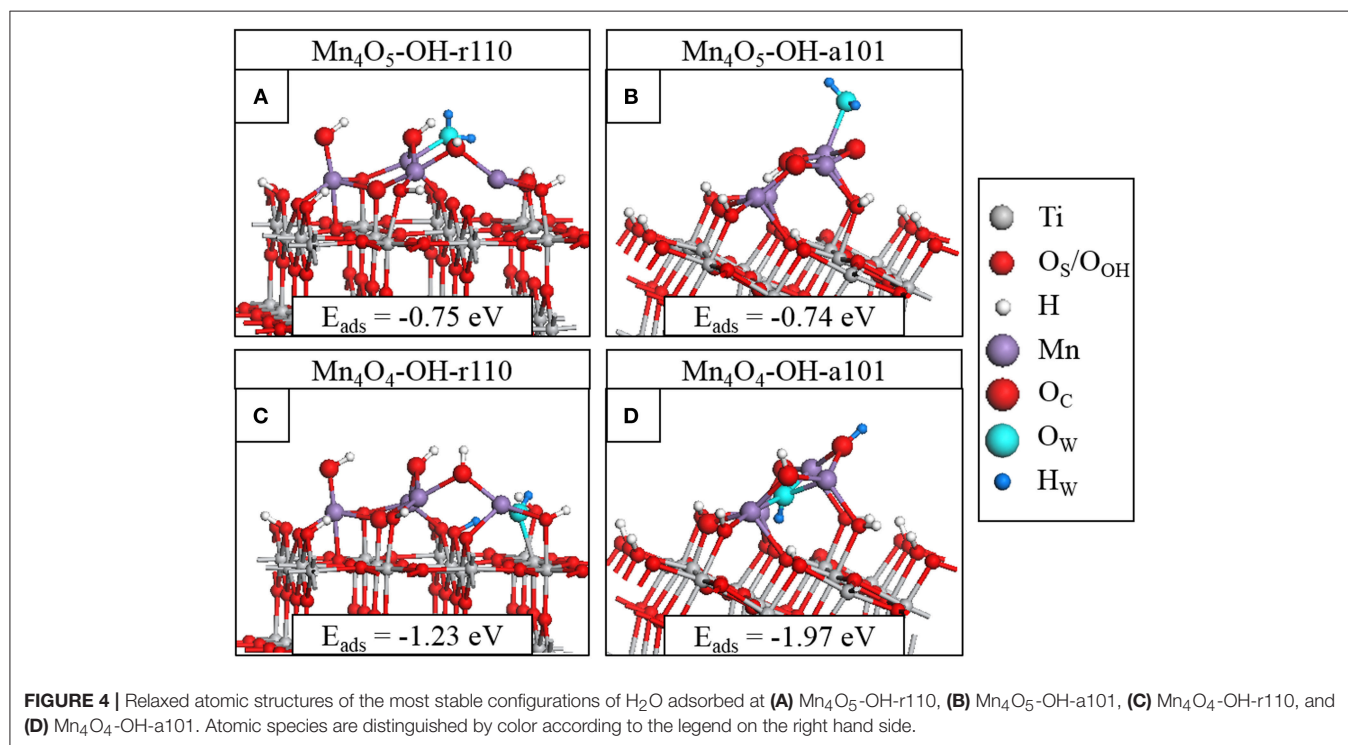
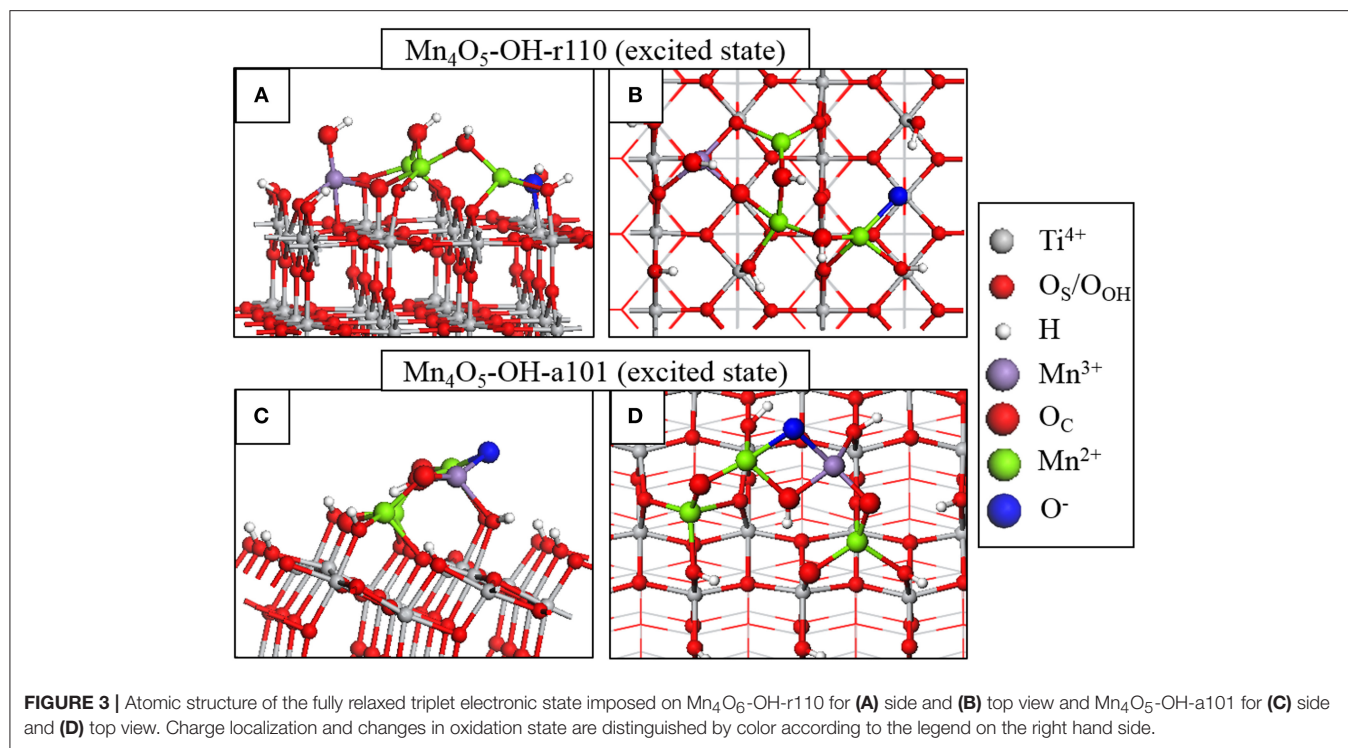
For $Mn_4O_4-OH-a101$, the water molecule adsorbs at an O_V site and after dissociation an H atom migrates to an O_C ion which shows an increase in Bader charge, from 7.1 to 7.6 electrons. The OH_W group binds to three Mn^{2+} ions; the Bader charges and spin magnetizations for cation sites are unchanged so that these ions are not involved in the charge transfer. However, for the water adsorption configuration shown in Figure S5D, an Mn^{2+} ion is re-oxidized to Mn^{3+} after dissociation of the water molecule. In this instance the adsorption energy is -1.89 eV and the OH_W group is singly-coordinated to the re-oxidized Mn ion.

CONCLUSIONS

The properties of Mn_4O_x -modified hydroxylated titania surfaces and their reduction and interaction with water depend on the phase of the TiO_2 substrate. For Mn_4O_6 adsorbed at the hydroxylated anatase (101) surface, one interfacial bond is established between a cluster oxygen ion and the surface and Mn ions bind mostly to oxygen ions of the surface bound hydroxyls. Conversely, for Mn_4O_6 at hydroxylated rutile (110), the nanocluster-surface interaction is more intimate, with Mn ions binding to bridging and in-plane oxygen ions of the rutile surface.

Our results indicate that both $Mn_4O_x-OH-r110$ and $Mn_4O_x-OH-a101$ favor non-stoichiometry, in contrast to unhydroxylated modified TiO_2 surfaces, as oxygen vacancies form spontaneously and both composites can be considered highly reducible with moderate energy costs for subsequent oxygen vacancy formation. Bader charge analysis shows that Mn ions are present in a mixture of oxidation states of at the hydroxylated surfaces. Both Mn and Ti ions are reduced in response to vacancy formation.

Modification with Mn_4O_x has a significant impact on the light absorption properties. Occupied Mn 3d states extend the



VBM of the composites to higher energies relative to that of the titania support and empty states emerge below the CBM. The consequent red shift in the light absorption edge is confirmed by our model for the photoexcited state. In particular, the vertical energy, analogous to the optical energy gap, decreases significantly for Mn₄O₅-OH-a101 relative to that computed for

the unmodified, hydroxylated anatase (101) surface. Analysis of this model shows that electrons and holes localize at Mn and neighboring O_C sites, respectively, so that modification may not promote separation of photoexcited charges, but the trapping energies of the electron and hole are quite high, suggesting high stability.

With regard to water adsorption and activation, the formation of oxygen vacancies has an impact on the strength of interaction and the most favorable adsorption mode of H₂O at the modified surfaces. For Mn₄O₅-OH-a101, with a spontaneously formed O_V, water adsorbs only in molecular form. With formation of reducing oxygen vacancies, water adsorption becomes more exothermic and leads to spontaneous dissociation to surface bound hydroxyls, similar to observations made for water interacting at reduced TiO₂ (Schaub et al., 2001; Henderson et al., 2003) and CeO₂ (Mullins et al., 2012) surfaces.

AUTHOR CONTRIBUTIONS

MN devised the research. SR performed the DFT modeling. Both authors analyzed the results and prepared, reviewed and approved the text of the paper.

REFERENCES

- Anisimov, V. I., Zaanen, J., and Andersen, O. K. (1991). Band theory and mott insulators: hubbard U instead of stoner I. *Phys. Rev. B* 44, 943–954.
- Bhachu, D. S., Sathasivam, S., Carmalt, C. J., and Parkin, I. P. (2014). PbO-modified TiO₂ thin films: a route to visible light photocatalysts. *Langmuir* 30, 624–630. doi: 10.1021/la4038777
- Bhatia, S., and Verma, N. (2017). Photocatalytic activity of ZnO nanoparticles with optimization of defects. *Mater. Res. Bull.* 95, 468–476. doi: 10.1016/j.materresbull.2017.08.019
- Blöchl, P. E. (1994). Projector augmented-wave method. *Phys. Rev. B* 50, 17953–17979.
- Boppana, V., Jiao, B. R., Newby, F., Laverock, D., Smith, J., Lobo, K. E., et al. (2013). Analysis of visible-light-active Sn(II)-TiO₂ photocatalysts. *Phys. Chem. Chem. Phys.* 15, 6185–6189. doi: 10.1039/c3cp44635b
- Boppana, V., Lobo, B. R. (2011). SnOx-ZnGa2O4 photocatalysts with enhanced visible light activity. *ACS Catal.* 1, 923–928. doi: 10.1021/cs200137h
- Chae, S. Y., Lee, C. S., Jung, H., Joo, O. S., Min, B. K., et al. (2017). Insight into charge separation in WO₃/BiVO₄ heterojunction for solar water splitting. *ACS Appl. Mater. Interfaces* 9, 19780–19790. doi: 10.1021/acsami.7b02486
- Chand, R., Obuchi, E., Katoh, K., Luitel, H. N., and Nakano, K. (2011). Enhanced photocatalytic activity of TiO₂/SiO₂ by the influence of Cu-doping under reducing calcination atmosphere. *Catal. Commun.* 13, 49–53. doi: 10.1016/j.catcom.2011.04.024
- Cui, Y., Du, H., and Wen, L. (2008). Doped-TiO₂ photocatalysts and synthesis methods to prepare TiO₂ films. *J. Mater. Sci. Technol.* 24, 675–689.
- Czoska, A. M., Livraghi, S., Chiesa, M., Giamello, E., Agnoli, S., Granozzi, G., et al. (2008). The nature of defects in fluorine-doped TiO₂. *J. Phys. Chem. C* 112, 8951–8956. doi: 10.1021/jp8004184
- Di Valentin, C., Finazzi, E., Pacchioni, G., Selloni, A., Livraghi, S., and Paganini, M. C. (2007). N-doped TiO₂: theory and experiment. *Chem. Phys.* 339, 44–56. doi: 10.1016/j.chemphys.2007.07.020
- Di Valentin, C., and Selloni, A. (2011). Bulk and surface polarons in photoexcited anatase TiO₂. *J. Phys. Chem. Lett.* 2, 2223–2228. doi: 10.1021/jz2009874
- Dimitrijevic, N. M., Vijayan, B. K., Poluektov, O. G., Rajh, T., Gray, K. A., and He, H. (2011). Role of water and carbonates in photocatalytic transformation of CO₂ to CH₄ on Titania. *J. Am. Chem. Soc.* 133, 3964–3971. doi: 10.1021/ja108791u
- Dudarev, S. L., Botton, G. A., Savrasov, S. Y., Humphreys, C. J., and Sutton, A. P. (1998). Electron-energy-loss spectra and the structural stability of nickel oxide: an LSDA+U study. *Phys. Rev. B* 57, 1505–1509. doi: 10.1103/PhysRevB.57.1505
- Etacheri, V., Di Valentin, C., Schneider, J., Bahnemann, D., and Pillai, S. C. (2015). Visible-light activation of TiO₂ photocatalysts: advances in theory

ACKNOWLEDGMENTS

We acknowledge support from Science Foundation Ireland through the US-Ireland R and D Partnerships program, Grant number SFI/US/14/E2915 and the ERA.Net for Materials Research and Innovation (M-ERA.Net 2), Horizon 2020 grant agreement number 685451, SFI Grant Number SFI/16/M-ERA/3418 (RATOCAT). We acknowledge access to SFI funded computing resources at Tyndall Institute and the SFI/HEA funded Irish Centre for High End Computing. We are grateful for support from the COST Action CM1104 Reducible Metal Oxides, Structure and Function.

SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: <https://www.frontiersin.org/articles/10.3389/fchem.2019.00067/full#supplementary-material>

- and experiments. *J. Photochem. Photobiol. C: Photochem. Rev.* 25, 1–29. doi: 10.1016/j.jphotochemrev.2015.08.003
- Franchini, C., Podloucky, R., Paier, J., Marsman, M., and Kresse, G. (2007). Ground-state properties of multivalent manganese oxides: density functional and hybrid density functional calculations. *Phys. Rev. B* 75:195128. doi: 10.1103/PhysRevB.75.195128
- Fronzi, M., Daly, W., and Nolan, M. (2016b). Reactivity of metal oxide nanocluster modified rutile and anatase TiO₂: oxygen vacancy formation and CO₂ interaction. *App. Catal. A* 521, 240–249. doi: 10.1016/j.apcata.2015.11.038
- Fronzi, M., Iwaszuk, A., Lucid, A., and Nolan, M. (2016a). Metal oxide nanocluster-modified TiO₂ as solar activated photocatalyst materials. *J. Phys.: Condens. Matter* 28, 074006. doi: 10.1088/0953-8984/28/7/074006
- Fronzi, M., and Nolan, M. (2017). Surface modification of perfect and hydroxylated TiO₂ rutile (110) and anatase (101) with chromium oxide nanoclusters. *ACS Omega* 2, 6795–6808. doi: 10.1021/acsomega.7b01118
- Fujishima, A., and Honda, K. (1972). Electrochemical photolysis of water at a semiconductor electrode. *Nature* 238, 37–38. doi: 10.1038/238037a0
- Fujishima, A., Zhang, X., and Tryk, D. A. (2008). TiO₂ photocatalysis and related surface phenomena. *Surf. Sci. Rep.* 63, 515–582. doi: 10.1016/j.surfrep.2008.10.001
- Furthmüller, J., Hafner, J., and Kresse, G. (1996). Dimer reconstruction and electronic surface states on clean and hydrogenated diamond (100) surfaces. *Phys. Rev. B* 53, 7334–7351.
- Gai, Y., Li, J., Li, S. S., Xia, J. B., and Wei, S. H. (2009). Design of narrow-gap TiO₂: a passivated codoping approach for enhanced photoelectrochemical activity. *Phys. Rev. Lett.* 102:036402. doi: 10.1103/PhysRevLett.102.036402
- Ganduglia-Pirovano, M. V., Hofmann, A., and Sauer, J. (2007). Oxygen vacancies in transition metal and rare earth oxides: current state of understanding and remaining challenges. *Surf. Sci. Rep.* 62, 219–270. doi: 10.1016/j.surfrep.2007.03.002
- Gordon, T. R., Cargnello, M., Paik, T., Mangolini, F., Weber, R. T., Fornasiero, P., et al. (2012). Nonaqueous synthesis of TiO₂ nanocrystals using TiF₄ to engineer morphology, oxygen vacancy concentration, and photocatalytic activity. *J. Am. Chem. Soc.* 134, 6751–6761. doi: 10.1021/ja300823a
- Graciani, J., Plata, J. J., Sanz, J. F., Liu, P., and Rodriguez, J. A. (2010). A theoretical insight into the catalytic effect of a mixed-metal oxide at the nanometer level: the case of the highly active metal/CeO_x/TiO₂(110) catalysts. *J. Chem. Phys.* 132:104703. doi: 10.1063/1.3337918
- Guo, M., and Du, J. (2012). First-principles study of electronic structures and optical properties of Cu, Ag, and Au-doped anatase TiO₂. *Physica B* 407, 1003–1007. doi: 10.1016/j.physb.2011.12.128
- Habisreutinger, S. N., Schmidt-Mende, L., and Stolarczyk, J. K. (2013). Photocatalytic reduction of CO₂ on TiO₂ and other semiconductors. *Angew. Chem. Int. Ed.* 52, 7372–7408. doi: 10.1002/anie.201207199

- Haowei, P., Jingbo, L., Shu-Shen, L., and Jian-Bai, X. (2008). First-principles study of the electronic structures and magnetic properties of 3d transition metal-doped anatase TiO₂. *J. Phys.* 20:125207. doi: 10.1088/0953-8984/20/12/125207
- He, Y., Dulub, O., Cheng, H., Selloni, A., and Diebold, U. (2009). Evidence for the predominance of subsurface defects on reduced anatase TiO₂(101). *Phys. Rev. Lett.* 102:106105. doi: 10.1103/PhysRevLett.102.106105
- Henderson, M. A., Epling, W. S., Peden, C. H. F., and Perkins, C. L. (2003). Insights into photoexcited electron scavenging processes on TiO₂ obtained from studies of the reaction of O₂ with OH groups adsorbed at electronic defects on TiO₂(110). *J. Phys. Chem. B.* 107, 534–545. doi: 10.1021/jp0262113
- Herrmann, J. M. (2012). Detrimental cationic doping of titania in photocatalysis: why chromium Cr³⁺-doping is a catastrophe for photocatalysis, both under UV- and visible irradiations. *New J. Chem.* 36, 883–890. doi: 10.1039/c2nj20914d
- Ikeda, T., Nomoto, T., Eda, K., Mizutani, Y., Kato, H., Kudo, A., et al. (2008). Photoinduced dynamics of TiO₂ doped with Cr and Sb. *J. Phys. Chem. C.* 112, 1167–1173. doi: 10.1021/jp0752264
- Iwaszuk, A., and Nolan, M. (2011). Reactivity of sub 1 nm supported clusters: (TiO₂)_n clusters supported on rutile TiO₂ (110). *Phys. Chem. Chem. Phys.* 13, 4963–4973. doi: 10.1039/C0CP02030C
- Iwaszuk, A., and Nolan, M. (2013). Lead oxide-modified TiO₂ photocatalyst: tuning light absorption and charge carrier separation by lead oxidation state. *Catal. Sci. Technol.* 3, 2000–2008. doi: 10.1039/c3cy00194f
- Iwaszuk, A., Nolan, M., Jin, Q., Fujishima, M., and Tada, H. (2013). Origin of the visible-light response of nickel(ii) oxide cluster surface modified titanium(iv) dioxide. *J. Phys. Chem. C.* 117, 2709–2718. doi: 10.1021/jp306793r
- Jiang, C., Moniz, S. J. A., Wang, A., Zhang, T., and Tang, J. (2017). Photoelectrochemical devices for solar water splitting - materials and challenges. *Chem. Soc. Rev.* 46, 4645–4660. doi: 10.1039/C6CS00306K
- Jin, Q., Fujishima, M., Nolan, M., Iwaszuk, A., and Tada, H. (2012). Photocatalytic activities of tin(iv) oxide surface-modified titanium(iv) dioxide show a strong sensitivity to the TiO₂ crystal form. *J. Phys. Chem. C.* 116, 12621–12626. doi: 10.1021/jp302493f
- Jin, Q., Fujishima, M., and Tada, H. (2011). Visible-light-active iron oxide-modified anatase titanium(iv) dioxide. *J. Phys. Chem. C.* 115, 6478–6483. doi: 10.1021/jp201131t
- Kitchaev, D. A., Peng, H., Liu, Y., Sun, J., Perdew, J. P., and Ceder, G. (2016). Energetics of MnO₂ polymorphs in density functional theory. *Phys. Rev. B.* 93:045132. doi: 10.1103/PhysRevB.93.045132
- Kresse, G., and Hafner, J. (1994). *Ab initio* molecular-dynamics simulation of the liquid-metal-amorphous-semiconductor transition in germanium. *Phys. Rev. B.* 49, 14251–14269.
- Kresse, G., and Joubert, D. (1999). From ultrasoft pseudopotentials to the projector augmented-wave method. *Phys. Rev. B.* 59, 1758–1775.
- Li, W. (2015). Influence of electronic structures of doped TiO₂ on their photocatalysis. *Phys. Status Solidi (RRL) - Rapid Res. Lett.* 9, 10–27. doi: 10.1002/pssr.201409365
- Libera, J. A., Elam, J. W., Sather, N. F., Rajh, T., and Dimitrijevic, N. M. (2010). Iron(iii)-oxo centers on TiO₂ for visible-light photocatalysis. *Chem. Mater.* 22, 409–413. doi: 10.1021/cm902825c
- Lira, E., Wendt, S., Huo, P., Hansen, J. Ø., Streber, R., Porsgaard, S., et al. (2011). The importance of Bulk Ti³⁺ defects in the oxygen chemistry on titania surfaces. *J. Am. Chem. Soc.* 133, 6529–6532. doi: 10.1021/ja200884w
- Long, R., and English, N. J. (2010a). First-principles calculation of synergistic (N, P)-codoping effects on the visible-light photocatalytic activity of anatase TiO₂. *J. Phys. Chem. C.* 114, 11984–11990. doi: 10.1021/jp100802r
- Long, R., and English, N. J. (2010b). Synergistic effects on band gap-narrowing in titania by codoping from first-principles calculations. *Chem. Mater.* 22, 1616–1623. doi: 10.1021/cm903688z
- Lucid, A., Iwaszuk, A., and Nolan, M. (2014). A first principles investigation of Bi₂O₃-modified TiO₂ for visible light activated photocatalysis: the role of TiO₂ crystal form and the Bi³⁺ stereochemical lone pair. *Mater. Sci. Semicond. Process.* 25, 59–67. doi: 10.1016/j.mssp.2014.01.005
- Maeda, K., and Domen, K. (2010). Photocatalytic water splitting: recent progress and future challenges. *J. Phys. Chem. Lett.* 1, 2655–2661. doi: 10.1021/jz1007966
- Mars, P., and van Krevelen, D. W. (1954). Oxidation carried out by means of vanadium oxide catalysts. *Chem. Eng. Sci.* 3, 41–59.
- Morgan, B. J., and Watson, G. W. (2007). A DFT + U description of oxygen vacancies at the TiO₂ rutile (1 1 0) surface. *Surf. Sci.* 601, 5034–5041.
- Mu, R., Zhao, Z.-J., Dohnálek, Z., and Gong, J. (2017). Structural motifs of water on metal oxide surfaces. *Chem. Soc. Rev.* 46, 1785–1806. doi: 10.1039/C6CS00864J
- Muhich, C. L., Ehrhart, B. D., Al-Shankiti, I., Ward, B. J., Musgrave, C. B., and Weimer, A. W. (2016). A review and perspective of efficient hydrogen generation via solar thermal water splitting. *Wiley Interdis. Rev.* 5, 261–287. doi: 10.1002/wene.174
- Mullins, D. R., Albrecht, P. M., Chen, L., Calaza, F. C., Biegalski, M. D., Overbury, S. H., et al. (2012). Water dissociation on CeO₂(100) and CeO₂(111) thin films. *J. Phys. Chem. C.* 116, 19419–19428. doi: 10.1021/jp306444h
- Na Phattalung, S., Limpijumngong, S., and Yu, J. (2017). Passivated co-doping approach to bandgap narrowing of titanium dioxide with enhanced photocatalytic activity. *Appl. Catal. B.* 200, 1–9. doi: 10.1016/j.apcatb.2016.06.054
- Ni, M., Leung, M., Leung, K. H., and Sumathy, K. (2007). A review and recent developments in photocatalytic water-splitting using TiO₂ for hydrogen production. *Renew. Sustain. Energy Rev.* 11, 401–425. doi: 10.1016/j.rser.2005.01.009
- Nie, X., Zhuo, S., Maeng, G., and Sohlberg, K. (2009). Doping of polymorphs for altered optical and photocatalytic properties. *Int. J. Photoenergy.* 2009:294042. doi: 10.1155/2009/294042
- Nolan, M. (2011a). Electronic coupling in iron oxide-modified TiO₂ leads to a reduced band gap and charge separation for visible light active photocatalysis. *Phys. Chem. Chem. Phys.* 13, 18194–18199. doi: 10.1039/c1cp21418g
- Nolan, M. (2011b). Surface modification of TiO₂ with metal oxide nanoclusters: a route to composite photocatalytic materials. *Chem. Commun.* 47, 8617–8619. doi: 10.1039/c1cc13243a
- Nolan, M. (2012). First-principles prediction of new photocatalyst materials with visible-light absorption and improved charge separation: surface modification of rutile TiO₂ with nanoclusters of MgO and Ga₂O₃. *ACS Appl. Mater. Interfaces.* 4, 5863–5871. doi: 10.1021/am301516c
- Nolan, M. (2018). Alkaline earth metal oxide nanocluster modification of rutile TiO₂ (110) promotes water activation and CO₂ chemisorption. *J. Mater. Chem. A.* 6, 9451–9466. doi: 10.1039/C8TA01789A
- Nolan, M., Elliott, S. D., Mulley, J. S., Bennett, R. A., Basham, M., and Mulheran, P. (2008). Electronic structure of point defects in controlled self-doping of the TiO₂ (110) surface: combined photoemission spectroscopy and density functional theory study. *Phys. Rev. B.* 77:235424. doi: 10.1103/PhysRevB.77.235424
- Nolan, M., Iwaszuk, A., and Gray, K. A. (2014). Localization of photoexcited electrons and holes on low coordinated Ti and O sites in free and supported TiO₂ nanoclusters. *J. Phys. Chem. C.* 118, 27890–27900. doi: 10.1021/jp508822v
- Nolan, M., Iwaszuk, A., Lucid, A. K., Carey, J. J., and Fronzi, M. (2016). Design of novel visible light active photocatalysts: surface modified TiO₂. *Adv. Mater.* 28, 5425–5446. doi: 10.1002/adma.201504894
- Nolan, M., Iwaszuk, A., and Tada, H. (2012). Molecular metal oxide cluster-surface modified titanium dioxide photocatalysts. *Aust. J. Chem.* 65, 624–632. doi: 10.1071/CH11451
- Ong, C. B., Ng, L. Y., and Mohammad, A., W. (2018). A review of ZnO nanoparticulates as solar photocatalysts: synthesis, mechanisms and applications. *Renew. Sustain. Energy Rev.* 81, 536–551. doi: 10.1016/j.rser.2017.08.020
- Park, J. B., Graciani, J., Evans, J., Stacchiola, D., Ma, S., Liu, P., et al. (2009). High catalytic activity of Au/CeO_x/TiO₂ (110) controlled by the nature of the mixed-metal oxide at the nanometer level. *Proc. Natl. Acad. Sci. U. S. A.* 106, 4975–4980. doi: 10.1073/pnas.0812604106
- Pelaez, M., Nolan, N. T., Pillai, S. C., Seery, M. K., Falaras, P., Kontos, A. G., et al. (2012). A review on the visible light active titanium dioxide photocatalysts for environmental applications. *Appl. Catal. B.* 125, 331–349. doi: 10.1016/j.apcatb.2012.05.036
- Perdew, J. P., Burke, K., and Ernzerhof, M. (1996). Generalized gradient approximation made simple. *Phys. Rev. Lett.* 77, 3865–3868.
- Rhatigan, S., and Nolan, M. (2018a). Impact of surface hydroxylation in MgO-/SnO-nanocluster modified TiO₂ anatase (101) composites on visible light absorption, charge separation and reducibility. *Chin. Chem. Lett.* 29, 757–764. doi: 10.1016/j.ccl.2017.11.036

- Rhatigan, S., and Nolan, M. (2018b). CO₂ and water activation on ceria nanocluster modified TiO₂ rutile (110). *J. Mater. Chem. A*, 6, 9139–9152. doi: 10.1039/C8TA01270A
- Ruiz Puigdollers, A., Schlexer, P., Tosoni, S., and Pacchioni, G. (2017). Increasing oxide reducibility: the role of metal/oxide interfaces in the formation of oxygen vacancies. *ACS Catal.* 7, 6493–6513. doi: 10.1021/acscatal.7b01913
- Scanlon, D. O., Dunnill, C. W., Buckleridge, J., Shevlin, S. A., Logsdail, A. J., Woodley, S. M., et al. (2013). Band alignment of rutile and anatase TiO₂. *Nat. Mater.* 12, 798–801. doi: 10.1038/nmat3697
- Schaub, R., Thostrup, P., Lopez, N., Lægsgaard, E., Stensgaard, I., and Nørskov, J. K. (2001). Oxygen vacancies as active sites for water dissociation on rutile TiO₂(110). *Phys. Rev. Lett.* 87, 266104. doi: 10.1103/PhysRevLett.87.266104
- Scheiber, P., Fidler, M., Dulub, O., Schmid, M., Diebold, U., Hou, W., et al. (2012). (Sub)Surface Mobility of Oxygen Vacancies at the TiO₂ Anatase (101) Surface. *Phys. Rev. Lett.* 109:136103. doi: 10.1103/PhysRevLett.109.136103
- Schwartzberg, K. C., Hamilton, J., Lucid, W. J., Weitz, A. K., Notestein, E., Gray, K. A., et al. (2017). Multifunctional photo/thermal catalysts for the reduction of carbon dioxide. *Catal. Today* 280, 65–73. doi: 10.1016/j.cattod.2016.06.002
- Setvin, M., Aschauer, U., Hulva, J., Simschitz, T., Daniel, B., Schmid, M., et al. (2016). Following the reduction of oxygen on TiO₂ anatase (101) step by step. *J. Am. Chem. Soc.* 138, 9565–9571. doi: 10.1021/jacs.6b04004
- Shannon, R. D., and Prewitt, C. T. (1969). Effective ionic radii in oxides and fluorides. *Acta Crystallogr. Sec. B*, 25, 925–946. doi: 10.1107/S0567740869003220
- Sotelo-Vazquez, C., Quesada-Cabrera, R., Ling, M., Scanlon, D. O., Kafzas, A., Thakur, P. K., et al. (2017). Evidence and effect of photogenerated charge transfer for enhanced photocatalysis in WO₃/TiO₂ heterojunction films: a computational and experimental study. *Adv. Funct. Mater.* 27:1605413. doi: 10.1002/adfm.201605413
- Tada, H., Jin, Q., Iwaszuk, A., and Nolan, M. (2014). Molecular-scale transition metal oxide nanocluster surface-modified titanium dioxide as solar-activated environmental catalysts. *J. Phys. Chem. C*, 118, 12077–12086. doi: 10.1021/jp412312m
- Valdés, Á., Qu, Z. W., Kroes, G. J., Rossmel, J., and Nørskov, J. K. (2008). Oxidation and photo-oxidation of water on TiO₂ surface. *J. Phys. Chem. C*, 112, 9872–9879. doi: 10.1021/jp711929d
- Valentin, C. D., Pacchioni, G., Onishi, H., and Kudo, A. (2009). Cr/Sb codoped TiO₂ from first principles calculations. *Chem. Phys. Lett.* 469, 166–171. doi: 10.1016/j.cplett.2008.12.086
- Wang, F., Wei, S., Zhang, Z., Patzke, G. R., and Zhou, Y. (2016). Oxygen vacancies as active sites for H₂S dissociation on the rutile TiO₂(110) surface: a first-principles study. *Phys. Chem. Chem. Phys.* 18, 6706–6712. doi: 10.1039/C5CP06835E
- Wang, J., Li, H., Meng, S., Zhang, L., Fu, X., and Chen, S. (2017). One-pot hydrothermal synthesis of highly efficient SnO_x/Zn₂SnO₄ composite photocatalyst for the degradation of methyl orange and gaseous benzene. *Appl. Catal. B*, 200, 19–30. doi: 10.1016/j.apcatb.2016.06.070
- Xiong, L. B., Li, L., Yang, B., and Yu, Y. (2012). Ti³⁺ in the surface of titanium dioxide: generation, properties and photocatalytic application. *J. Nanomater.* 2013:831524. doi: 10.1155/2012/831524
- Xu, J.-, Li, P., Lv, L., Zhang, Y., Chen, S., Du, W., et al. (2009). Structural and magnetic properties of Fe-doped anatase TiO₂ films annealed in vacuum. *Chin. Phys. Lett.* 26:097502. doi: 10.1088/0256-307X/26/9/097502
- Yang, K., Dai, Y., Huang, B., and Whangbo, M.- H. (2009). Density functional characterization of the visible-light absorption in substitutional C-anion- and C-cation-doped TiO₂. *J. Phys. Chem. C*, 113, 2624–2629. doi: 10.1021/jp808483a
- Yu, J., Xiang, Q., and Zhou, M. (2009). Preparation, characterization and visible-light-driven photocatalytic activity of Fe-doped titania nanorods and first-principles study for electronic structures. *Appl. Catal. B*, 90, 595–602. doi: 10.1016/j.apcatb.2009.04.021
- Zhang, H., Yu, X., McLeod, J. A., and Sun, X. (2014). First-principles study of Cu-doping and oxygen vacancy effects on TiO₂ for water splitting. *Chem. Phys. Lett.* 612, 106–110. doi: 10.1016/j.cplett.2014.08.003
- Zhang, J., Pan, C., Fang, P., Wei, J., and Xiong, R. (2010). Mo + C codoped TiO₂ using thermal oxidation for enhancing photocatalytic activity. *ACS Appl. Mater. Interf.* 2, 1173–1176. doi: 10.1021/am100011c
- Zhang, J., Salles, I., Pering, S., Cameron, P. J., Mattia, D., and Eslava, S. (2017a). Nanostructured WO₃ photoanodes for efficient water splitting via anodisation in citric acid. *RSC Adv.* 7, 35221–35227. doi: 10.1039/C7RA05342H
- Zhang, Y., Dai, R., and Hu, S. (2017b). Study of the role of oxygen vacancies as active sites in reduced graphene oxide-modified TiO₂. *Phys. Chem. Chem. Phys.* 19, 7307–7315. doi: 10.1039/C7CP00630F
- Zheng, J. W., Bhattacharyya, A., Wu, P., Chen, Z., Highfield, J., Dong, Z., et al. (2010). The origin of visible light absorption in chalcogen element (S, Se, and Te)-doped anatase TiO₂ photocatalysts. *J. Phys. Chem. C*, 114, 7063–7069. doi: 10.1021/jp9115035
- Zhu, W., Qiu, X., Iancu, V., Chen, X. Q., Pan, H., Zhang, Z., et al. (2009). Band gap narrowing of titanium oxide semiconductors by noncompensated anion-cation codoping for enhanced visible-light photoactivity. *Phys. Rev. Lett.* 103:226401. doi: 10.1103/PhysRevLett.103.226401

Conflict of Interest Statement: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

Copyright © 2019 Rhatigan and Nolan. This is an open-access article distributed under the terms of the Creative Commons Attribution License (CC BY). The use, distribution or reproduction in other forums is permitted, provided the original author(s) and the copyright owner(s) are credited and that the original publication in this journal is cited, in accordance with accepted academic practice. No use, distribution or reproduction is permitted which does not comply with these terms.