

# SCIENTIFIC REPORTS



OPEN

## A pathway of nanocrystallite fabrication by photo-assisted growth in pure water

Received: 10 February 2015

Accepted: 26 May 2015

Published: 16 June 2015

Melbert Jeem<sup>1</sup>, Muhammad Rafiq Mirza bin Julaihi<sup>1</sup>, Junya Ishioka<sup>2</sup>, Shigeo Yatsu<sup>1,2</sup>, Kazumasa Okamoto<sup>1,2</sup>, Tamaki Shibayama<sup>1,2</sup>, Tomio Iwasaki<sup>3</sup>, Takahiko Kato<sup>2,3</sup> & Seiichi Watanabe<sup>2,2</sup>

We report a new production pathway for a variety of metal oxide nanocrystallites via submerged illumination in water: submerged photosynthesis of crystallites (SPSC). Similar to the growth of green plants by photosynthesis, nanocrystallites shaped as nanoflowers and nanorods are hereby shown to grow at the protruded surfaces via illumination in pure, neutral water. The process is photocatalytic, accompanied with hydroxyl radical generation via water splitting; hydrogen gas is generated in some cases, which indicates potential for application in green technologies. Together with the aid of *ab initio* calculation, it turns out that the nanobumped surface, as well as aqueous ambience and illumination are essential for the SPSC method. Therefore, SPSC is a surfactant-free, low-temperature technique for metal oxide nanocrystallites fabrication.

New approaches to manufacturing the nanocrystallites of metallic oxides are desired due to their emerging applications in a wide range of high-technology applications<sup>1–8</sup>. In the case of liquid-solid phase crystallisation studies, the surface morphology control plays an essential role in influencing the nucleation of nanocrystallites<sup>9</sup>. Recent studies have demonstrated the role of surface chemistry and morphology utilizing various mechanisms gained from polymeric substrates<sup>10,11</sup>. Achieving those understandings provides a powerful means to widespread reports in nanocrystallites research.

Our interest is in the easy feasible design of nanocrystallites fabrication beneficial for the nanotechnology and environment. With regard to obtaining the desired products, we herein report a new pathway of production for variety of metal oxides nanocrystallites via SPSC. We chose various metals (Zn, W, Cu and Ce) to demonstrate the effectiveness of the submerged photosynthesis of crystallites (SPSC) method in obtaining a variety of morphologies of metal oxide nanocrystallites (NCs). The present report primarily focuses on ZnO nanofabrication due to its promising environmental<sup>12</sup> applications and broad range of modern device applications, including light-emitting diodes<sup>13</sup>, photo-detectors<sup>14</sup>, gas sensors<sup>15</sup>, and solar cells<sup>16</sup>.

The functions of these semiconductive ZnO (with a wide bandgap of 3.37 eV) devices are crucially dependent on the nanostructure morphology. Hence, it is important to tune and stabilise the syntheses parameter for improved performance. To this end, there has been a substantial increase in the number of reports on hydrothermally synthesised ZnO nanostructures<sup>17,18</sup>, including lasing<sup>1</sup> and the addition of metal-ion impurity<sup>19</sup> techniques. Expanding on these efforts, this study develops a rational and environmentally benign approach to synthesise a plethora of ZnO nanomorphologies.

We employed our initial metal surface treatment based on the utilisation of the submerged liquid plasma process<sup>20,21</sup>. The reaction of plasma in an aqueous solution facilitates the synthesis of metal oxide powder<sup>22,23</sup>. Hence, we adopted this technique for the direct (one-step) synthesis of ZnO “seeds”. Our intention was to create a semiconductive reformed layer with protruding characteristics (nanobumps) on

<sup>1</sup>Graduate School of Engineering, Hokkaido University, N13, W8, Kita-ku, Sapporo, Hokkaido 060-8628, Japan.

<sup>2</sup>Faculty of Engineering, Hokkaido University, N13, W8, Kita-ku, Sapporo, Hokkaido 060-8628, Japan. <sup>3</sup>Hitachi Research Laboratory, Hitachi Ltd., 7-1-1 Omika, Hitachi, Ibaraki, 319-1292, Japan. Correspondence and requests for materials should be addressed to S.W. (email: swoo4@eng.hokudai.ac.jp)

the material. For this purpose, a raw Zn metallic plate was used as the target material for the formation of nanobumps. The NC growth was then completed by a “photosynthesis” reaction, where the irradiation of UV light (typically  $\lambda = 365$  nm and  $I = 28$  mWcm<sup>-2</sup>) on the nanobumps inside pure water assisted the growth of ZnO NCs.

The SPSC method is different from previous nanofabrication reports<sup>24,25</sup>, in which a hydrothermal decomposition process using UV-Vis light irradiation directed the formation of NCs. Instead, we found that the illumination in ultrapure water induced an apical growth characteristic via water radiolysis, as will be described later. Furthermore, we were able to demonstrate working NC fabrication at room temperature (RT) under surfactant- and contamination-free conditions by eliminating the need for organometallic or other organic solution phases. Similar to the growth of green plants via botanical photosynthesis, our SPSC method requires only light, water and nanobumps (crystallite seeds). Interestingly, the SPSC end products were accompanied by the generation of hydrogen gas, which gives rise to a potential application of this methodology as a green technology for energy, chemistry and nanotechnology.

## Results

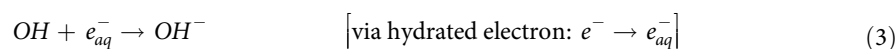
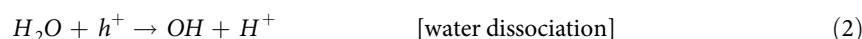
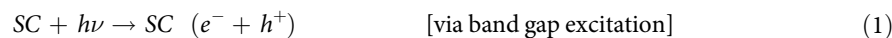
Figure 1(a) presents a scanning electron microscopy (SEM) image of ZnO nanobumps that were tailored by submerged liquid plasma treatment and depicts the metal surface having average of two or more protrusions per 10  $\mu\text{m}^2$ . The protruded surface exhibits an average diameter of 1  $\mu\text{m}$  or less. The higher-magnified image indicates the small ZnO seeds homogeneously localised on the protruded surface; these seeds have an average diameter of approximately 20 nm. Here, a drastic change in the seed clustering was observed after subsequent UV irradiation in ultrapure water and ambient temperature: widely spread ZnO NCs covered the metal substrate. The previously localised seeds grew outward to form a bunch of nanorods that formed into nanoflowers (dandelion-like), dendrites (tree-like), and aligned nanorods (lawn grass-like) (Fig. 1(b–d), respectively). The SEM-energy-dispersive X-ray spectroscopy (EDS) analysis (Fig. 2(a,b)) and X-ray diffraction (XRD) analysis (Fig. 2(c)) for the ZnO NCs illustrated that the NCs were synthesised on a Zn substrate, which contained Zn and O. As evidenced by the selected area electron diffraction (SAED) pattern and high-resolution transmission electron micrograph (HRTEM) of a nanorod examined along the [1 $\bar{1}$ 0] axis (Fig. 2(d)), the nanorods were single crystallites (a wurtzite structure). The apical growth direction was in the *c*-axis  $\langle 001 \rangle$ . This result is consistent with previous ZnO crystal growth reports<sup>24,26,27</sup>.

## Discussion

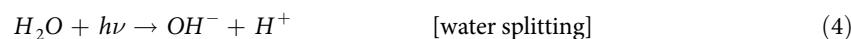
To consider the SPSC mechanism (as illustrated in Fig. 3(a)) via dissociation of water molecules (H<sub>2</sub>O) on nanobumped ZnO surfaces, the electron density and bond-dissociation energy required to alter H<sub>2</sub>O into OH and H radicals were calculated using *ab initio* simulations<sup>28,29</sup>. To clarify the effect of the apical growth of NCs accompanied by the dissociation of water, the bond-dissociation energy was calculated for the flat surface (Fig. 3(b)) and nanobumped surface (Fig. 3(c) for curvature radius,  $R = 0.5$  nm). The high electron density appeared to be localised near the top surface of a nanobump (Fig. 3(c)). The dissociation energy for each surface was evaluated as the difference in the total energy of the dissociated state with H and OH and that of the equilibrated bonded state of H<sub>2</sub>O. The calculated value for the flat surface was 5.03 eV without considering the photoexcitation effect, corresponding to the experimentally measured direct dissociation energy of a water molecule (5.1 eV)<sup>30</sup>. In contrast, the dissociation energy for the nanobumped surface model was 0.323 eV when  $R = 0.5$  nm (subsequently, 0.409 eV for  $R = 1.0$  nm and 0.552 eV for  $R = 2.0$  nm). Notably, the dissociation energy of a water molecule was lower for the nanobumped surface. Overall, the local electron density and dissociation energy reduction at the top of the NC bumps played a key role in the SPSC process.

We now suggest that the mechanism for photosynthesised NCs in water via the SPSC process (Fig. 3(a)) can, *in principle*, be described by the following photo-induced reactions:

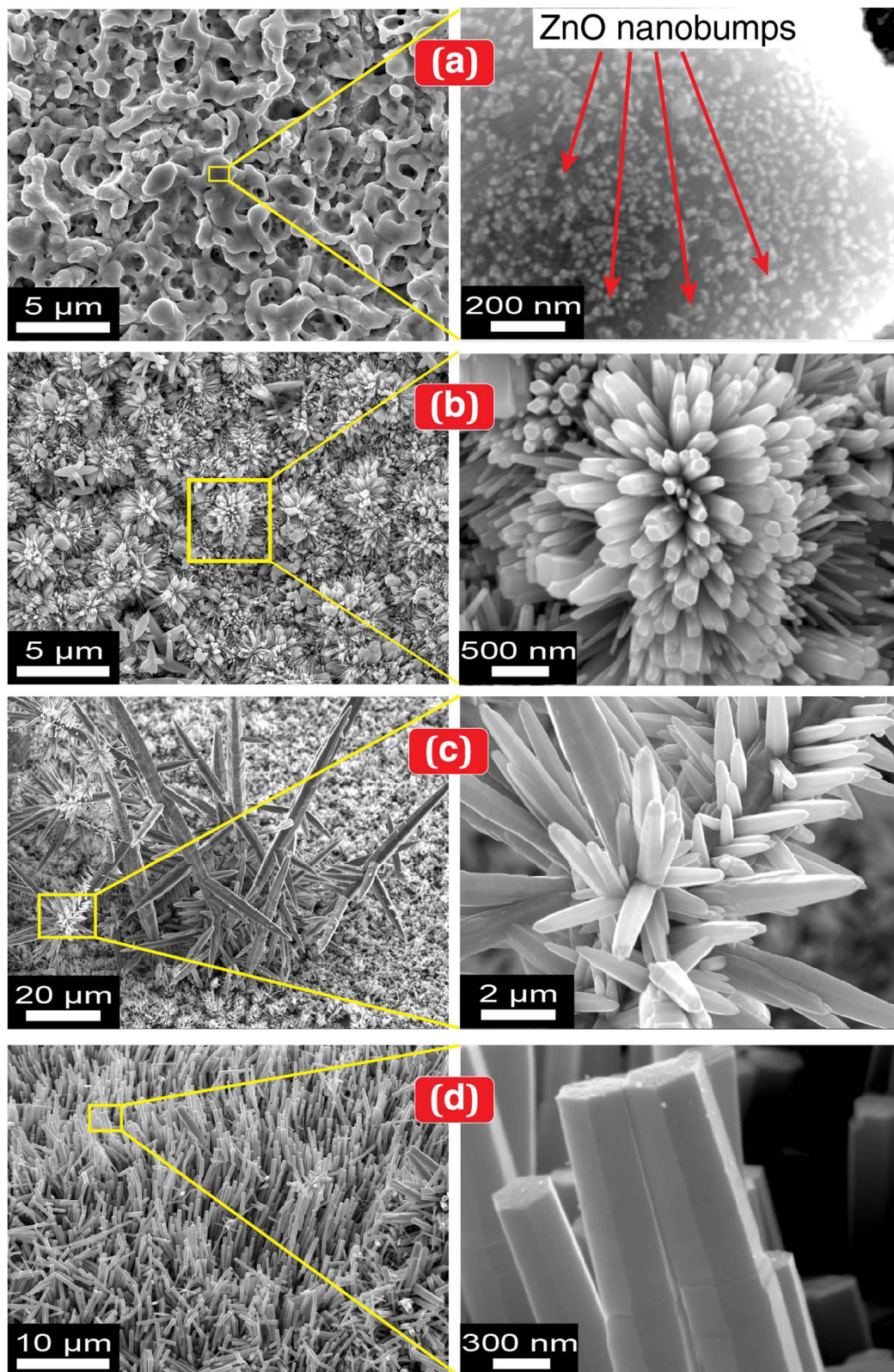
The mechanism starts with water splitting into ion species in the vicinity of a semiconductive (SC) surface:



Hence, the water splits into ions by photoinduction (1)–(3):

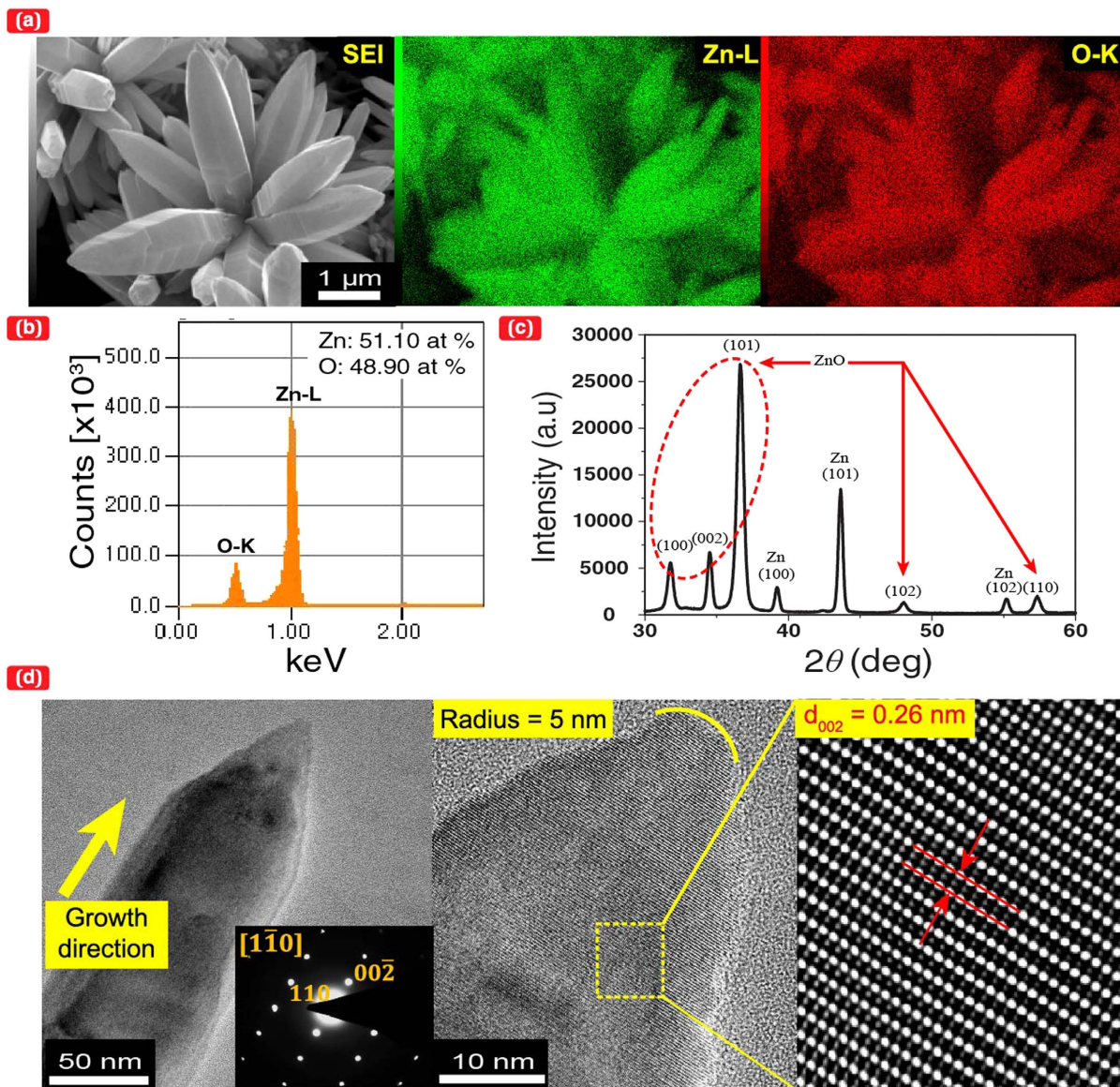


The formation of transient species (H, OH,  $e_{aq}^-$ ) and other molecular byproducts (H<sub>2</sub>, H<sub>2</sub>O<sub>2</sub>) (see Fig. 3(c) for  $H + H \rightarrow H_2$ ) can be well understood as water radiolysis in radiation chemistry<sup>31</sup>. These reactions occur in short times of less than micro-second-order. Presumably, assisted by the



**Figure 1.** Surface morphology after SPSC on a Zn substrate plate. (a) ZnO nanobumps (plasma 140 V, 10 min, UV<sub>0h</sub>). (b) ZnO nanoflowers (UV<sub>24h</sub>). (c) ZnO dendrites (UV<sub>24h</sub>). (d) ZnO nanorods (UV<sub>72h</sub>). The right panel images are the respective magnified FE-SEM micrographs. The heterogeneous growth is due to the local morphology variation via the plasma treatment. Typically, a fine structure of NCs can be obtained at room temperature after 24h of UV irradiation. Extended irradiation increased the size and diameter but terminated the apical growth to yield flat, hexagonal tip ends (at UV<sub>72h</sub> irradiation, as shown in (d) and Supplementary Fig. S2).





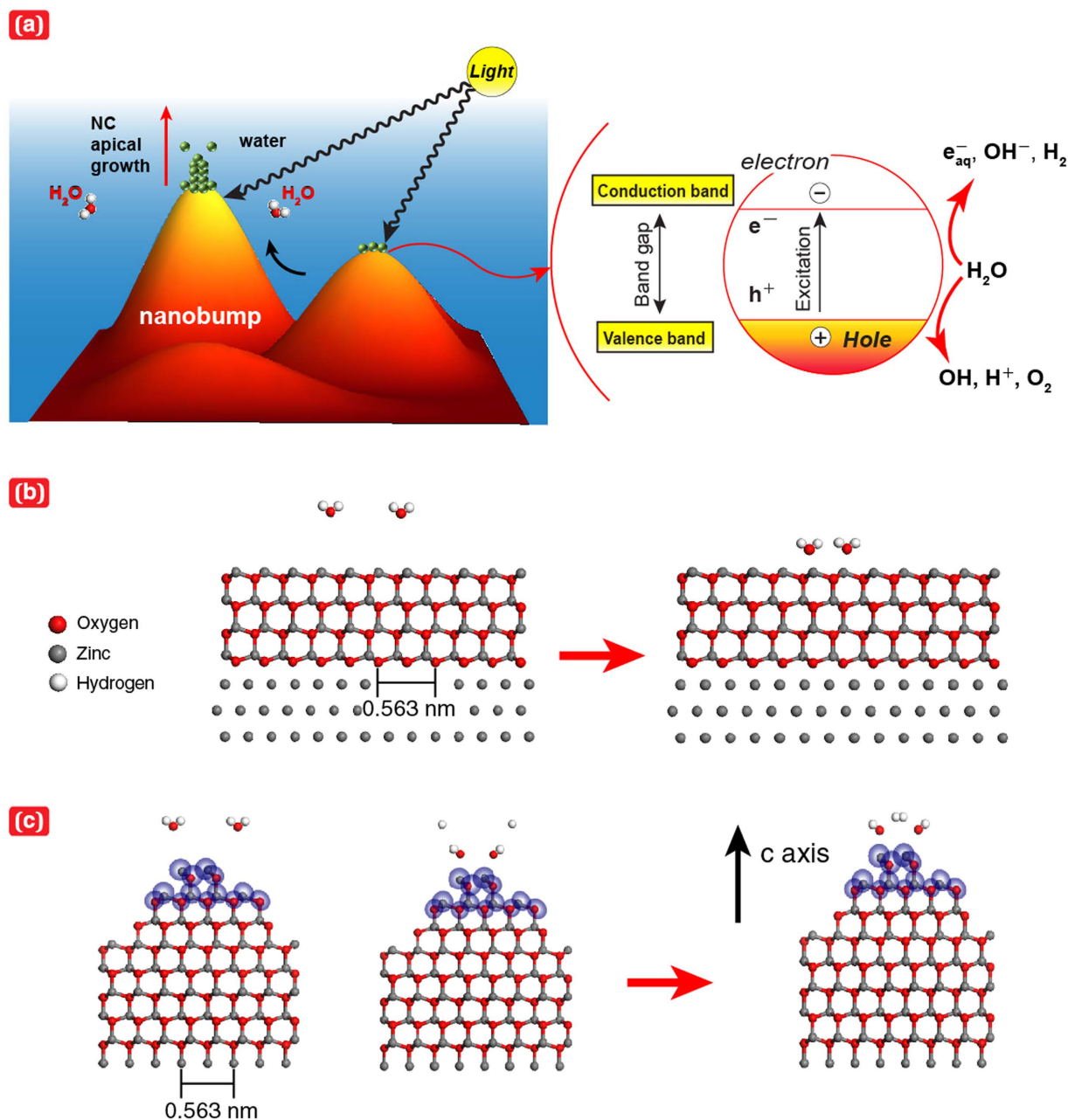
**Figure 2. Structural characterisation of ZnO NCs.** (a) 2-D EDS map of ZnO nanoflowers (UV<sub>48h</sub>); the left panel is a secondary electron image of ZnO crystals grown on the Zn substrate, centre: Zn, and right: O. (b) EDS spectrum and quantitative composition of ZnO. (c) XRD pattern of Zn and ZnO NCs. (d) TEM micrograph of a ZnO nanorod on a carbon thin film; the left panel is the TEM image (200 × 200 nm) of the ZnO nanorod, and the inset is the SAED pattern obtained along the [110] direction; the centre and right panels are the HRTEM image (40 nm × 40 nm) and its magnified image after inverted Fourier transformation, respectively. Fig. 2(d) exhibits the apical growth direction of ZnO in the c-axis.

aforementioned morphology effect,  $H^+$  and  $OH^-$  ions are then locally separated, e.g.,  $H^+$  at a valley in equation (2) and  $OH^-$  at an apical hill in equation (3) throughout a protruded surface. Otherwise,  $H_2O$  will be immediately reproduced in the reverse of equation (4). Such locality-assigned ion production gives rise to a local pH imbalance: alkaline at the hill and acidic near the valley. Therefore, one can expect NC growth at a hill in association with hydrothermal reactions for ZnO generation in an alkaline solution<sup>26,27</sup>:



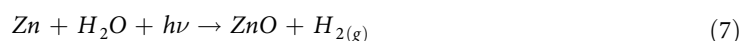
The metal may resolve into an ion at the valley:





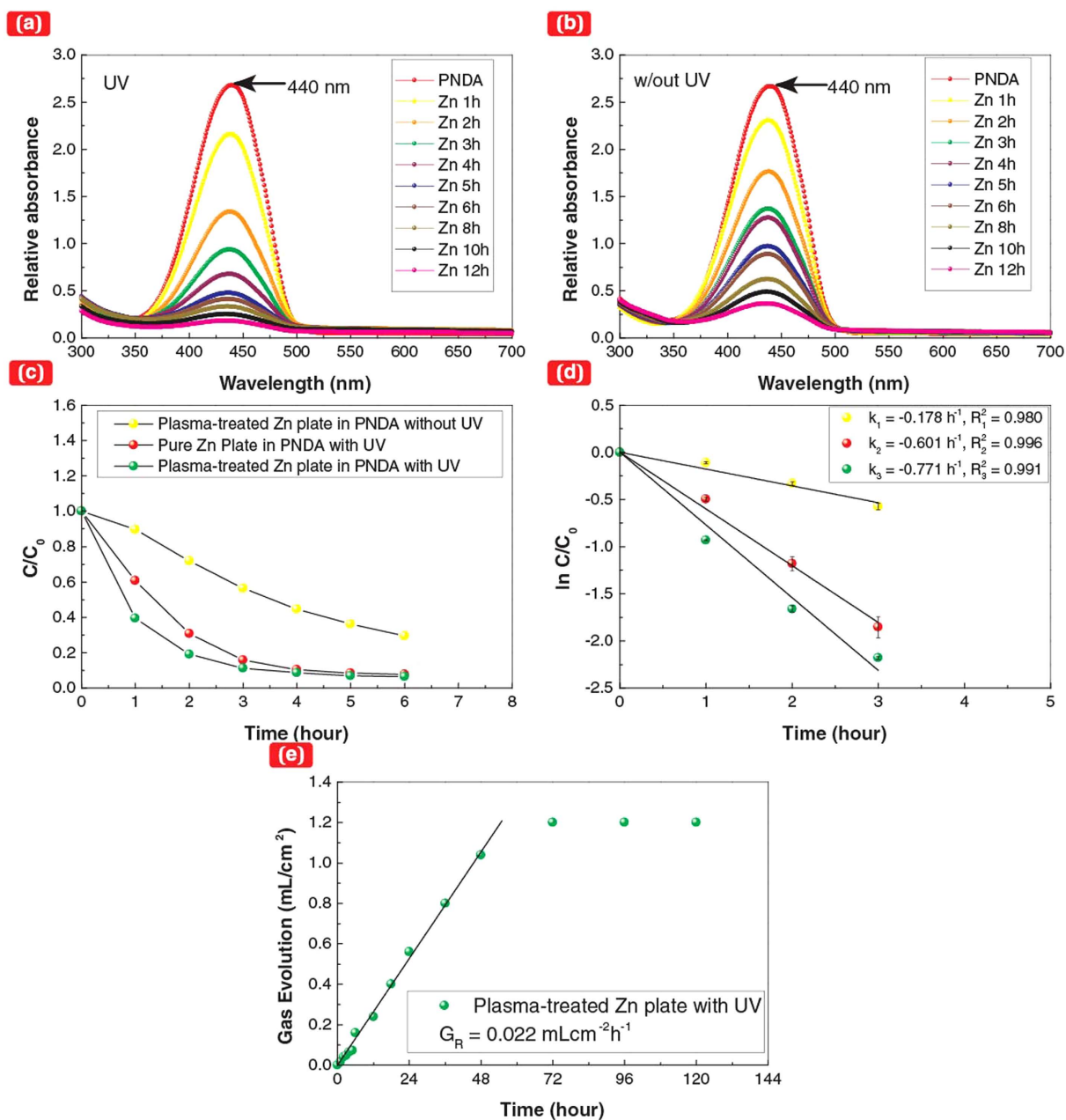
**Figure 3. Nanobumped surface effect in the SPSC model and simulation by molecular dynamical *ab initio* calculations.** (a) A schematic of SPSC with a nanobumped surface. (b) H<sub>2</sub>O molecules with a flat surface of ZnO. Water molecules localise at stable places near the flat surface. (c) Water molecule stabilised on the top of a nanobump (radius of curvature, R = 0.5 nm). Electron density isosurfaces of 2.0 electron/Å<sup>3</sup> (purple coloured) are observed only near the top of a nanobump. The state was relaxed to a state where a H<sub>2</sub> molecule is formed by equilibrating the state with the dissociation of H<sub>2</sub>O molecules into OH and H.

Hence, the net SPSC reaction is completed with



Similarly to known hydrothermal mechanism, NC growth by equations (5)-(6) might be accompanied by aggregation and recrystallisation processes of metal oxide nanoparticles. Thus, the SPSC is completed with three *principles*. Firstly, a photo-induced water splitting process. Secondly, separation of H<sup>+</sup> and OH<sup>-</sup> ions due to nanobumps protruded surface. Finally, aggregation and recrystallisation of metal oxide nanoparticles (superimposed hydrothermal reactions) result in the nanocrystallites growth.

The SPSC process characteristically predicts hydrogen gas (H<sub>2(g)</sub>) and hydroxyl radicals (OH) as intermediate products (see Fig. 4). To confirm our model, we detected H<sub>2</sub> gas using gas chromatography

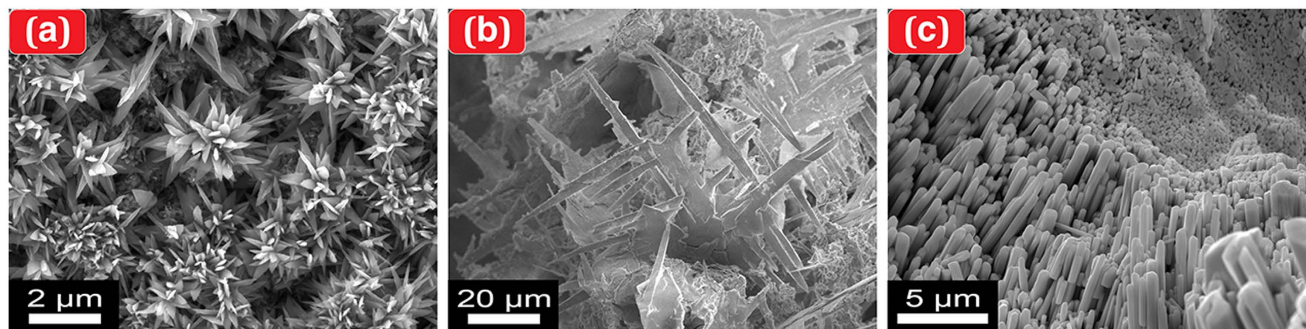


**Figure 4. Detection of OH radicals and measurement of gas evolution.** (a–b) PNDa peak intensity decrease at  $\lambda = 440$  nm to observe the effect of UV irradiation. (a) Plasma-treated Zn plate (UV). (b) Plasma-treated Zn plate (No UV). (c,d) Decomposition trend for PNDa by evaluating the concentration change at the  $\lambda = 440$  nm peak. (e) Time dependence of the gas evolution on extended UV-irradiated, plasma-treated Zn. The gas volume increase slowed down after UV<sub>72h</sub>, which is consistent with the terminated apical growth of a nanorod in Fig. 1(d).

(GC) after the SPSC experiment on Zn. The ratio of H<sub>2</sub>/O<sub>2</sub> in the collected gas was evaluated to be nearly 10. H<sub>2</sub> gas can also be produced by a typical photocatalytic reaction<sup>12</sup> ( $2H + 2e^- \rightarrow H_2$ ), as well as via water radiolysis (with hydrogen radicals, Fig. 3(c)) and hydrothermal reaction (equation (6)). In the present study, OH radical generation was also investigated (Fig. 4(a–d)) by monitoring the bleaching of *p*-nitrosodimethylaniline (PNDa) as the intensity of the characteristic absorption peak at  $\lambda = 440$  nm decreased<sup>32,33</sup>. These results confirm the photochemical reactions proposed for the SPSC process, in which H<sub>2</sub> gas and OH radicals were generated during the photo-assisted growth of ZnO nanorods.

In general, hydrothermal reactions in equations (5) and (6) are known to occur in alkaline solutions at higher temperatures<sup>18,26,27</sup>. Noting this, we tested the SPSC process at two different water temperatures (10 °C and 60 °C) other than room temperature. The NC growth was clearly enhanced at higher water





**Figure 5.** Various nanostructures of metallic oxides produced by the SPSC method. (a) CuO nanoflowers (UV<sub>24h</sub>). (b) CeO<sub>2</sub> dendrites (Vis<sub>24h</sub>,  $\lambda \approx 500$  nm). (c) WO<sub>3</sub> nanorods (UV<sub>48h</sub>).

temperatures. Nonetheless, the overall SPSC process is to occur assumed in a pure, neutral water environment rather than in acidic or alkaline aqueous solutions. However, one can observe the pH dependence or impurity additive influence on the final NC morphology. Moreover, the photocorrosion effect<sup>34</sup>, e.g.,  $ZnO + 2h^+ \rightarrow Zn^{2+} + \frac{1}{2}O_2$ , which causes ZnO to redissolve into Zn ions or  $Zn(OH)_2$  or  $Zn(OH)_4^{2-}$  (equation (5)) and  $Zn(OH)_2^{2-}$  (equation (6)). They may suppress the reaction processes and thus reduce the yield of SPSC in equation (7). However, the re-dissolution effect results in the possibility of ZnO recrystallisation at the initial fine nanocrystallites to regrow into thicker and wider NCs by re-precipitation, as shown in Fig. 1(b–d).

With a slight modification of reactions in equations (5–7), our SPSC method can also be applied to derive a wide variety of metal oxide NCs (e.g., metals of Cu, Ce, and W, as shown in Fig. 5). These modifications revealed the difference in the SPSC morphology, resulting in various NCs. In the present work, we typically employed the submerged liquid plasma technique as an initial surface nanostructure treatment of natively oxidised NC seeds. Nevertheless, other possible alternative seeding methods can also be applied, including laser processing, ion irradiation, and tribological, mechanical scratching.

In summary, SPSC requires *light*, ranging from UV to visible (as shown in Fig. 5(b)), to assist the apical growth of NCs. Secondly, the use of *water* (specifically, ultrapure water) will deliver a fine structure of NCs. Because the NC morphology is sensitive to pH changes and water impurities, the additive effect resulted in different morphologies. For example, Si from tap water resulted in sphere-like crystallites and NaCl from natural seawater resulted in plate-like crystallites. These results are presumably caused by the alteration of the electronic state of the apical surfaces. Finally, metal oxide surface nanobumps act as the NC *seeds* and enclose the nucleation for the apical growth reaction.

## Methods

**Surface pretreatment.** In the submerged liquid plasma experiment devices (Supplementary Fig. S1(a)), the anode was a  $\varnothing 0.5 \times 1$  mm platinum wire (Nilaco, Tokyo, Japan) with purity of 99.9% arched into a hemispherical glass mesh ( $R = 30$  mm). The cathode (target material) was a raw metal Zn plate (Nilaco, Japan, 99.5%), cut into a size of  $35 \times 5 \times 1$  mm. A  $60 \text{ mm}^2$  contact area with a wrapped  $\varnothing 0.5$  mm Cu wire (Nilaco, Japan, 99.9%) on the tip of the Zn plate was used to prepare the working electrode. A solution of 0.1 mol/l  $K_2CO_3$  with pH 11.5 was used as the electrolyte. Deionised water was used as the washing solution. Prior to the experiments, both of the electrodes were washed with deionised water, and the electrolyte was preheated to 90 °C. Insulation of the contact area between the Cu wire and Zn plate was achieved by a  $\varnothing 10$  mm glass tube, ensuring that the exposed Zn plate length was approximately 25 mm. Then, both of the electrodes were immersed in the  $K_2CO_3$  solution (300 ml) and separated by distance of 30 mm. A discharge voltage of 140 V (current: 1.6–1.8 A) was applied across the electrodes using a direct current power supply (KIKUSUI, PWR1600H, Japan). The synthesis of nanobumps using submerged liquid plasma was conducted for a fixed reaction time of 10 min, appreciating the simple and time- and cost-efficient technique. At the end of the plasma reaction, the cathode was collected and washed with deionised water, and the length was cut to 25 mm. A white film surface, confirmed to constitute of ZnO, was obtained on the electrode surface. For further experimentation and analysis, the specimen was allowed to dry at ambient temperature.

**SPSC experiment.** In the UV irradiation experiment (Supplementary Fig. S1(b)), the plasma-treated Zn plate was inserted into a polymethylmethacrylate (PMMA) cuvette, which was then filled with 4 ml of ultrapure water (Wako Pure Chemical, pH 7–7.5, resistivity  $18 \text{ M}\Omega$ ) and capped. Prior experiments, the ultrapure water was degassed to remove the dissolved gas. A UV lamp (UVP, B-100AP, USA) with 100 W longwave UV ( $\lambda = 365$  nm, 3.4 eV) was mainly used for SPSC. Visible light irradiation ( $\lambda \approx 500$  nm) was employed for Fig. 5(b) using spot light source (Hamamatsu LightningCure LC8, L9588, Japan). The irradiance orientation was set to the horizontal position, and the distance between the specimen and

UV lamp was set to 100 mm. In the typical synthesis of nanocrystallites, the UV irradiation was performed in a dark chamber for a fixed reaction time (24 h) at room temperature. Extended UV irradiation times (48 h and 72 h) were also applied to clarify the NC growth characteristics (Fig. 1(d), Fig. 5(c), and Supplementary Fig. S2). At the end of the UV irradiation, the specimen was collected, and the ultrapure water pH change was recorded using a pH meter (Horiba, D-51). For ZnO, the final pH of the water solution exhibited a typical increase to 8.5 in ambient temperature. The final water temperature increase was measured to be less than 10 °C.

**Crystallite characterisation.** The surface morphology and elemental composition analysis of the substrates were monitored using a field emission scanning electron microscopy (FE-SEM, JEOL, JSM-7001FA). The chemical properties analysis was performed using X-ray diffraction (XRD, Rigaku, Tokyo, Japan, RINT2500HLB) with a Cu K $\alpha$  line of 1.5406 Å and a scanning field of  $2.5^\circ \leq 2\theta \leq 100^\circ$ . Peak fitting was performed in referenced to JCPDS card 4-0831 and 5-0664. TEM micrographs, SAED patterns and HRTEM micrographs for the NCs were obtained using a double Cs-corrected-TEM (FEI, Titan cubed) operated at 300 kV.

**Ab initio calculation.** The simulation models of a flat surface and a nanobumped surface were constructed using Materials Studio<sup>®</sup> atomic simulation software (Accelrys Software Inc.). The calculations were performed based on density functional theory (DFT)<sup>35,36</sup>. The radii of curvature of the nanobumped surface, shown in Fig. 3(c) and Supplementary Fig. S3(a) and S3(b), were set to 0.5, 1.0, and 2.0 nm, respectively. The grey, red, and white spheres in these figures represent zinc, oxygen, and hydrogen atoms, respectively. Initially, two free H<sub>2</sub>O molecules were placed in positions where strong interatomic forces were not exerted on each atom. The position of each atom in the equilibrated state was obtained using a dynamic simulated annealing method<sup>28,29</sup>. In this method, the electronic states can be calculated by solving the quantum mechanical equation. In the actual experiment for the effect of nanobumps, the reduced dissociation energy of the water molecules can be larger than 0.323 eV because the radius of curvature of the nanobumped (or apical) surfaces might be larger than 0.5 nm (e.g., approximately, 5 nm in Fig. 2(d)). However, that value was considered to be considerably smaller than 5.03 eV for the flat surface because the apical radius of the curved surface was still extremely small (typically approximately 10 nm). Additionally, an illumination, e.g., a typical UV light ( $\lambda = 350$  nm, 3.54 eV), further enhanced the dissociation of water molecules on the nanobumped surface because the photon energy was considerably larger than the dissociation energy for the nanobumped surface (0.323 eV). This energy difference resulted in the localised SPSC reaction increment via a photo-electron excitation effect, namely, the enhancement of equation (1) for the generation of additional excited electrons. Therefore, for the dissociated state of water molecules, which was induced by illumination, followed by an equilibration process, we obtained a final state, as shown in Fig. 3(c), in which H<sub>2</sub> molecule (gaseous) formation was predicted.

**OH radical analysis.** During the UV light irradiation of the plasma-treated Zn plate, 4 ml of PNDA with a concentration of 1.5 mg/l was used as a scavenger in OH radical detection. Based from estimated O<sub>2</sub> production in Fig. 4(e), the greater factor of ~24 from the experiment results can neglect the dissolved gas effect in generated radicals: photo-induced water splitting governed the OH radical production. The absorption spectra before and after UV irradiation were compared: the concentration of PNDA was measured using a JASCO V-630 UV-Vis spectrophotometer. Then, the time vs. exponential decay was plotted, and the first-order reaction rate (k) was calculated.

**H<sub>2</sub> gas analysis.** The gas captured after UV<sub>120h</sub> of six plasma-treated Zn substrates was used for GC analysis. The analysis of H<sub>2</sub> and O<sub>2</sub> gases was performed using a Shimadzu GC 8-A (thermal conductivity detector, molecular sieve 13X, N<sub>2</sub> carrier for H<sub>2</sub>, and He carrier for O<sub>2</sub>). For H<sub>2</sub> gas detection, 100% H<sub>2</sub> gas (100  $\mu$ l) was injected into the GC using a microsyringe, and the calibration curve was plotted. Then, 100  $\mu$ l of the captured gas was injected into the GC, and its concentration was compared with the calibration curve. The same steps were repeated when recording the O<sub>2</sub> concentration. The obtained result gives in the H<sub>2</sub>/O<sub>2</sub> ratio of 10. The value is five times more than two in the case of normal water splitting.

## References

- Huang, M. H. *et al.* Room-temperature ultraviolet nanowire nanolasers. *Science* **292**, 1897–1899 (2001).
- Wang, X., Zhuang, J., Peng, Q. & Li, Y. A general strategy for nanocrystal synthesis. *Nature* **437**, 121–124 (2005).
- Morales, A. M. & Lieber, C. M. A laser ablation method for the synthesis of crystalline semiconductor nanowires. *Science* **279**, 208–211 (1998).
- Kubota, Y., Watanabe, K., Tsuda, O. & Taniguchi, T. Deep ultraviolet light-emitting hexagonal boron nitride synthesized at atmospheric pressure. *Science* **317**, 932–934 (2007).
- Pan, Z. W., Dai, Z. R. & Wang, Z. L. Nanobelts of semiconducting oxides. *Science* **291**, 1947–1949 (2001).
- Huo, Q. *et al.* Generalized synthesis of periodic surfactant/inorganic composite materials. *Nature* **368**, 317–321 (1994).
- Salunkhe, R. R. *et al.* Large-scale synthesis of coaxial carbon nanotube/Ni(OH)<sub>2</sub> composites for asymmetric supercapacitor application. *Nano Energy* **11**, 211–218 (2015).
- Lin, J. *et al.* Mesoporous anatase single crystals for efficient Co<sup>(2+/3+)</sup>-based dye-sensitized solar cells. *Nano Energy* **11**, 557–567 (2015).
- Briseno, A. L. *et al.* Patterning organic single-crystal transistor arrays. *Nature* **444**, 913–917 (2006).
- Diao, Y. *et al.* Controlled nucleation from solution using polymer microgels. *J. Am. Chem. Soc.* **133**, 3756–3759 (2011).



11. Diao, Y., Myerson, A. S., Hatton, T. A. & Trout, B. L. Surface design for controlled crystallization: The role of surface chemistry and nanoscale pores in heterogeneous nucleation. *Langmuir* **27**, 5324–5334 (2011).
12. Fujishima, A. & Honda, K. Electrochemical photolysis of water at a semiconductor electrode. *Nature* **238**, 37–38 (1972).
13. Saito, N. *et al.* Low-temperature fabrication of light-emitting zinc oxide micropatterns using self-assembled monolayers. *Adv. Mater.* **14**, 418–421 (2002).
14. Tian, W. *et al.* Low-cost fully transparent ultraviolet photodetectors based on electrospun ZnO-SnO<sub>2</sub> heterojunction nanofibers. *Adv. Mater.* **25**, 4625–4630 (2013).
15. Golego, N., Studenikin, S. A. & Cocivera, M. Sensor photoresponse of thin-film oxides of zinc and titanium to oxygen gas. *J. Electrochem. Soc.* **147**, 1592–1594 (2000).
16. Keis, K., Magnusson, E., Lindström, H., Lindquist, S. E. & Hagfeldt, A. A 5% efficient photoelectrochemical solar cell based on nanostructured ZnO electrodes. *Sol. Energy Mater. Sol. Cells* **73**, 51–58 (2002).
17. Govender, K., Boyle, D. S., Kenway, P. B. & O'Brien, P. Understanding the factors that govern the deposition and morphology of thin films of ZnO from aqueous solution. *J. Mater. Chem.* **14**, 2575–2591 (2004).
18. Lincot, D. Solution growth of functional zinc oxide films and nanostructures. *MRS Bull.* **35**, 778–789 (2010).
19. Tian, Z. R. *et al.* Complex and oriented ZnO nanostructures. *Nat. Mater.* **2**, 821–826 (2003).
20. Toriyabe, Y., Watanabe, S., Yatsu, S., Shibayama, T. & Mizuno, T. Controlled formation of metallic nanoballs during plasma electrolysis. *Appl. Phys. Lett.* **91**, 041501–041503 (2007).
21. Julaihi, M. R. M. b., Yatsu, S., Jeem, M. & Watanabe, S. Synthesis of stainless steel nanoballs via submerged glow-discharge plasma and its photocatalytic performance in methylene blue decomposition. *J. Exp. Nanosci.* **10**, 965–982, (2014).
22. Sōmiya, S. & Roy, R. Hydrothermal synthesis of fine oxide powders. *Bull. Mater. Sci.* **23**, 453–460 (2000).
23. Lee, M. J. *et al.* A fast, high-endurance and scalable non-volatile memory device made from asymmetric Ta<sub>2</sub>O<sub>5-x</sub>/TaO<sub>2-x</sub> bilayer structures. *Nat. Mater.* **10**, 625–630 (2011).
24. Shi, L., Bao, K., Cao, J. & Qian, Y. Sunlight-assisted fabrication of a hierarchical ZnO nanorod array structure. *CrysEngComm* **11**, 2009–2014 (2009).
25. Wu, J. M. & Chen, Y. R. Ultraviolet-light-assisted formation of ZnO nanowires in ambient air: Comparison of photoresponsive and photocatalytic activities in zinc hydroxide. *J. Phys. Chem. C* **115**, 2235–2243 (2011).
26. Jang, J. H., Park, J. H. & Oh, S. G. Effects of dodecyl sulfate anionic surfactants on the crystal growth of ZnO through hydrothermal process. *J. Ceram. Process. Res.* **10**, 783–790 (2009).
27. Shi, R. *et al.* Growth of flower-like ZnO via surfactant-free hydrothermal synthesis on ITO substrate at low temperature. *CrysEngComm* **14**, 5996–6003 (2012).
28. Car, R. & Parrinello, M. Unified approach for molecular dynamics and density-functional theory. *Phys. Rev. Lett.* **55**, 2471–2474 (1985).
29. Segall, M. D. *et al.* First-principles simulation: Ideas, illustrations and the CASTEP code. *J. Phys.: Condens. Matter* **14**, 2717 (2002).
30. Petrik, N. G., Alexandrov, A. B. & Vall, A. I. Interfacial energy transfer during gamma radiolysis of water on the surface of ZrO<sub>2</sub> and some other oxides. *J. Phys. Chem. B* **105**, 5935–5944 (2001).
31. Le Caër, S. Water radiolysis: Influence of oxide surfaces on H<sub>2</sub> production under ionizing radiation. *Water* **3**, 235–253 (2011).
32. Baxendale, J. H. & Khan, A. A. The pulse radiolysis of p-nitrosodimethylaniline in aqueous solution. *Int. J. Radiat. Phys. Ch.* **1**, 11–24 (1969).
33. Simonsen, M. E., Muff, J., Bennedsen, L. R., Kowalski, K. P. & Søgaard, E. G. Photocatalytic bleaching of p-nitrosodimethylaniline and a comparison to the performance of other AOP technologies. *J. Photochem. Photobiol. A: Chem.* **216**, 244–249 (2010).
34. Domenech, J. & Prieto, A. Stability of zinc oxide particles in aqueous suspensions under uv illumination. *J. Phys. Chem.* **90**, 1123–1126 (1986).
35. Kohn, W. & Sham, L. J. Self-consistent equations including exchange and correlation effects. *Phys. Rev.* **140**, A1133–A1138 (1965).
36. Hohenberg, P. & Kohn, W. Inhomogeneous electron gas. *Phys. Rev.* **136**, B864–B871 (1964).

## Acknowledgements

This work was supported by the “Nanotechnology Platform” from The Ministry of Education, Culture, Sports, Science and Technology (MEXT). The authors are grateful to Professor N. Sakaguchi, Mr. K. Ohkubo, and R. Ohta for their helpful discussions and technical assistance in operating the TEM system. We also thank Mr. Yukiyasu Yamakoshi for his technical assistance in operating the gas chromatography system.

## Author Contributions

M.J. and M.R.M.J. performed most of the fabrication experiments and microstructural analyses. M.R.M.J., S.Y. and K.O. conducted the OH radical and gas analyses. T.K. performed the XRD analysis. M.J., J.I. and T.S. performed the TEM observation. T.I. and T.K. conducted the *ab initio* calculation. S.W. conceived and supervised the project and developed the model. M.J., J.I. and S.W. wrote the paper. All of the authors discussed the results and contributed to the manuscript.

## Additional Information

**Supplementary information** accompanies this paper at <http://www.nature.com/srep>

**Competing financial interests:** The authors declare no competing financial interests.

**How to cite this article:** Jeem, M. *et al.* A pathway of nanocrystallite fabrication by photo-assisted growth in pure water. *Sci. Rep.* **5**, 11429; doi: 10.1038/srep11429 (2015).



This work is licensed under a Creative Commons Attribution 4.0 International License. The images or other third party material in this article are included in the article's Creative Commons license, unless indicated otherwise in the credit line; if the material is not included under the Creative Commons license, users will need to obtain permission from the license holder to reproduce the material. To view a copy of this license, visit <http://creativecommons.org/licenses/by/4.0/>