

SCIENTIFIC REPORTS

OPEN

One-Pot Solvothermal Synthesis of $\text{Bi}_4\text{V}_2\text{O}_{11}$ as A New Solar Water Oxidation Photocatalyst

Received: 10 December 2015

Accepted: 17 February 2016

Published: 07 March 2016

Zaiyong Jiang¹, Yuanyuan Liu¹, Mengmeng Li², Tao Jing², Baibiao Huang¹, Xiaoyang Zhang¹, Xiaoyan Qin¹ & Ying Dai²

$\text{Bi}_4\text{V}_2\text{O}_{11}$ was prepared via a one-pot solvothermal method and characterized via XRD, Raman, XPS, Electrochemical impedance spectroscopy. The as-prepared $\text{Bi}_4\text{V}_2\text{O}_{11}$ sample displays excellent photocatalytic activity towards oxygen evolution under light irradiation. The hierarchical structure is in favour of the spatial separation of photogenerated electrons and holes. Furthermore, the internal polar field also plays a role in improving the charge separation. Both of the two results are responsible for excellent activity of O_2 evolution. The resulting hierarchical $\text{Bi}_4\text{V}_2\text{O}_{11}$ sample should be very promising photocatalyst for the application of photocatalytic O_2 evolution in the future.

Since the first report about photocatalytic water splitting via TiO_2 electrode by Honda *et al.* in 1972¹, how to convert solar energy into renewable clean chemical energy has become one of the most profound challenges^{2,3}. Over the past few years, various effort has been put into the research on water splitting via semiconductor photocatalysis⁴⁻⁷, and some excellent results have been achieved^{8,9}. Nevertheless, the efficiencies are still low and do not meet the requirements of practical applications¹⁰. As far as we know, the photocatalytic oxygen evolution is considered as the crucial step for the water splitting¹¹. Therefore, the major challenge for water splitting is how to work out the problem of O_2 evolution.

Currently, BiVO_4 is the most widely investigated photocatalyst for the water splitting to produce O_2 , due to its advantage, such as nontoxicity, low cost and high chemical and thermal stability etc. Various methods were proposed to enhance the photocatalytic activity of BiVO_4 , including doping¹², semiconductor recombination^{13,14}, and cocatalyst depositing¹¹, etc. Based on the above discussion, we can find that the current studies on O_2 evolution mainly focused on the modification of BiVO_4 . However, the reports about exploration of new photocatalyst with the performance of oxygen evolution are limited. Therefore, further explorations on new photocatalysts with photocatalytic oxygen evolution activity are of great importance for both scientific research and practical applications.

$\text{Bi}_4\text{V}_2\text{O}_{11}$ is one of newly exploited Bi-based semiconductor photocatalyst with the features of nontoxicity, low cost and high chemical and thermal stability. It has a layered structure that consists of $[\text{Bi}_2\text{O}_2]$ slabs which are interleaved by $(\text{VO}_{3.5}\square_{0.5})^{2-}$ (Fig. 1a)^{15,16}. The unique layered structure is in favour of enhancing the separation of photo-generated electron-hole pairs¹⁷, which can improve the photocatalytic property. Moreover, this compound exhibits strong polar response^{18,19}, which can form the internal polar electric field to further enhance the separation of photoinduced carriers^{20,21}. So, $\text{Bi}_4\text{V}_2\text{O}_{11}$ photocatalyst has attracted widespread attention for applications in the degradation of organic compounds under solar light illumination^{22,23}. For example, Liu *et al.*¹⁵ prepared $\text{Bi}_4\text{V}_2\text{O}_{11}$ hierarchical hollow microspheres by a facile template-free solvothermal route and exhibited excellent photocatalytic activity for the degradation of Rhodamine-B.

However, to the best of our knowledge, researches on the photocatalytic activity of $\text{Bi}_4\text{V}_2\text{O}_{11}$ are limited to organic molecules degradation. Up to now, there is no a report whether the $\text{Bi}_4\text{V}_2\text{O}_{11}$ photocatalyst can split water to generate oxygen. Through the preliminary observation, we could find that the $\text{Bi}_4\text{V}_2\text{O}_{11}$ possess same element compositions with the BiVO_4 , which suggests their properties to be possibly similar. In order to further confirm the possibility, we have calculated density of states of $\text{Bi}_4\text{V}_2\text{O}_{11}$. As shown in Fig. 1b, the VBM of $\text{Bi}_4\text{V}_2\text{O}_{11}$ are mainly contributed by O 2p states, and slightly contributed by Bi 6s states, moreover, the CBM are mainly composed of V 3d states and Bi 6p states, which is basically consistent with that of BiVO_4 (Figure S1). Based on above discussion, we speculate that $\text{Bi}_4\text{V}_2\text{O}_{11}$ might have excellent performance of O_2 evolution like BiVO_4 . Besides that,

¹State Key Laboratory of Crystal Materials, Shandong University, Jinan 250100, P. R. China. ²School of Physics, Shandong University, Jinan 250100, P. R. China. Correspondence and requests for materials should be addressed to Y.L. (email: yyliu@sdu.edu.cn) or B.H. (email: bbhuang@sdu.edu.cn)

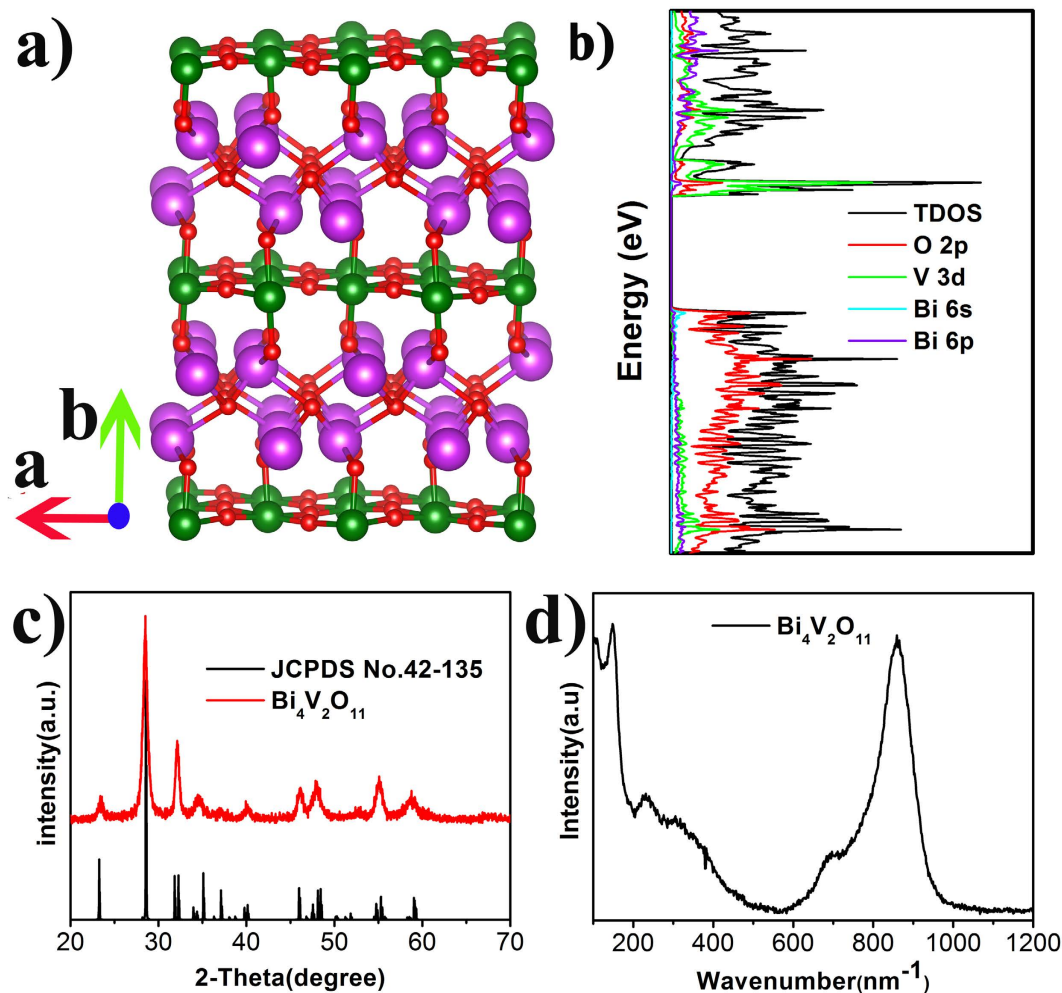


Figure 1. A perspective view of the $\text{Bi}_4\text{V}_2\text{O}_{11}$ slab (a) (The big purple circles represent Bi atoms, the small red circles represent O atoms and the green circles represent V atoms, respectively.), Density of states of $\text{Bi}_4\text{V}_2\text{O}_{11}$ (b) XRD patterns of $\text{Bi}_4\text{V}_2\text{O}_{11}$ (c) and Raman spectra for $\text{Bi}_4\text{V}_2\text{O}_{11}$ (d).

building internal polar electric field was already believed to be a useful way to improve photocatalytic activity. Based on above discussion, the exploration of O_2 evolution of $\text{Bi}_4\text{V}_2\text{O}_{11}$ is of great importance for both scientific research and practical applications.

In this paper, we synthesized the $\text{Bi}_4\text{V}_2\text{O}_{11}$ photocatalyst via a one-pot facile solvothermal method and efficient photocatalytic oxygen evolution from water was observed over $\text{Bi}_4\text{V}_2\text{O}_{11}$. First principle calculation suggests that both the layered structure and the spontaneous electric polarization of $\text{Bi}_4\text{V}_2\text{O}_{11}$ play important roles in suppressing the recombination of photogenerated charge carriers, which enable $\text{Bi}_4\text{V}_2\text{O}_{11}$ to display efficient photocatalytic activity towards oxygen evolution from water.

Results and Discussion

The XRD pattern of as-prepared $\text{Bi}_4\text{V}_2\text{O}_{11}$ is shown in Fig. 1c, in which all the diffraction peaks can be perfectly indexed to the standard data for orthorhombic crystal structure of $\text{Bi}_4\text{V}_2\text{O}_{11}$ (JCPDS No. 42-135). No impurity peaks were observed, indicating the high purity of $\text{Bi}_4\text{V}_2\text{O}_{11}$. Raman spectra were further investigated so as to obtain more structural information of $\text{Bi}_4\text{V}_2\text{O}_{11}$. The result was shown in Fig. 1d. A broad peak between 600 and 950 cm^{-1} was observed, which could be attributed to stretching modes of V-O band of VO_6 groups. Moreover, two peaks located at about 232 and 303 cm^{-1} were found, which can be regarded as lattice modes or vibration chain bending, respectively²⁴. This result also suggests the successful preparation of $\text{Bi}_4\text{V}_2\text{O}_{11}$, which is in good consistent with the foregoing XRD result. SEM images (Figure S2) suggest that the morphology of the as-prepared $\text{Bi}_4\text{V}_2\text{O}_{11}$ is irregular nanoparticle. And EDS results (Figure S3) suggest that the atom ratio of Bi to V is 2.14, which is in accordance with the stoichiometric value, i.e. 2.

X-ray photoelectron spectra (XPS) were further investigated to identify the surface compositions and valence states of $\text{Bi}_4\text{V}_2\text{O}_{11}$, which is shown in Figure S4. Two peaks at 159.20 eV and 164.48 eV are detected, which can be attributed to $\text{Bi}^{3+} 4f_{7/2}$ and $\text{Bi}^{3+} 4f_{5/2}$, respectively²⁵. There are three binding energy locates at 529.88, 531.25 and 532.71 eV, respectively, which are assigned to the lattice oxygen, hydroxyl groups adsorbed onto the surface of sample and oxygen vacancies. Furthermore, the binding energy peaks at 516.72 and 524.09 eV are ascribed to the

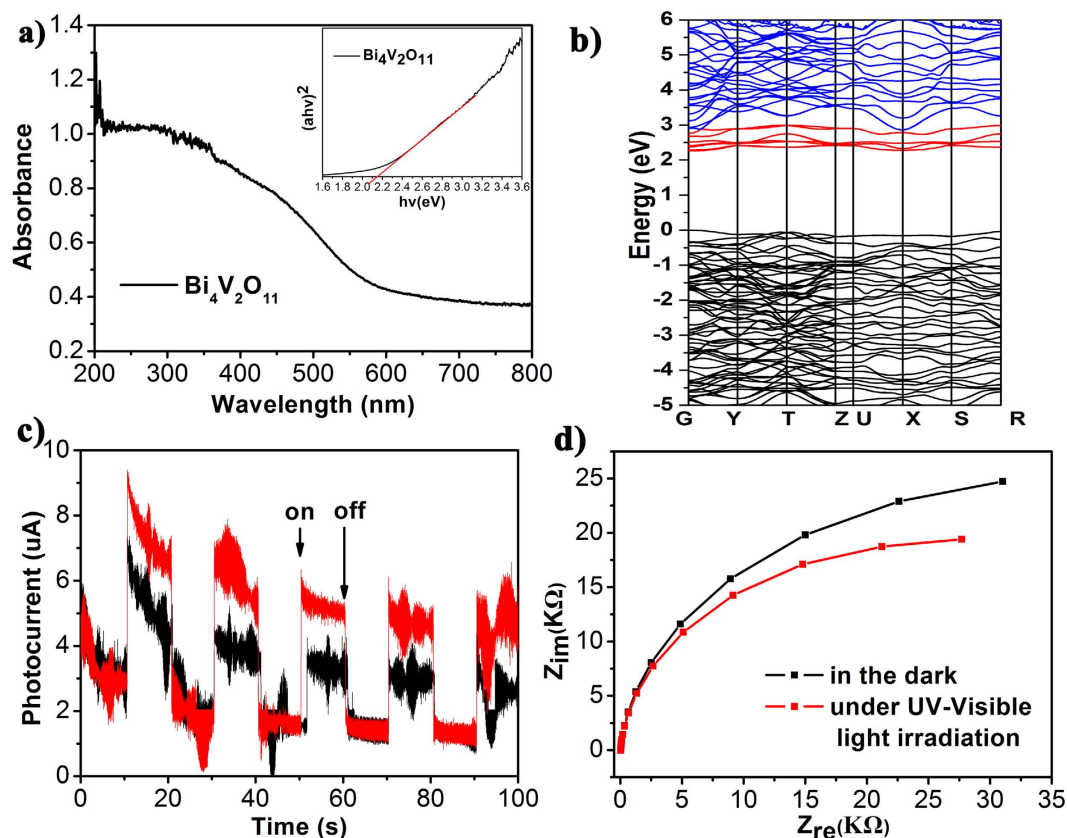


Figure 2. (a) UV-Visible DRS spectra, (inset of a) plots of $(ah\nu)^2$ versus the photon energy ($h\nu$), (b) the images of energy band structure of $\text{Bi}_4\text{V}_2\text{O}_{11}$, (c) Transient photocurrent responses of $\text{Bi}_4\text{V}_2\text{O}_{11}$ under a bias of 0.4V and 0.8 V and (d) EIS Nyquist plots the $\text{Bi}_4\text{V}_2\text{O}_{11}$ electrode in the dark and under UV-Visible light irradiation. The reference electrode was Ag/AgCl, and the electrolyte was 0.2 M Na_2SO_4 .

$\text{V } 2p_{3/2}$ and $\text{V } 2p_{1/2}$, respectively, which is identical to the results of V^{5+} in $\text{Bi}_4\text{V}_2\text{O}_{11}$ reported in literature²². The XPS results indicate that $\text{Bi}_4\text{V}_2\text{O}_{11}$ with high purity was successfully synthesized.

Figure 2a shows the UV-Visible (UV-Vis) diffuse reflection spectra (DRS) spectra of $\text{Bi}_4\text{V}_2\text{O}_{11}$. It is observed that $\text{Bi}_4\text{V}_2\text{O}_{11}$ exhibits strong visible-light absorption with the absorption edge of ca. 580nm. The band gap of $\text{Bi}_4\text{V}_2\text{O}_{11}$ is determined to be 2.15eV (the inset of Fig. 2a). In addition, the potentials of conduction band (CB) and valence band (VB) edges of $\text{Bi}_4\text{V}_2\text{O}_{11}$ were calculated using the Mulliken electronegativity theory²⁶, whose equations are listed in equation 2–5. The calculated values were $E_{\text{CB}} = 0.61 \text{ eV}$ and $E_{\text{VB}} = 2.76 \text{ eV}$, respectively. This result suggests that $\text{Bi}_4\text{V}_2\text{O}_{11}$ is able to oxidize water giving rise to oxygen thermodynamically. The energy band structure of $\text{Bi}_4\text{V}_2\text{O}_{11}$ were also calculated and the result was shown in Fig. 2b. It is observed that both the VBM and the CBM locate at Gama point, which indicates that $\text{Bi}_4\text{V}_2\text{O}_{11}$ is a direct band gap semiconductor. The calculated band gap is about 2.26eV, which is good agreement with the experimental value.

Transient photocurrent responses were performed to investigate the photoelectric properties of $\text{Bi}_4\text{V}_2\text{O}_{11}$ and the result is shown in Fig. 2c. The dark current is very low while the photocurrent increases sharply upon light illumination, and reaches a steady state quickly. As expected, the current returns quickly to its darkcurrent state when the light is turned off. This result suggests that free electron-hole pairs can be generated and efficiently separated when $\text{Bi}_4\text{V}_2\text{O}_{11}$ is irradiated by UV-Vis light. The result of electrochemical impedance spectroscopy (EIS) further supports the above conclusion (Fig. 2d). The arc radius in the EIS Nyquist plot under UV-Vis light illumination is smaller than that in the dark, suggesting that the photo-generated electron-hole pairs can be well separated and efficiently transferred to the surface of $\text{Bi}_4\text{V}_2\text{O}_{11}$ ²⁷. Based on the above results, it is reasonable to conclude that $\text{Bi}_4\text{V}_2\text{O}_{11}$ is a potential photocatalyst.

To check above assumption, photocatalytic oxygen evolution reaction was carried out over $\text{Bi}_4\text{V}_2\text{O}_{11}$ in 100 mL of aqueous solution containing AgNO_3 as a sacrificial reagent. As shown in Fig. 3a, $\text{Bi}_4\text{V}_2\text{O}_{11}$ exhibits an excellent O_2 evolution activity under both UV-Vis and Vis light irradiation. Figure 3b shows the photocatalytic O_2 evolution, as a function of reaction time under UV-Vis light irradiation. Obviously, the amount of O_2 evolution increases with increasing the reaction time.

The electric dipole moments were calculated in order to check whether the internal polar electric field plays an important role in the excellent photocatalytic activity of $\text{Bi}_4\text{V}_2\text{O}_{11}$. The bond valences of $[\text{BiO}_4]$ and $[\text{VO}_6]$ unit were calculated using the following equation:

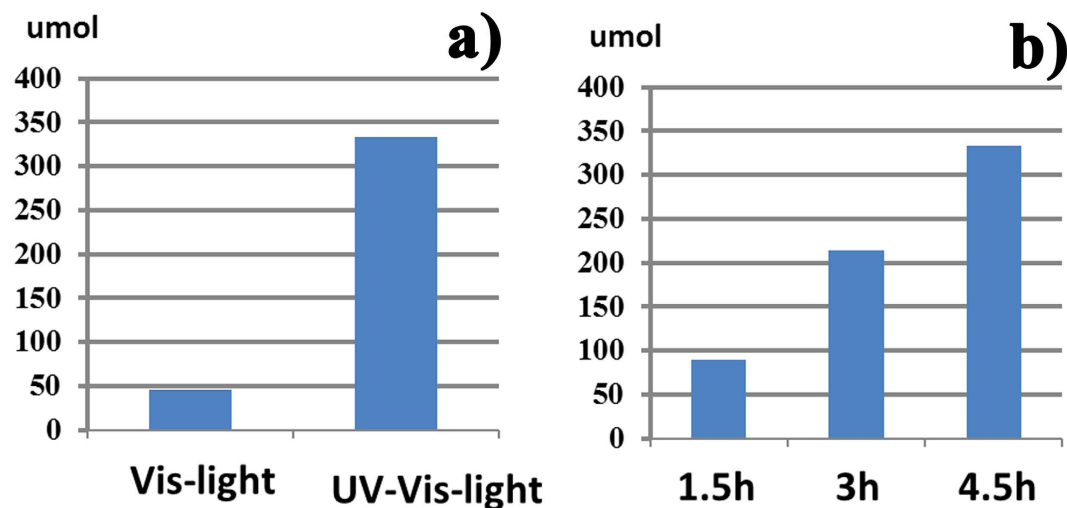


Figure 3. (a) Photocatalytic O₂ evolution of Bi₄V₂O₁₁ and under UV-Visible and Visible light irradiation, (b) the detailed image of O₂ evolution of Bi₄V₂O₁₁ upon UV-Visible light irradiation.

	μ_x	μ_y	μ_z
[VO ₆]	0	0	-7.102
[BiO ₄]-1	4.930	16.234	14.930
[BiO ₄]-2	4.930	-16.234	14.930
[BiO ₄]-3	-4.930	16.234	14.9303
[BiO ₄]-4	-4.930	-16.234	14.9303
Net dipole moments	0	0	91.032

Table 1. The total value of dipole moments of VO₆ and BiO₄ groups in unit cell of Bi₄V₂O₁₁.

$$V_i = \sum_j S_{ij} = \sum_j \exp\left[\frac{R_0 - R_{ij}}{B}\right] \quad (1)$$

In equation, R_0 represents the average bond length (Bi-O or V-O) and $B = 0.37$ is a constant. R_{ij} is the actual bond length between i and j . Moreover, S_{ij} is defined as the valence of the bond i - j . And, V_i is the bond valence sum of the cation i . Through the Debye equation $\mu = neR$, the net dipole moments of [BiO₄] and [VO₆] units were calculated and given in Table 1. Here, n represents the total number of electrons, e is the electron charge, R is the difference between the “centroids” of positive and negative charge and μ is the net dipole moment in Debye. As shown in table 1, it is observed that the dipole moments in ab plane offset mutually. Therefore, the direction of net dipole moment of the Bi₄V₂O₁₁ in a unit cell is along the c axis (0, 0, 1) and the value is determined to be 91.032 D by calculation. From the DOS image (Fig. 1b), it is observed that V atoms makes the mainly contribution to the conduction band. So, the photo-generated electrons will transfer along V-O-V layer direction. While, the holes can transfer along Bi-O-Bi layer direction due to their significant contribution to the valence band. It is noteworthy to point out that the separated direction of photo-generated electron and hole pairs is in accordance with that of internal polar electric field. Therefore, the internal polar field would facilitate the transfer of photogenerated electron and hole pairs along opposite directions, which is advantageous of the separation of photogenerated charge carriers and consequently in favour of high photocatalytic O₂ evolution. Time-resolved PL spectrum of Bi₄V₂O₁₁ (Figure S5) indicates that the excited state of Bi₄V₂O₁₁ displays a two exponential decay. i.e. 0.205 ns (82.46%), and 1.085 ns (17.54%), respectively. The longer lifetime (1.085 ns) is mostly likely due to the effect of the internal polar field.

Based on both the theoretical and experimental results, a possible photocatalytic mechanism is proposed to explain the process of photocatalytic O₂ evolution over Bi₄V₂O₁₁, which is shown in Fig. 4. Under light irradiation, Bi₄V₂O₁₁ semiconductor was excited. The photo-generated electrons and holes gather in the (VO_{3.5}□_{0.5})²⁻ layers and (Bi₂O₂)²⁺ layers, respectively, resulting in the photo-generated electrons and holes spatially separated. Moreover, the internal polar field accelerates the separation process. The photo-generated electrons on (VO_{3.5}□_{0.5})²⁻ layers can be captured by Ag⁺ in the AgNO₃ aqueous solution, generating metallic Ag. At the same time, the photo-generated holes on (Bi₂O₂)²⁺ layers could directly oxidize water to produce O₂. In our opinion, both Bi₄V₂O₁₁ and BiVO₄ are bismuth vanadium based semiconductors, and both of them are stable during the photocatalytic process. Moreover, Bi₄V₂O₁₁ displays two advantages compared with BiVO₄. First, the band gap of Bi₄V₂O₁₁ (2.15 eV) is narrower than that of BiVO₄ (2.4 eV), which means that Bi₄V₂O₁₁ can absorb more solar

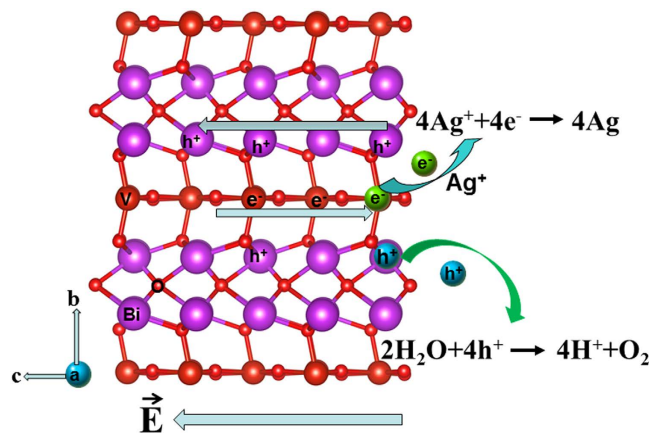


Figure 4. Schematic illustration of the photocatalytic process of $\text{Bi}_4\text{V}_2\text{O}_{11}$.

light. Second, the internal polar field of $\text{Bi}_4\text{V}_2\text{O}_{11}$ can facilitate the effective separation of photo-generated electron and hole pairs, which is in favour of high photocatalytic O_2 evolution.

In summary, the hierarchical $\text{Bi}_4\text{V}_2\text{O}_{11}$ photocatalyst was synthesized via one pot solvothermal method, which is verified to display excellent photocatalytic O_2 evolution activity under light irradiation. The layered structure can effectively facilitate the spatial separation of photoinduced charges. Moreover, the intrinsic internal polar field makes the photogenerated electrons and holes move along opposite directions. The two factors are believed to be advantageous for the separation of photogenerated charge carriers and therefore responsible for excellent photocatalytic performance of $\text{Bi}_4\text{V}_2\text{O}_{11}$ towards O_2 evolution.

Methods

Hydrothermal synthesis of $\text{Bi}_4\text{V}_2\text{O}_{11}$. All reagents used in this study were purchased from the Sinopharm Chemical Reagent Corporation (Shanghai, China) and they were of analytical grade without further purification. $\text{Bi}_4\text{V}_2\text{O}_{11}$ photocatalyst was prepared via one-pot facile solvothermal method according to ref. 15. In a typical process, 5 mmol $\text{Bi}(\text{NO}_3)_3 \cdot 5\text{H}_2\text{O}$ and 1.0 g urea were dissolved into 70 mL ethylene glycol (EG) solution with vigorous stirring. Afterwards, 2.5 mmol NH_4VO_3 were added into the above solution. The pH was adjusted to 7.5 by the addition of diluent ammonia water. The resulting precursor suspension was transferred into a 100 mL Teflon-lined stainless autoclave and maintained at 453 K for 24 h. And then, the reactor was allowed to cool to room temperature naturally. The resulting product $\text{Bi}_4\text{V}_2\text{O}_{11}$ was washed with deionized water and absolute ethanol for several times and dried at 333 K for 6 h in oven.

Characterization. XRD pattern of the $\text{Bi}_4\text{V}_2\text{O}_{11}$ was recorded on a Bruker AXS D8 advance powder diffractometer with Cu K α X-ray radiation. The morphology was investigated by a scanning electron microscopy (SEM, Hitachi S-4800), and the diffuse reflection spectra (DRS) by a Shimadzu UV 2550 recording spectrophotometer. XPS measurement was carried out on a VG MicroTech ESCA 3000 X-ray photoelectron spectroscope with a monochromatic Al-K α source to explore the elements on the surface. The time-resolved fluorescence spectrum was measured via Edinburgh FLS920 PL. Steady state fluoresce was performed via using an Edinburgh FS920 high sensitivity fluorescence spectrometer.

Computation. The spin-polarized density function theory (DFT) calculations were performed by Vienna ab-initio simulation package (VASP). The generalized gradient approximation (GGA) of Perdew, Burke, and Ernzerhof (PBE) was employed for the exchange-correlation functional. The cut-off energy of 400 eV was adopted for the plane-wave basis set. The convergence threshold for self-consistent iteration is set at 10 $^{-6}$ eV, and all atomic structures were fully relaxed until the residual forces on all atoms were smaller than 0.01 eV/Å.

According to the Mulliken electronegativity theory, the potentials of the conduction band (CB) and valence band (VB) edges of $\text{Bi}_4\text{V}_2\text{O}_{11}$ were calculated, whose equations is following:

$$E_{\text{CB}}^0 \approx E_{\text{CB}} = \chi_{\text{comp}} - E^{\text{e}} - \frac{1}{2}E_{\text{g}} \quad (2)$$

$$\chi_{\text{comp}} = \sqrt[N]{\chi_1^r \chi_2^s \cdots \chi_{n-1}^p \chi_n^q} \quad (3)$$

$$N = r + s + \cdots + p + q \quad (4)$$

$$E_{\text{g}} = E_{\text{VB}} - E_{\text{CB}} \quad (5)$$

χ_{comp} represents the absolute electronegativity of the semiconductors depending on the species and number of the constituent atoms and given by geometric mean of the absolute electronegativity of each atom and the

total number of atoms (equations (3) and (4); E^e is defined as the energy of free electrons on the hydrogen scale ($E^e = \sim 4.5 \text{ eV}$), in addition, E_g is the band gap energy of the semiconductor. According to the above equation (3) and (4), χ_{comp} value is evaluated to be 6.30 eV. And then, the values of E_{CB} and E_{VB} are calculated according to the equation (2) and (5), respectively.

Photocatalytic water oxidation activity. The photocatalytic O_2 evolution reaction was performed in a quartz reactor at constant temperature with circulation of water through the internal U-type glass tube from a thermostatic bath. The typical experimental procedure is as follows: first, 100 mg samples were dispersed in 100 mL of aqueous AgNO_3 solution (0.015 M) and sealed the reactor. Subsequently, argon flow has been purged the solution for 30 min to drive away the residual oxygen. And then, the mixture solution was irradiated on the 300 W Xe lamp (PLS-SXE300, Beijing Trusttech Co., Ltd.). The evolution amount of O_2 gas was measured with the gas chromatography with a thermal conductivity detector.

Electrochemical performances. Photocurrent measurements for $\text{Bi}_4\text{V}_2\text{O}_{11}$ were carried out with a CHI 660 C electrochemical workstation. A 300 W Xe arc lamp was utilized as the light source. The $\text{Bi}_4\text{V}_2\text{O}_{11}$ sample (10 mg) was spin coated on a $1.5 \times 1.5 \text{ cm}^2$ ITO glass electrode. The ITO glass was used as working electrodes. In addition, a Pt was employed as the counter electrode and a saturated calomel electrode was used as reference electrode. 0.2 M Na_2SO_4 solution was used as the electrolyte. The electrochemical impedance spectroscopy (EIS) was performed at the open circuit potential.

References

- Fujishima, A. & Honda, K. Electrochemical Photolysis of Water at a Semiconductor Electrode. *Nature* **238**, 37–38 (1972).
- Ran, J. G., Zhang, J., Yu, J. G., Jaroniec, M. & Qiao, S. Z. Earth-abundant cocatalysts for semiconductor-based photocatalytic water splitting. *Chem. Soc. Rev.* **43**, 7787–7812 (2014).
- Chen, X., Shen, S., Guo, L. & Mao, S. S. Semiconductor-based Photocatalytic Hydrogen Generation. *Chem. Rev.* **110**, 6503–6570 (2010).
- Asahi, R., Morikawa, T., Ohwaki, T., Aoki, K. & Taga, Y. Visible-Light Photocatalysis in Nitrogen-Doped Titanium Oxides. *Science* **293**, 269–271 (2001).
- Hagiwara, H., Inoue, T., Kaneko, K. & Ishihara, T. Charge-Transfer Mechanism in $\text{Pt}/\text{KTa}(\text{Zr})\text{O}_3$ Photocatalysts Modified with Porphyrinoids for Water Splitting. *Chem. Eur. J.* **15**, 12862–12870 (2009).
- Zhu, L. L., Hong, M. H. & Ho, G. W. Hierarchical Assembly of SnO_2/ZnO Nanostructures for Enhanced Photocatalytic Performance. *Sci. Rep.* **5**, 11609 (2015).
- Wang, G. *et al.* Synthesis and characterization of ZnS with controlled amount of S vacancies for photocatalytic H_2 production under visible light. *Sci. Rep.* **5**, 8544 (2015).
- Hou, H. L. *et al.* Efficient Photocatalytic Activities of TiO_2 Hollow Fibers with Mixed Phases and Mesoporous Walls. *Sci. Rep.* **5**, 15228 (2015).
- Kudo, A. & Miseki, Y. Heterogeneous photocatalyst materials for water splitting. *Chem. Soc. Rev.* **38**, 253–278 (2009).
- Wang, W. Z. *et al.* p-n junction CuO/BiVO_4 heterogeneous nanostructures: synthesis and highly efficient visible-light photocatalytic performance. *Dalton Trans.* **43**, 6735–6743 (2014).
- Wang, D. *et al.* Photocatalytic Water Oxidation on BiVO_4 with the Electrocatalyst as an Oxidation Cocatalyst: Essential Relations between Electrocatalyst and Photocatalyst. *J. Phys. Chem. C.* **116**, 5082–5089 (2012).
- Li, Y. Q. *et al.* Enhancing the Efficiency of Water Oxidation by Boron-Doped BiVO_4 under Visible Light: Hole Trapping by BO_4 Tetrahedra. *Chempluschem* **80**, 1113–1118 (2015).
- Hong, S. J., Lee, S., Jang, J. S. & Lee, J. S. Heterojunction $\text{BiVO}_4/\text{WO}_3$ electrodes for enhanced photoactivity of water oxidation. *Energy Environ. Sci.* **4**, 1781–1787 (2011).
- Ho-Kimura, S. M., Moniz, S. J. A., Handoko, A. D. & Tang, J. W. Enhanced photoelectrochemical water splitting by nanostructured $\text{BiVO}_4\text{-TiO}_2$ composite electrodes. *J. Mater. Chem. A.* **2**, 3948–3953 (2014).
- Chen, X. F. *et al.* One-step approach to novel $\text{Bi}_4\text{V}_2\text{O}_{11}$ hierarchical hollow microspheres with high visible-light-driven photocatalytic activities. *J. Mater. Chem. A.* **1**, 877–883 (2013).
- Al-Areqi, N. A. S., Al-Alas, A., Al-Kamali, A. S. N., Ghaleb, K. A. S. & Al-Mureish, K. Photodegradation of 4-SPPN dye catalyzed by Ni(II)-substituted $\text{Bi}_2\text{VO}_{5.5}$ system under visible light irradiation: Influence of phase stability and perovskite vanadate-oxygen vacancies of photocatalyst. *J. Mol. Catal. A-Chem.* **381**, 1–8 (2014).
- Jiang, J., Zhao, K., Xiao, X. Y. & Zhang, L. Z. Synthesis and facet-dependent photoreactivity of BiOCl single-crystalline nanosheets. *J. Am. Chem. Soc.* **134**, 4473–4476 (2012).
- Prasad, K. V. R. & Varma, K. B. R. Pyroelectric properties of $\text{Bi}_2\text{VO}_{5.5}$ ceramic. *J. Phys. D: Appl. Phys.* **24**, 1858–1860 (1991).
- Lu, Y. T. *et al.* On structure and methylene blue degradation activity of an Aurivillius-type photocatalyst of $\text{Bi}_4\text{V}_2\text{O}_{11}$ nanoparticles. *Appl. Surf. Sci.* **347**, 719–726 (2015).
- Zhang, R. *et al.* Layered photocatalyst $\text{Bi}_2\text{O}_2[\text{BO}_2(\text{OH})]$ nanosheets with internal polar field enhanced photocatalytic activity. *CrystEngComm* **16**, 4931–4934 (2014).
- Wang, W. J. *et al.* Efficient Separation of Photogenerated Electron-Hole Pairs by the Combination of a Heterolayered Structure and Internal Polar Field in Pyroelectric BiOIO_3 Nanoplates. *Chem. Eur. J.* **19**, 14777–14780 (2013).
- Liu, Z. S., Niu, J. N., Feng, P. Z., Sui, Y. W. & Zhu, Y. B. One-pot synthesis of $\text{Bi}_{24}\text{O}_{31}\text{Br}_{10}/\text{Bi}_4\text{V}_2\text{O}_{11}$ heterostructures and their photocatalytic properties. *RSC Adv.* **4**, 43399–43405 (2014).
- Lv, C. *et al.* Realizing nanosized interfacial contact via constructing $\text{BiVO}_4/\text{Bi}_4\text{V}_2\text{O}_{11}$ element-copied heterojunction nanofibres for superior photocatalytic properties. *Appl. Catal. B-Environ.* **179**, 54–60 (2015).
- Trzcinski, K., Borowska-Centkowska, A., Sawczak, M. & Lisowska-Oleksiak, A. Photoelectrochemical properties of BIMEVOX (ME = Cu, Zn, Mn) electrodes in contact with aqueous electrolyte. *Solid State Ionics.* **271**, 63–68 (2015).
- Long, M. C. *et al.* Efficient Photocatalytic Degradation of Phenol over $\text{Co}_3\text{O}_4/\text{BiVO}_4$ Composite under Visible Light Irradiation. *J. Phys. Chem. B.* **110**, 20211–20216 (2006).
- Lin, X. P. *et al.* Photocatalytic Activities of Heterojunction Semiconductors $\text{Bi}_2\text{O}_3/\text{BaTiO}_3$: A Strategy for the Design of Efficient Combined Photocatalysts. *J. Phys. Chem. C.* **111**, 18288–18293 (2007).
- Huang, Q. W. *et al.* Enhanced Photocatalytic Activity of Chemically Bonded $\text{TiO}_2/\text{Graphene}$ Composites Based on the Effective Interfacial Charge Transfer through the C-Ti Bond. *ACS Catal.* **3**, 1477–1485 (2013).

Acknowledgements

This work was financially supported by the National Basic Research Program of China (the 973 Program, No. 2013CB632401), the National Natural Science Foundation of China (No. 21573135, 21333006, 21007031,

11374190 and 51002091), Taishan Scholar Foundation of Shandong Province, China and the Shandong Province Natural Science Foundation (ZR2014JL008).

Author Contributions

Z.Y.J., Y.Y. and B.B.H. designed the research and carried out the experiments. M.M.L., T.J. and Y.D. performed the theory calculations. Z.Y.J., Y.Y.L., X.Y.Z., X.Y.Q. and B.B.H. analysed the experimental data. Z.Y.J., Y.Y.L. and B.B.H. prepared the manuscript. All authors contributed to the scientific discussions.

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: Jiang, Z. *et al.* One-Pot Solvothermal Synthesis of $\text{Bi}_4\text{V}_2\text{O}_{11}$ as A New Solar Water Oxidation Photocatalyst. *Sci. Rep.* **6**, 22727; doi: 10.1038/srep22727 (2016).



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/>