

Supporting Information for

**Ultrathin BiVO₄ Nanobelts: Controllable Synthesis and Improved
Photocatalytic Oxidation of Water**

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Experimental Section

Chemical Reagents and Instruments: Ammonium metavanadate (NH_4VO_3), bismuth nitrate ($\text{Bi}(\text{NO}_3)_3$), ammonia ($\text{NH}_3 \cdot \text{H}_2\text{O}$), nitric acid (HNO_3), methanol (CH_3OH), silver nitrate (AgNO_3) were purchased from Sinopharm Chemical Reagent Co., Ltd. Deionized water (18 M Ω , Molecular) was used for all solution preparations.

The morphology and size of the as-prepared products were characterized by using a field-emission scanning electron microscope (JSM-6701F, JEOL). The X-ray diffraction spectra (XRD) measurements were performed on a PANalytical X'Pert PRO instrument using Cu K α radiation (40 kV). The XRD patterns were recorded from 10° to 90° with a scanning rate of 0.067 °/s. UV-visible diffuse reflectance spectra were taken on a UV-2550 (Shimadzu) spectrometer by using BaSO_4 as the reference. HRTEM imaging was carried out using an FEI Tecnai TF20 microscope operated at 200 kV.

Preparation of BiVO_4 nanobelts: Firstly, 0.036 g NH_4VO_3 was dissolved in 1 M ammonia solution (solution A) because it is slight soluble in pure water. 0.03 g BiNO_3 was dissolved into 1 M dilute nitric acid (solution B) since it could hydrolyze to BiONO_3 precipitates in aqueous solution. Secondly, solution A was added into solution B drop by drop. Along with the consumption of ammonia and nitric acid, NH_4VO_3 and BiNO_3 would react and produce tetragonal BiVO_4 nanoparticles. The PH is measured to be 5.1 after the full mixture of solution A with solution B. At last, the solution was transferred into a Teflon-lined stainless steel autoclave, which was kept at 180 °C for one day. After the hydrothermal reaction, the BiVO_4 nanobelts were obtained finally.

Preparation of BiVO_4 nanoparticles: The pure BiVO_4 nanoparticles was synthesized via a typical procedure, 1 mmol of $\text{Bi}(\text{NO}_3)_3 \cdot 5\text{H}_2\text{O}$ and 1mmol of NH_4VO_3 was added to 30 mL of 2 M nitric acid solution at room temperature and remained continuous stirring for 30 minutes. Then, an appropriate amount of NH_4OH (25~28%) was also added into the prepared solution to fit the pH value to 8. After stirring for 30 min, a clear orange solution was obtained. This solution was poured into a 50 mL Teflon-lined autoclave and maintained at 180°C for 24 h under autogenous pressure, and then naturally cooled to room temperature. The resulting precipitates were collected and washed with ethanol and deionized water thoroughly and dried at 80°C in air.

Photoelectrochemical measurements: The BiVO_4 nanobelts suspended in water and the

mixtures were ultrasonically scattered for 15 min to form homogeneous solution. Then, the solution was spin coated on a Fluorine-doped tin oxide (FTO) substrate with a rate of 300 rpm for 30 s. This procedure was repeated for 3 times. Photoelectrochemical properties were measured in a three-electrode configuration. A Pt foil, saturated calomel electrode (SCE), and 0.2 M sodium sulfate with methanol were used as the counter-electrode, the reference electrode, and the electrolyte, respectively. The current-time (i-t) curves were collected at 0.4 V vs SCE. A 300 W Xe lamp was utilized as the light source.

Photocatalytic reactions: The photocatalytic O₂ evolution was carried out with 0.3 g photocatalyst suspending in AgNO₃ solution in a Pyrex glass reaction cell. The reaction cell was connected to a gas-closed system with a gas-circulated pump. A 300 W Xe arc lamp was employed for the light source of photocatalytic reaction. The evolved O₂ was analyzed by an online gas chromatograph (GC-8A; Shimadzu Corp., Japan) equipped with a thermal conductivity detector.

Additional Figures

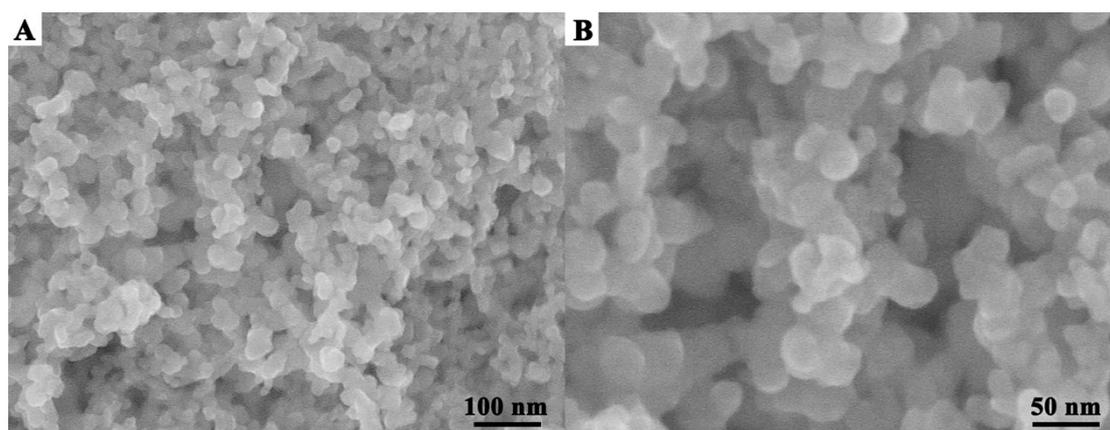


Fig. S1. SEM images of BiVO₄ nanoparticles before hydrothermal reaction.

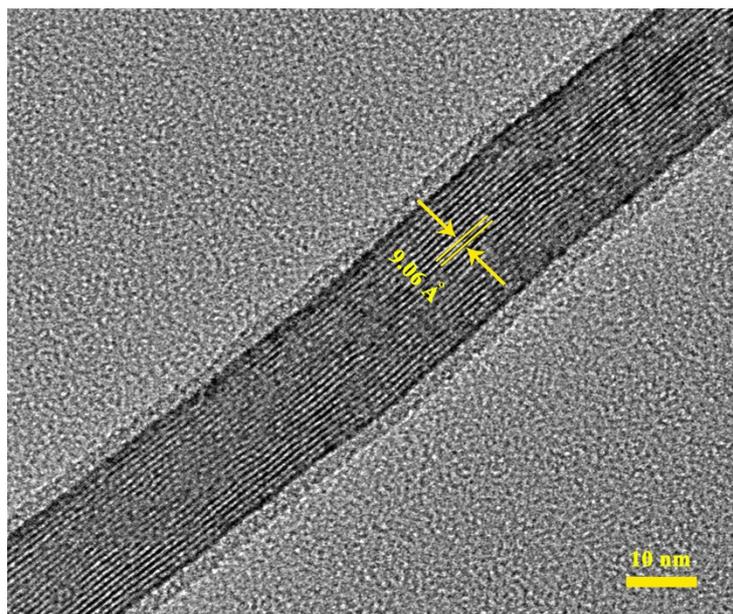


Fig. S2. Sideview TEM image of BiVO₄ nanobelts.

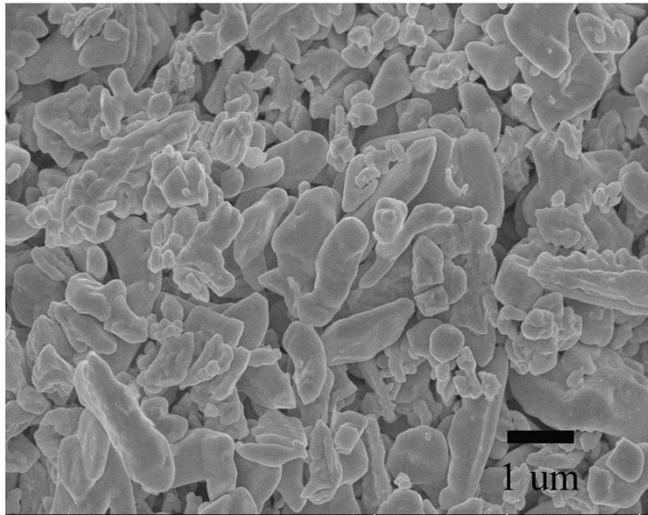


Fig. S3. SEM image of BiVO₄ nanoparticles.

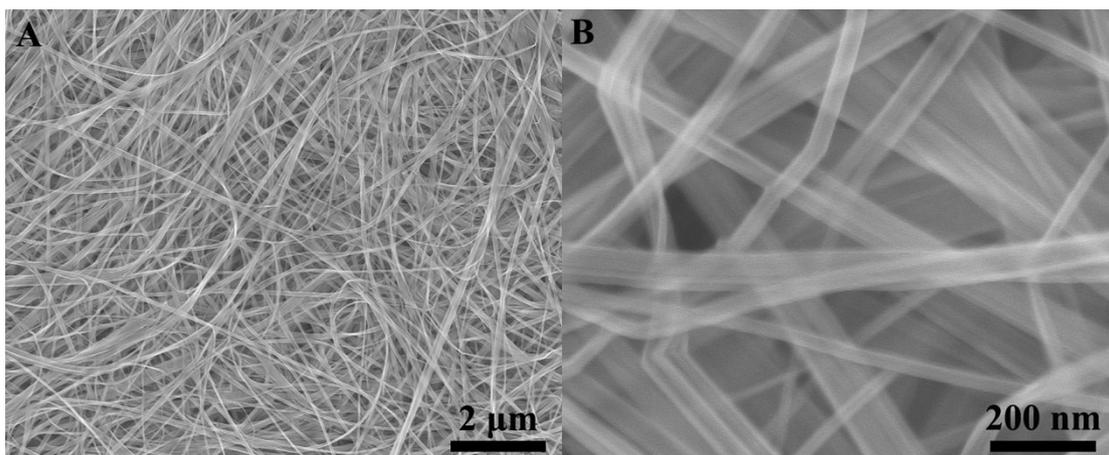


Fig. S4. SEM images of ultrathin BiVO_4 nanobelts after photocatalytic oxidation of water.