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Supporting Information

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Layered VS₂ Nanosheet-Based Aqueous Zn Ion Battery Cathode

Pan He, Mengyu Yan, Guobin Zhang, Ruimin Sun, Lineng Chen, Qinyou An,* and Liqiang Mai*

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Experimental Section

Synthesis of VS_2 nanosheets: VS_2 nanosheets were prepared via a simple hydrothermal method. Briefly, 2 mmol NH₄VO₃ was dissolved in 30 ml deionized water and 2 mL NH₃.H₂O in a glass jar. Then, 15 mmol thioacetamide (TAA) was added in the homogeneous solution with continuous magnetic stirring at room temperature for 1 h. After that, the mixture was transferred to a 50 mL Teflon-lined sealed autoclave and maintained at 180°C for 20 h. Afterward, the system was cooled down to room temperature naturally and the samples were washed with distilled water and ethanol thoroughly for 3 times, respectively. The final product was dried at 60°C for 8 h in vacuum, and the black powder was obtained.

Material characterizations: The as-prepared samples were characterized by power Xray diffraction (XRD, D8 Discover X-ray diffractometer with Cu K_{α} radiation), X-ray photoelectron spectroscopy (XPS, Thermo Scientific Escalab 250Xi), Raman spectra (Renishaw INVIA), field emission scanning electron microscopy (FESEM, JSM-7100F) transmission electron microscopy (TEM) and energy dispersive spectroscopy (JEM-2100F, STEM/EDS).

Electrochemical characterizations: The VS₂ electrode was prepared by mixing VS₂ (60 wt%), acetylene black (Super-P, 30 wt%), and poly tetrafluoroethylene (PTFE, 10 wt%), then the slurry was evenly grinded, tableted and cut into Φ 10 mm electrodes. Zinc foil and glass fiber membrane were used as the anode and separator, respectively, and 1 M zinc sulfate electrolyte solution was used as the electrolyte. A CR2016-type coin cell was assembled in the air atmosphere to evaluate the electrochemical performance on a LAND battery testing system (CT2001A). Cyclic voltammograms (CV) were test on a CHI600E electrochemical workstation. All of the tests were performed at room temperature.

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Figure S1. TEM-EDS of VS_2 nanosheets



Figure S2. The growth process of VS_2 nanosheets at different reaction times (from 0 h to 20 h).



Figure S3. a) The charge and discharge curves of VS₂ at 0.1 A g^{-1} , b) Cyclic properties



Figure S4. The charge and discharge curves of VS_2 of different cycles at a current density of 0.5 A g⁻¹.



Figure S5. a) The CV curves of VS_2 at a scan rate of 0.1 mV/s, b) The contribution ratio of the capacitive capacities and diffusion-limited capacities at 0.1 mV/s.



Figure S6. The charge and discharge curves of VS₂ at 0.05 A g^{-1} .

In the first step: $VS_2 + xZn^{2+} + 2xe^- \leftrightarrow Zn_xVS$ (x = 0.09)

In the second step: $Zn_xVS_2 + yZn^{2+} + 2ye^- \leftrightarrow Zn_{x+y}VS_2$ (y = 0.14)

x and y are calculated based on the following equations: (F = N_A*e = 96500 C/mol, N_A = 6.02×10^{23} , 1 A h = 1 A × 3600s = 3600 C, C₀ = 26.8 nm/M)