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Electronic Supplementary Information

3D Porous Hierarchical Li₂FeSiO₄/C for Rechargeable Lithium Batteries

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Materials characterization

Crystallographic information of the samples was identified by X-ray diffraction (XRD) using a Rigaku Dmax-2400 automatic diffractometer (Cu $K\alpha$). Material shape and morphology were observed by scanning electron microscopy (SEM, Hitachi S-4800) and transmission electron microscopy (TEM, FEI Tecnai G2 T20). Raman spectroscopy measurements were conducted on a JY HR800 spectrometer using an excited HeNe laser (633 nm). Thermogravimetric (TG) analysis was performed on a TG/DTA-7300 thermal analyzer (Seko). Nitrogen adsorption and desorption isotherms were measured on a Tristar 3020 micropore analyzer (Micromeritics).

Electrochemical evaluation

Electrochemical tests were performed on a 2032-type coin cell. The working electrodes were composed of 80 wt% active material, 10 wt% acetylene black, and 10 wt% polyvinylidene fluoride binder. The mass loading for the composite electrode is 2.0–2.5 mg cm⁻². Li foil was used as the counter electrode and GFA glass fiber filters (Whatman) as separator. The electrolyte is 1 mol l^{-1} LiPF₆ solution dissolved in ethylene carbonate/dimethyl carbonate (1:1 by volume). Cyclic voltammetry and electrochemical impedance spectroscopy were conducted on a CHI 660E electrochemical workstation. Galvanostatic tests were carried out on a Land battery test system.



Figure S1. SEM images of 3D porous hierarchical Li₂FeSiO₄/C.



Figure S2. Energy dispersive X-ray spectroscopy of 3D porous Li_2FeSiO_4/C .



Figure S3. SEM images of (a) hydrothermal intermediate and (b) 3D Li₂FeSiO₄.



Figure S4. TG and DTG curves of 3D porous Li₂FeSiO₄/C.



Figure S5. Nyquist plots of the impedance spectra of the 3D porous Li_2FeSiO_4/C after one and 20 cycles.



Figure S6. Typical galvanostatic curves of 3D porous Li₂FeSiO₄ after activation.



Figure S7. Rate capability of 3D porous Li₂FeSiO₄.