Dendritic nanostructured FeS$_2$-based high stability and capacity Li-ion cathodes†

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Introduction

In the quest for improved lithium-ion batteries (LIBs), considerable effort has been made to enhance the electrochemical properties of cathode materials.1 Along with development of new materials,2–5 an increasing number of groups are paying attention to the relationship between the electrochemical properties and the morphology of the active material.6,7 Nanostructuring offers a number of advantages, including shorter solid state ionic and electron transport pathways, and an increased contact area with the electrolyte, which improves the rate performance.8–11 However, this increased surface area can also lead to an increase in solid electrolyte interphase formation, which may be detrimental to the cycle life. Additionally, maintaining both good electron and ion transport pathways in a nanostructured system can be challenging.

†Electronic supplementary information (ESI) available: Energy Dispersive Spectroscopy (EDS), more electrochemical tests (including electrochemical impedance spectroscopy (EIS), cycling performance tests and cyclic voltammetry (CV) tests) and high-resolution transmission electron microscopy (HRTEM) images are available. See DOI: 10.1039/c8ra07606e
The pure iron pyrite nanowire cathode materials synthesized by Song et al.\textsuperscript{38} retained a discharge capacity of 350 mA h g$^{-1}$ after 50 cycles and 0.1C, possibly because of the improved reaction kinetics provided by the nanostructure. Liu et al.\textsuperscript{37} coated a carbon layer on octahedral FeS$_2$ nanocrystallites to enhance electrical conductivity and structure stability, and this system exhibited a capacity retention of 495 mA h g$^{-1}$ after 50 cycles at 0.5C. Wang et al.\textsuperscript{31} reported hierarchically structured hollow FeS$_2$ microspheres which showed reasonable capacity and cycling performance (392.7 mA h g$^{-1}$ after 30 cycles at 1C). All of these works emphasized the importance of nano-structures for improving the electrochemical properties of pyrite cathodes. Here, we describe a facile self-assembly route for preparing dendritically structured pyrite-based Li-ion cathodes. These dendritic FeS$_2$ composite cathodes exhibit high specific capacities and good cycling performance in the voltage window of both 0.7–2.4 V and 0.7–3.0 V. The Li$_{1.2-}$FeS$_2$ phase was observed by high-resolution transmission electron microscopy (HRETEM) and electron energy loss spectroscopy (EELS) in scanning transmission electron microscopy (STEM) mode. Identification of this phase may enable further understanding of the lithiation/delithiation mechanism of FeS$_2$.

### Experimental section

#### 1. Synthesis of the hierarchically structured FeS$_2$

Ferrous heptahydrate ($\text{FeSO}_4 \cdot 7\text{H}_2\text{O}$, Sigma Aldrich) was used to synthesize the dendritically structured $\alpha$-Fe by an electric-field-induced and electrochemical reduction method.\textsuperscript{34} The hierarchically structured dendritic pyrite was obtained via thermal sulfidation of the as-synthesized dendritic $\alpha$-Fe at 450 °C for 4 hours with a heating and cooling rate of 15 °C per minute under a continuous argon gas flow. Elemental sulfur (99.5–100.5%, Sigma Aldrich) was placed at the edge of the furnace, where it reached about 140 °C and formed a melt. The resulting sulphur vapour was carried by the argon gas flow over the dendritic nanostructured $\alpha$-Fe.

#### 2. Material characterization

Sample composition and microstructure were determined by: X-ray diffraction (Philips X’pert XRD system with Cu Kα radiation, λ = 0.15418 nm); scanning electron microscopy and energy dispersive spectroscopy (Hitachi S-4700 SEM, EDS); transmission electron microscopy and high-resolution transmission electron microscopy (JEOL 2100 cryo, TEM and HRETEM); and electron energy loss spectroscopy (EELS) during scanning transmission electron microscopy (JEOL, 2010F STEM). Cycling of TEM and STEM-EELS samples were performed by dispersing the dendritic nanostructured FeS$_2$ into ethanol, placing a couple of droplets on a copper grid with a carbon support film (EMS Acquisition Corp), covered by an identical copper grid and dried 2 h at 80 °C. The two grids were held by a toothless clamp and used as a cathode in a bottle cell. 1 M lithium hexafluorophosphate (LiPF$_6$) in a mixture of ethylene carbonate (EC), dimethyl carbonate (DMC) and diethyl carbonate (DEC) (2 : 1 : 2 in volume) was used as the electrolyte. 10 wt% fluoroethylene carbonate (FEC, Sigma Aldrich) was added to the electrolyte. The counter electrode is lithium foil (Alfa Aesar). After 7 charge/discharge cycles using about 0.2C rates between 0.7–2.4 V, the samples were removed from the electrolyte, and immersed into dimethyl carbonate (DMC) solvent for 30 min to remove the rest of electrolyte. The copper grids were taken out of the glove box, and immediately transferred to the TEM column for analysis.

#### 3. Fabrication of the composite cathode and electrochemical measurements

Composite electrodes were fabricated through mixing of the as-synthesized dendritic FeS$_2$, carbon black (Super P Conductive, Alfa Aesar), and poly(vinylidene fluoride) (PVDF, MTI) binder (65 : 25 : 15 weight ratio) using a homogenizer (GLH-115, OMNI International) for 15 min in N-methylpyrrolidone (NMP, MTI). The resulting slurry was printed onto a conductive carbon-coated copper foil current collector (MTI) with an average FeS$_2$ active material mass loading of 1–1.5 mg cm$^{-2}$. Then, the composite electrodes were dried for 10 h at 120 °C. 9.5 mm diameter electrode disks were punched and then dried at 100 °C in a vacuum oven for 12 h before cell assembly.

Two-electrode battery-type Swagelok cells were used to test the electrochemical properties of the composite electrode on a VMP3 (Bio-Logic) potentiostat. Cyclic voltammetry (CV) was performed between 0.7–2.4 V and 0.7–3.0 V at a scan rate of 0.2 mV s$^{-1}$. Electrochemical impedance spectroscopy (EIS) was performed on Swagelok cells after various cycle numbers at 2.4 V over a frequency range of 100 kHz to 10 mHz with an AC amplitude of 5.0 mV. The cells were kept at 2.4 V for 4 h, and allowed to rest for 4 h before taking EIS measurements.

### Results and discussion

Fig. 1a–c shows scanning electron microscopy (SEM) images of the as-synthesized FeS$_2$. The dendritic FeS$_2$ particles are 8 to 15...
μm in length and 4 to 6 μm in width, and possess a fine nanostructure on the branches and tips (in Fig. 1c). Energy dispersive spectroscopy (EDS) (Fig. S1†) indicates the atomic ratio of S to Fe is 2 : 1. All peaks in the X-ray diffraction (XRD) pattern (in Fig. 1d) can be indexed to pure cubic FeS₂ (JCPDS no. 65-7643, a = 0.5404 nm). No impurities such as elemental Fe, elemental S or marcasite were detected.

The electrochemical performance of as-synthesized dendritic nanostructured FeS₂ was studied in two-electrode battery-type Swagelok cells. The cycling and rate performance of the composite cathodes containing 60 wt% dendritic FeS₂ over the voltage range of 0.7–2.4 V and 0.7–3.0 V (in Fig. 2a and b) were investigated. Fig. 2a shows the cycling performance of the dendritic structural FeS₂ composite electrode between 0.7–2.4 V in an electrolyte with 10 wt% fluoroethylene carbonate (FEC) added. The specific capacities of the dendritic FeS₂ cathode are 566 mAh g⁻¹ (0.2C) and 449 mAh g⁻¹ (1.0C) in the 1st cycle, and 348 mAh g⁻¹ (0.2C) and 179 mAh g⁻¹ (1.0C) after 150 cycles, exceeding that reported for pyrite nano-wires over 150 cycles, exceeding that reported for pyrite nano-wires.

When cycling over 0.7–3.0 V, the specific capacities (in Fig. 2b) of a dendritic FeS₂ composite cathode could reach 796 mAh g⁻¹ (0.5C) and 717 mAh g⁻¹ (1.0C) after the 1st cycle, and 560 mAh g⁻¹ (0.5C) and 533 mAh g⁻¹ (1.0C) after 50 cycles, which is about 70% of the 2nd cycle capacity and superior to the performances of polyacrylonitrile (PNA)-FeS₂ (470 mAh g⁻¹ after 50 cycles at 0.1C) and FeS₂ stabilization with a bis[(trifluoromethanesulfonyl)imide (TFSI⁻) anion-based ionic liquid electrolyte over 1.0–3.0 V, but lower than that of pyrite nanocrystals (≥630 mAh g⁻¹) at the slow cycling rate of 200 mA g⁻¹. Importantly, the dendritic FeS₂ cathodes always show a high coulombic efficiency (CE) for different C rates. The average CE after 5 cycles exceeded 99.5% for 0.7–2.4 V and 98.9% for 0.7–3.0 V, indicating a good electrochemical reversibility. The electrochemical performance of some FeS₂-based materials are shown in Table S1.† Fig. 2c shows voltage profiles between 0.7–2.4 V after the 10th cycle at each cycling rate. As the current density increases, the specific capacity decreases from 540 mAh g⁻¹ at 0.2C to 490 mAh g⁻¹, 400 mAh g⁻¹, 220 mAh g⁻¹ and 80 mAh g⁻¹ at 0.5C, 1C, 1.5C and 3C, respectively. Meanwhile, the conversion plateaus become gradually sloped and shorter due to voltage polarization at higher C rates. There is the same change trend for the conversion plateaus in the 0.7–3.0 V range as shown in Fig. 2d. The consecutive rate capability tests (in Fig. 2e and f) also present good cycling stability at each rate. The capacity recovers even after high C rate charge/discharge processes, indicating that high C rates did not damage the electrode.

Cyclic voltammetry (CV) curves of this composite cathode provide some information on the charge/discharge mechanism of FeS₂.

As shown in Fig. 3a and c, both CV curves show a broad reduction peak at approximately 1.2 V and a broad oxidation peak at about 2.0 V in the 1st cycle. This broad reduction peak has been suggested to correspond to the following reactions:

$$\text{FeS}_2 + 2\text{Li}^+ + 2\text{e}^- \rightarrow \text{Li}_2\text{FeS}_2$$  \hspace{1cm} (1)

$$\text{Li}_2\text{FeS}_2 + 2\text{Li}^+ + 2\text{e}^- \rightarrow 2\text{Li}_2\text{S} + \text{Fe}^0$$  \hspace{1cm} (2)

Due to the relatively slow diffusion of Li⁺ into FeS₂, reactions (1) and (2) proceed simultaneously at room temperature (≤30 °C), and thus only single broad reduction peak appears. The oxidation peak includes the reactions.
2Li₂S + Fe⁰ → Li₂FeS₂ + 2Li⁺ + 2e⁻  \hspace{1cm} (3)

\begin{align*}
\text{Li}_2\text{FeS}_2 & \rightarrow \text{Li}_{2-x}\text{FeS}_2 + x\text{Li}^+ + xe^- (0 < x < 0.8) \hspace{1cm} (4)
\end{align*}

Generally, the hexagonal Li₂₋ₓFeS₂ phase begins to decompose into ortho-FeS₂ above 2.4 V as described in reaction (5),

\[\text{Li}_2₋ₓ\text{FeS}_2 \rightarrow 0.8\text{ortho-FeS}_2 + 0.2\text{FeS}_{8/7} + 0.175\text{S} + (2₋ₓ)\text{Li}^+ + (2₋ₓ)e^- \hspace{1cm} (5)\]

leading to another oxidation near 2.6 V (see Fig. 3c). Because the electrode does not fully revert to FeS₂ during oxidation, even above 2.4 V, the CE of the first few cycles is low.

After the first cycle, subsequent cycles between 0.7–2.4 V show a stable redox couple at 1.35 and 2.15 V corresponding to reactions (2), (3) and (4), respectively. The differential capacity versus voltage curves show that the redox reactions proceed at 1.50 and 1.85 V (Fig. 3b) when cycling at 0.2C. The CV curves of the composite cathode show two stable redox reactions during subsequent cycles when cycling over 0.7–3.0 V. In differential capacity curves (Fig. 3d), the peak at 2.50 V is from reaction (5), while the small broad reduction peak at 2.10 V corresponds to reduction of ortho-FeS₂, S and lithium sulphides.\(^{^{47}}\) The S and lithium sulphides are difficult to decompose and dissolve more easily in the electrolyte, which lead to the low reversibility of reaction (5), and probably is the main reason for the rapid capacity fade during cycling between 0.7–3.0 V.\(^{\text{38,48}}\) A 3.0 V cut off voltage does provide greater capacity, but also results in fast capacity fade.

We suggest the FeS₂ electrochemical performance over 0.7–2.4 V can be attributed to the high reversibility of the intermediate Li₁₋ₓFeS₂ and dendritic structure of FeS₂ which provides a large surface area,\(^{\text{50,51}}\) a good electron and ion transport network, and a mechanically robust structure (note, while we cannot extract the active material from a cycled electrode, as Fig. 4a shows, the dendritic structure is retained with cycling). The dendritic structure increases both the reaction area between the active materials and the electrolyte, and shortens the solid-state Li ion diffusion length. These are important parameters at high charge/discharge rates.

As result above, the dendritic nanostructured FeS₂ shows better cycling performance between 0.7 and 2.4 V than between 0.7 and 3.0 V, because of the high reversibility of the intermediate Li₁₋ₓFeS₂ phase. To study the Li ion lithiation/delithiation process and confirm the formation of Li₁₋ₓFeS₂, high-resolution transmission electron microscopy (HRTEM) images and scanning transmission electron microscopy energy loss spectroscopy (STEM-EELS) analyses in spot mode were performed on the dendritic FeS₂ after 7 cycling tests at low charge discharge rates (about 0.2C). The voltage vs. time curve (in Fig. S2†) shows clear charging/discharging conversion plateaus. Fig. 4a shows TEM images of cycled dendritic FeS₂, after the final charging to 2.4 V, indicating that the dendritic structure is retained after cycling. HRTEM images in Fig. 4b show a clear phase boundary on the edge of the dendritic FeS₂. The majority of inner part of the sample shows a 0.271 nm lattice spacing, corresponding to (200) crystal planes of cubic FeS₂. Another HRTEM image after cycling, as well as a HRTEM image before cycling tests is shown in Fig. S3.† These images indicate that after a small number of cycles, the inner parts of the dendrite are still cubic FeS₂ that has not reacted with Li. The lattice spacing near the surface is about 0.60 nm, very close to the theoretical value for the (001) planes in hexagonal-close-packed Li₂FeS₂. The fast Fourier transform (FFT) of the image, shown in top right of Fig. 4b, was taken from the outlined regions in the image and can be indexed to the [110] zone axis of the Li₂FeS₂ phase (crystal data adopted from the International Crystal Structural Database (ISCD #68380)). A simulated electron diffraction pattern of the Li₂FeS₂ phase along the [110] zone axis using the CrystalMaker™ software (right bottom inset in Fig. 4b) is in good agreement with the experimental FFT pattern, suggesting that the outside layer of the structure is the Li₂FeS₂ phase. Due to the low cut off voltage of 0.7 V, lithiation is incomplete, and the material is likely to be Li₁₋ₓFeS₂, which possess the same crystal structure as Li₂FeS₂.

Fig. 4d shows the EELS spectra taken from the body (position A) and edge (position B) of cycled dendritic FeS₂, as indicated in the STEM high angle annular dark field (HAADF) image in Fig. 4c. After background subtraction, a clear wide peak at about 63.5 eV is observed at the edge of cycled dendritic FeS₂, which corresponds to the Li K-edge.\(^{\text{32}}\) The corresponding peak is not found in the intact inner region, indicating that only the outside part of dendrite is lithiated. Our TEM data provide direct evidence that the lithium ions are incorporated into the dendritic FeS₂ nanostructure, resulting in the formation of the Li₁₋ₓFeS₂ phase during the lithiation process.

Besides of the high reversible intermediate Li₁₋ₓFeS₂ phase, the robust dendritic structure is the other key factor of dendritic FeS₂’s good cycling properties. The architecture of FeS₂ provides...
a large surface area, a good electron and ion transport network, and reduce losing of active materials during cycle process. In order to confirming the importance of dendritic structure, the cycling performance of dendritic FeS$_2$ composite cathode were tested in an electrolyte without FEC. The presence of FEC results in formation of a thin and stable SEI (Fig. S4 and Table S2†); which both enhances cycling reversibility, and appears to be an important contributor to retaining the dendritic structure of the electrode (Fig. 5). In a FEC-free electrolyte, the specific capacity of the dendritic FeS$_2$ cathode fades quickly, with the fall-off appearing to increase after about 100 cycles. After 150 cycles at 0.2C, the dendritic structure of FeS$_2$ appears considerably damaged (Fig. 5c) relative to the initial dendritic structure (Fig. 5b). In a 10 wt% FEC-containing electrolyte as already discussed, the dendritic FeS$_2$ composite cathode exhibits a much improved cycling performance (Fig. 2a). What we observe via SEM, is that the dendritic structure is retained in this case (Fig. 5d), a strong indication that the dendritic structure plays an important role in cycling performance.

**Conclusions**

We find that composite cathodes containing dendritic FeS$_2$ demonstrate good reversible capacity and stable cycling over 0.7–2.4 V, and an enhanced capacity but diminished cycling stability over 0.7–3.0 V. The reversible capacities of the dendritic FeS$_2$ cathodes were 560 mAh g$^{-1}$ (0.5C) and 533 mAh g$^{-1}$ (1.0C) after 50 cycles over 0.7–3.0 V, and 348 mAh g$^{-1}$ (0.2C) after 150 cycles over 0.7–2.4 V. We propose that the reasonable cycling performance of the dendritic FeS$_2$ cathode at different C rates is due to the hierarchical structure, which shortens the Li$^+$ diffusion length and increases the contact area with the electrolyte in an architecture that is able under volume changes due to lithiation and delithiation. The use of an FEC-containing electrolyte significantly improved the cycling performance, presumably by resulting in a thin and dense SEI. However, we still observe a steady capacity fade, indicating probably that a more stable SEI is still required, and that a robust artificial SEI, for example such as one formed via atomic layer deposition, might be required to stabilize the dendritic structure over many cycles. We postulate that the Li$_{1.4}$FeS$_2$ phase, which was observed by HRTEM and STEM-EELS, is one reason that the dendritic FeS$_2$ shows better cycling performance over 0.7–2.4 V than over 0.7–3.0 V, where electrolyte-soluble products are formed.

**Conflicts of interest**

There are no conflicts to declare.

**Acknowledgements**

The authors acknowledge support of an AFOSR MURI (FA9550–12–1–0471) and experimental assistance from Dr Ke Sun and Dr Yang Hu. Z. Yu is grateful for fellowship support from the China Scholarship Council.

**References**