Supporting Information

Low Volume Expansion Hierarchical Porous Sulfur-Doped Fe₂O₃@C

with High-Rate Capability for Superior Lithium Storage



Figure S1. (a, b) SEM images, (c) XRD patterns, and (d) TG curve of MIL-88B NSs



Figure S2. (a) SEM image and (b) XRD pattern of Fe₂O₃@C NSs



Figure S3. (a, b) SEM images of Fe₂O₃/C NSs



Figure S4. TEM images of (a) S₀-Fe₂O₃@C and (b) S_{0.15}-Fe₂O₃@C NSs



Figure S5. XPS survey spectra of S_0-Fe_2O_3@C and (b) S_{0.15}-Fe_2O_3@C NSs



Figure S6. Raman spectra of S0-Fe2O3@C and S0.15-Fe2O3@C NSs



Figure S7. TG curves of (a) $Fe_2O_3@C$, (b) S_0 - $Fe_2O_3@C$, and (c) $S_{0.15}$ - $Fe_2O_3@C$ NSs



Figure S8. The first three CV curves at 0.1 mV s⁻¹ of (a) S₀-Fe₂O₃@C and (b) S_{0.3}-Fe₂O₃@C electrodes



Figure S9. Galvanostatic discharge-charge profiles of (a) $S_0\text{-}Fe_2O_3@C$ and $S_{0.3}\text{-}Fe_2O_3@C$ electrodes at 200 mA $g^{-1}.$



Figure S10. Comparison of rate capabilities of $S_{0.15}$ -Fe₂O₃@C electrode and other reported Fe₂O₃-based LIB anode materials.



Figure S11. Cycle performance of $S_{0.15}$ -Fe₂O₃@C electrode at 5.0 A g⁻¹.



Figure S12. Discharge/charge state curves of GITT patterns for S_{0.15}-Fe_2O_3@C and S_0-Fe_2O_3@C electrodes at 200 mA g^{-1}

The Li-ion diffusion coefficients of $S_{0.15}$ -Fe₂O₃@C and S₀-Fe₂O₃@C electrodes can be measured according to the following Equation (1)

$$D_{\rm Li^+} = \frac{4L^2}{\pi\tau} \left(\frac{\Delta E_s}{\Delta E_\tau}\right)^2 \tag{1}$$

where L and τ are the thickness of the electrodes (cm) and the duration of the current pulse (s), respectively. ΔE_s is the change in steady-state voltage (V) for a single-step GITT measurement, and ΔE_{τ} is the voltage change (V) during one discharge pulse regardless of IR drop.

Table S1. The XPS composition analysis (O, S, Fe, C) of S_{0.15}-Fe₂O₃@C and S_{0.3}-Fe₂O₃@C.

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samples	O (at. %)	S (at. %)	Fe (at. %)	C (at. %)
S _{0.15} -Fe ₂ O ₃ @C	11.03	1.57	1.91	85.49
S _{0.3} -Fe ₂ O ₃ @C	9.03	2.17	1.87	86.93

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