

Supporting Information

In situ formation of MoS₂/C nanocomposite as an anode for high-performance lithium-ion batteries

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Supporting materials include:

Part I. Supplementary figures

Figure S1. XPS spectra of the as-prepared samples. High-resolution spectra of Mo3d, S2p, and C1s peaks for MoS₂-only((a),(e),(i)), MoS₂/C-1((b),(f),(j)), MoS₂/C-3((c),(g),(k)), and MoS₂/C-5((d),(h),(l)).

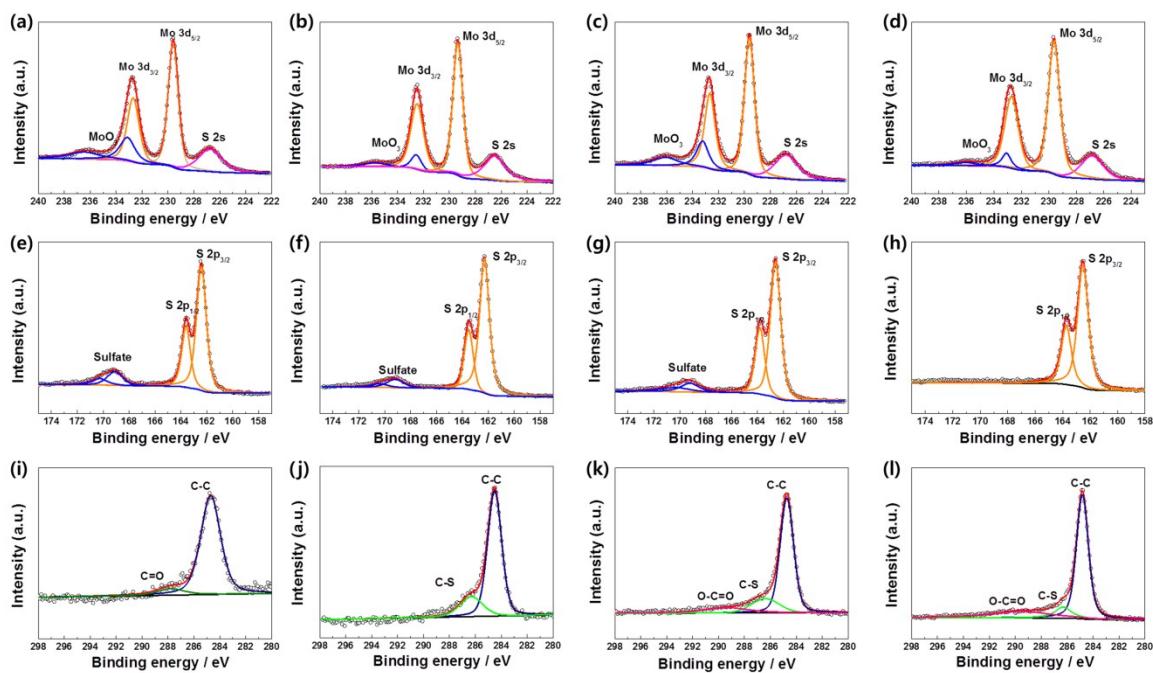


Figure S2. SEM and mapping images of (a) MoS₂-only, (b) MoS₂/C-1, (c) MoS₂/C-3, and (d) MoS₂/C-5.

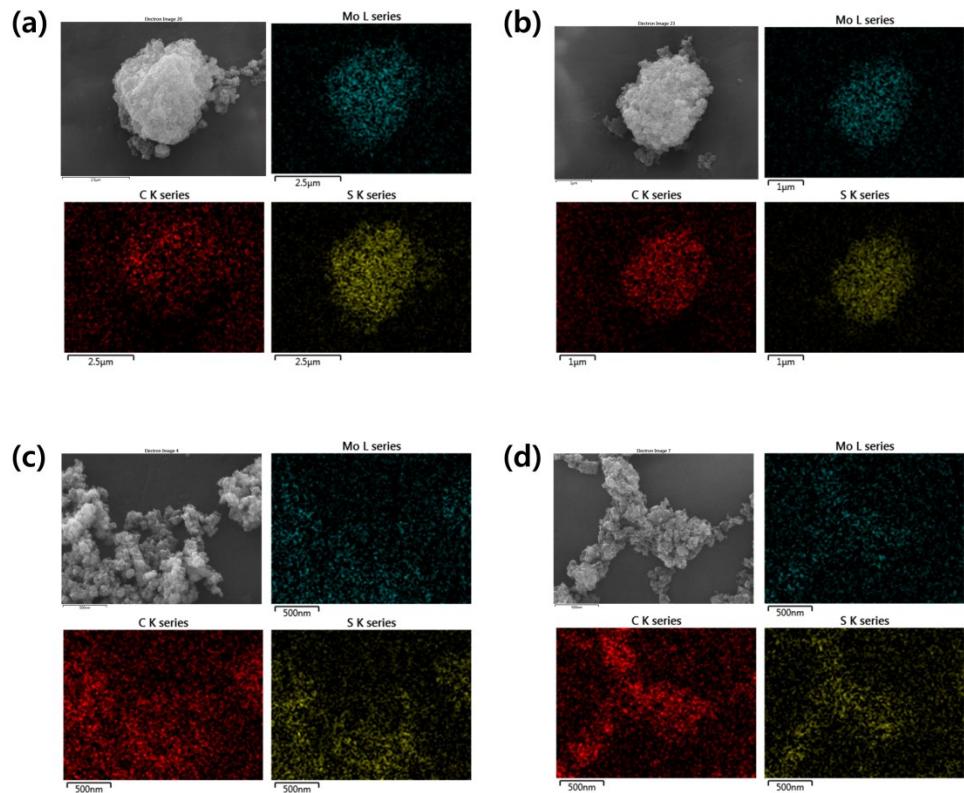


Figure S3. Nitrogen adsorption/desorption isotherms and pore size distributions of (a),(b) MoS₂-only, (c),(d) MoS₂/C-1, (e),(f) MoS₂/C-3, and (g),(h) MoS₂/C-5.

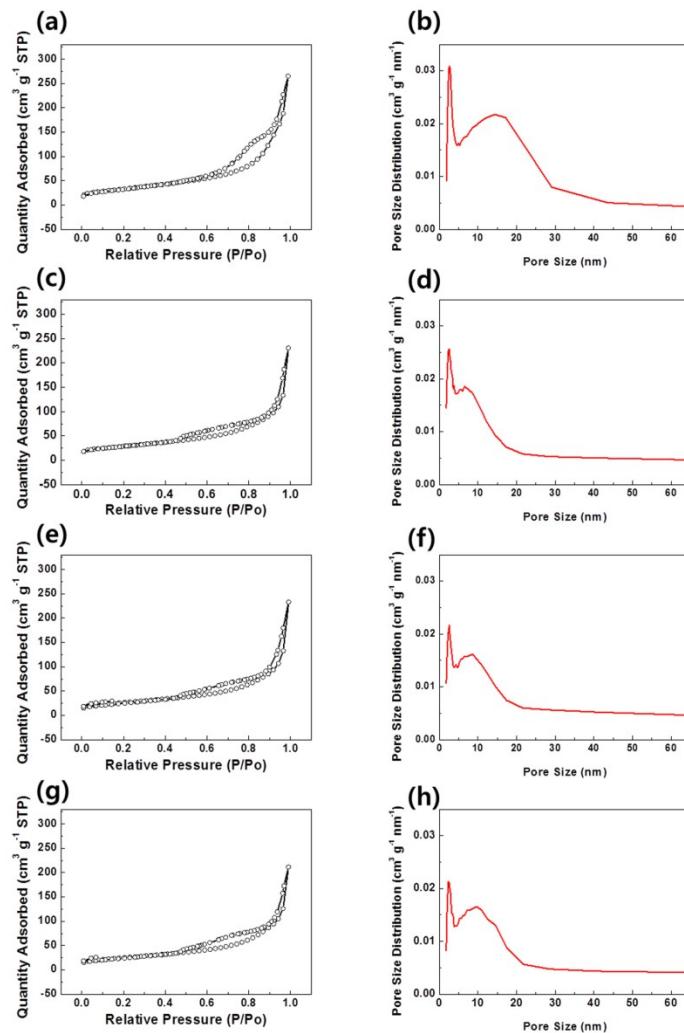


Figure S4. TGA curves of (a) MoS₂-only, (b) MoS₂/C-1, (c) MoS₂/C-3, and (d) MoS₂/C-5.

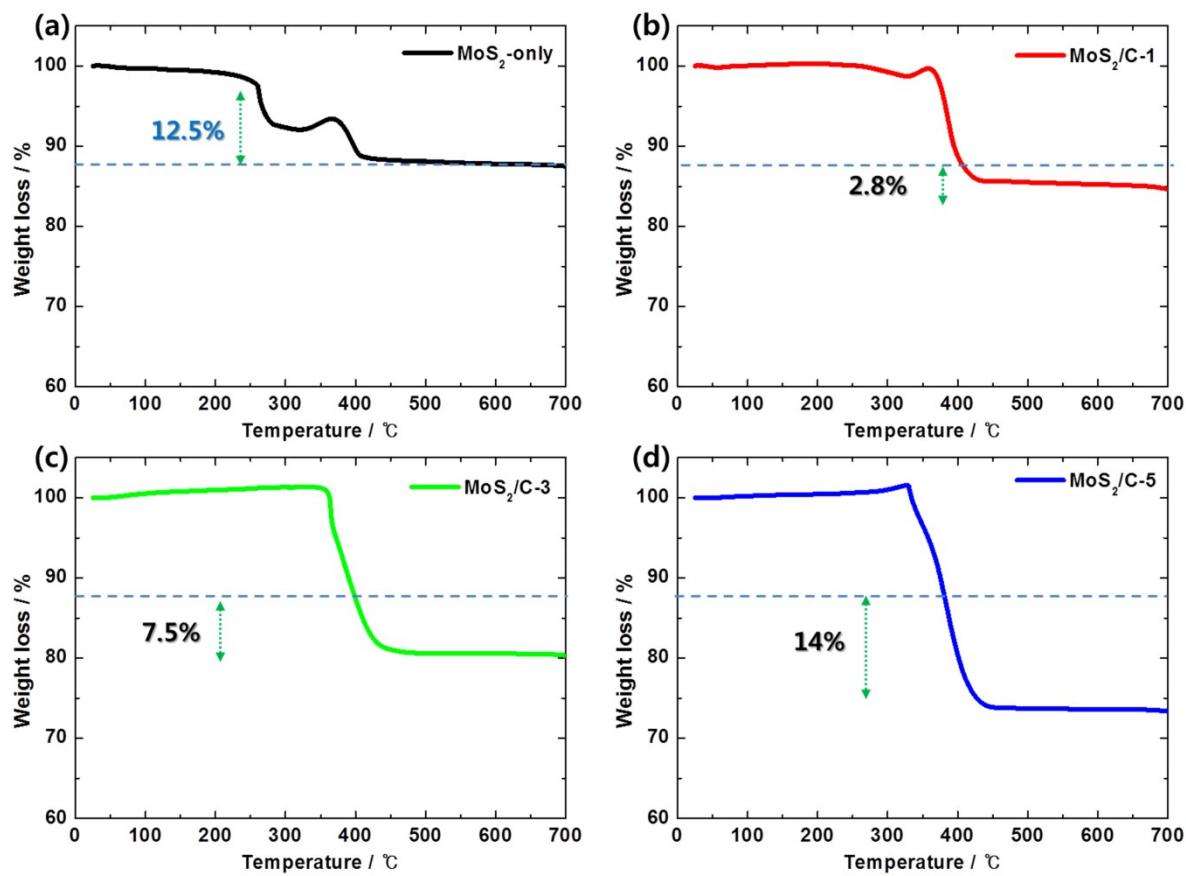


Figure S5. Comparison of electrical conductivity of the as-prepared samples.

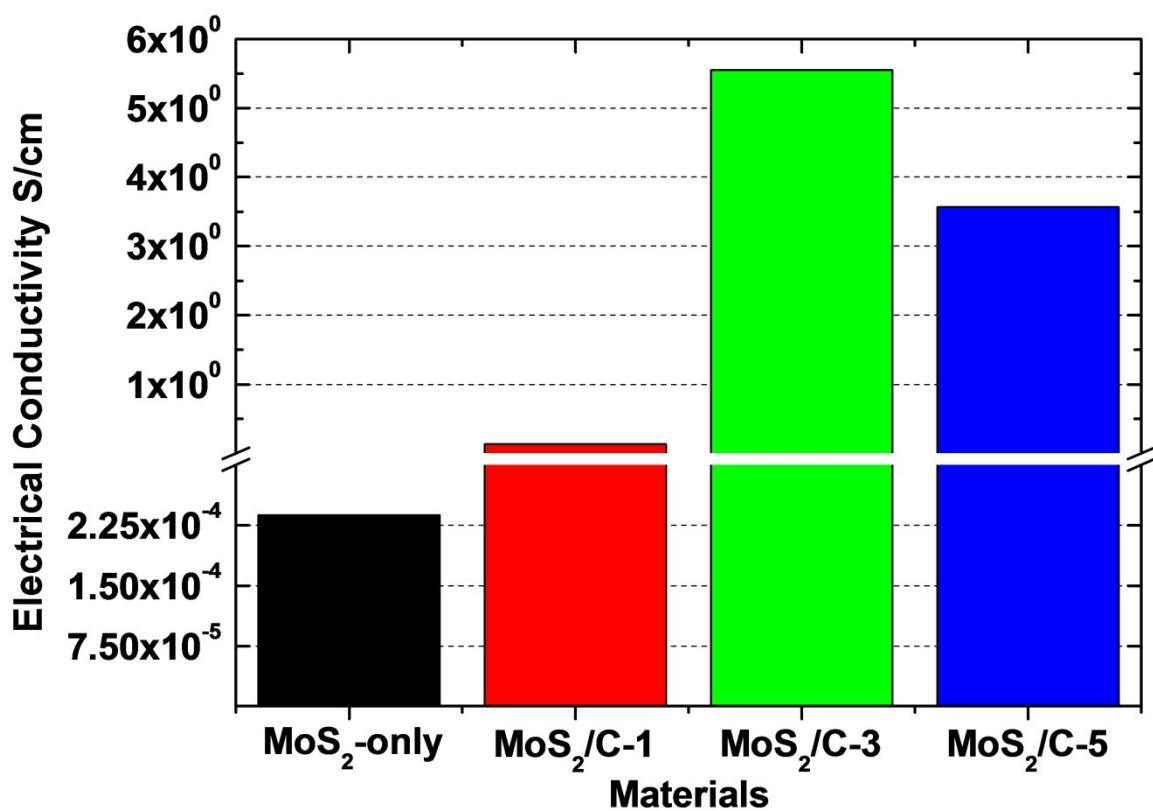


Figure S6. SEM images of (a) MoS₂-only, (c) MoS₂/C-1, (e) MoS₂/C-3, and (g) MoS₂/C-5 before the cycling process. (b) MoS₂-only, (d) MoS₂/C-1, (f) MoS₂/C-3, and (h) MoS₂/C-5 after 100 cycles at a current density of 500 mA g⁻¹.

