

Electronic Supplementary Information

Cylindrical nanostructured MoS₂ directly grown on CNT composites for lithium ion batteries

*HeeJoun Yoo^{†3}, Anand P. Tiwari^{†1}, JeongTaik Lee², Doyoung Kim³, JongHyeok Park² and Hyoyoung Lee^{*1}*

1. Department of Chemistry, Sungkyunkwan University, 300 Cheoncheon-Dong, Jangan-Gu, Suwon, Gyeonggi-Do 440-746, South Korea
E-mail: hyoyoung@skku.edu
2. School of Chemical Engineering, Sungkyunkwan University, Suwon 440-746, Republic of Korea
3. Department of Energy Science, Sungkyunkwan University, 2066 Seoburo, Jangan-gu, Suwon, Gyeonggi-do 440-746, Republic of Korea

† Equally contributed

* Corresponding Author: hyoyoung@skku.edu (Prof. Hyoyoung Lee)

In this Supplement, we present six figures that support results in the main text. The list is

Figure S1: X-ray photospectroscopy patterns of (a) MDGC-E and (b) MDGC-D.

Figure S2: Raman Spectra of (a) CNT (b) Sulfur coated CNT (c) MDGC-E and (d) MDGC- D samples.

Figure S3: TGA analysis of MDGC-E sample.

Figure S4: HRTEM of (a) MDGC-E and (b) MDGC-D samples.

Figure S5: Galvanostatic charge-discharge profile of MDGC-D.

Figure S6: Cycling performance of MDGC-D and corresponding charge discharge curve with different rate condition.

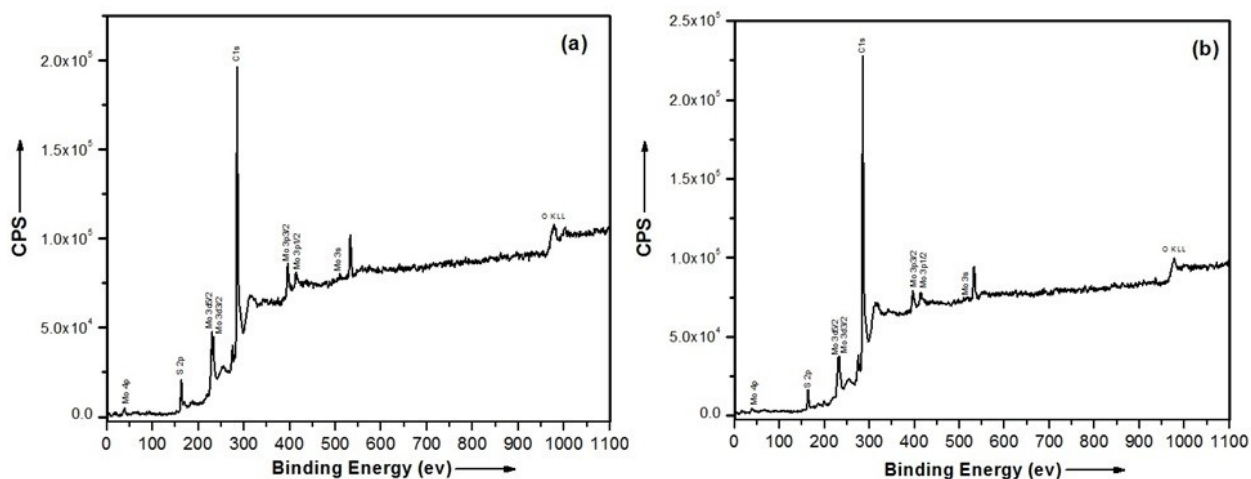


Figure S1: XPS patterns of (a) MDGC-E and (b) MDGC-D: It can be seen from Figures S1(a) and (b) that the sample contains the C, Mo and S elements. The calculated atomic ratio of Mo to S element is 1 to 2.14 and 1 to 2.20 for MDGC-E and MDGC-D samples, respectively, approaching the theoretical value of MoS_2 . The high-resolution of C1s can be seen apart from the “C = C” bond.

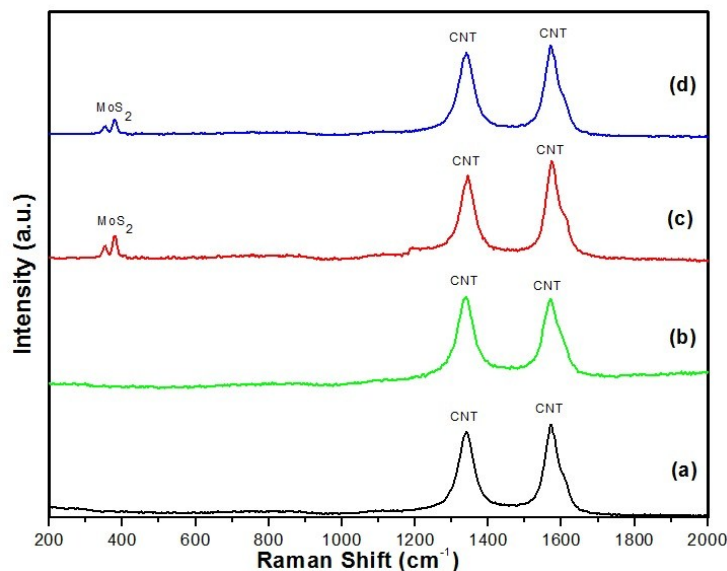


Figure S2: Raman Spectra of (a) CNT (b) Sulfur coated CNT (c) MDGC-E and (d) MDGC-D samples: It is evident from the Raman spectra S2(C) and S2 (d) that MoS_2 sheets have well

layered structure on the CNT surfaces. It is also illustrated that the D to G peak ratio of sulfur coated CNT increases with respect to bare CNT due to sulfur created some defects on the CNT surfaces.

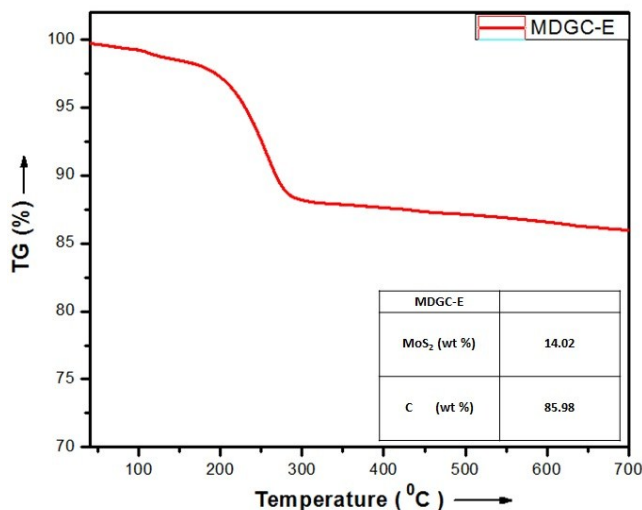


Figure S3: TGA curve of MDGC-E: The MDGC-E composite exhibits two weight losses. The first one appears at approximately between 230-250 °C, which can probably be attributed to the removal of oxygen-containing groups. The second is only one large continuous weight loss in the range of approximately 300-450 °C. This thermal behavior might be caused by the decomposition of the amorphous carbon and CNT, and oxidation of MoS₂ in the composites. The mass fraction of carbon and MoS₂ in the MDGC-E can be determined to be around 85.98 wt% and 14.02 wt%, respectively, assuming the complete conversion from MoS₂ to MoO₃.

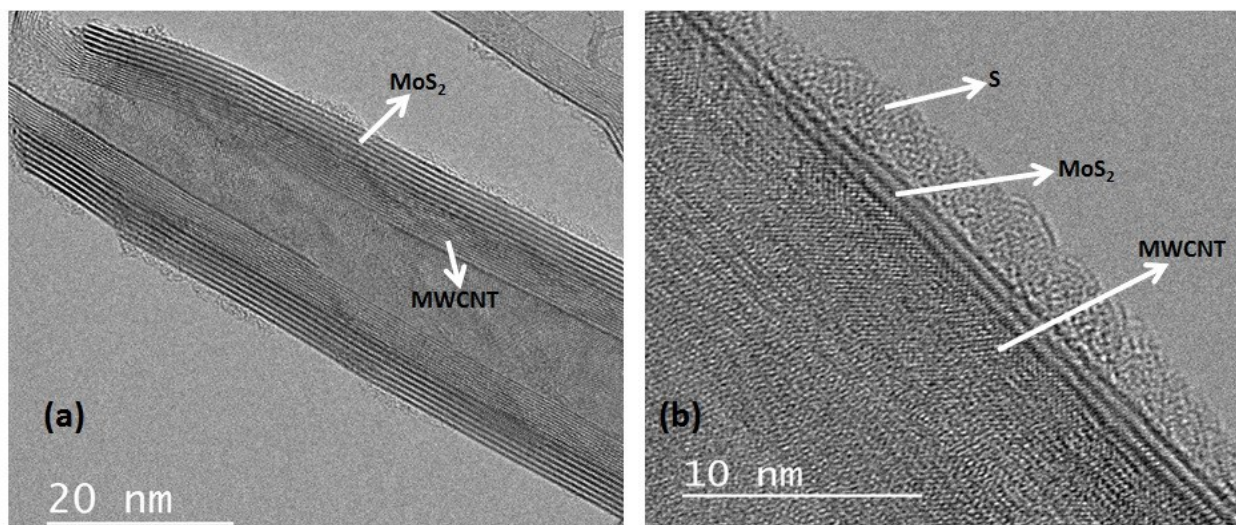


Figure S4: HRTEM of (a) MDGC-E and (b) MDGC-D samples. It can be clearly seen from Figures S4 (a) that MoS₂ sheets are well coated on the CNT surface. The d-values between the sheets are 0.62 nm which agrees with XRD analysis. It is illustrated from the Figures S4 (a) that the MDGC-E sample contains the 4-8 layers of MoS₂. The amorphous phase of residual S can be seen in the MDGC-D sample (Figure S4 (b)).

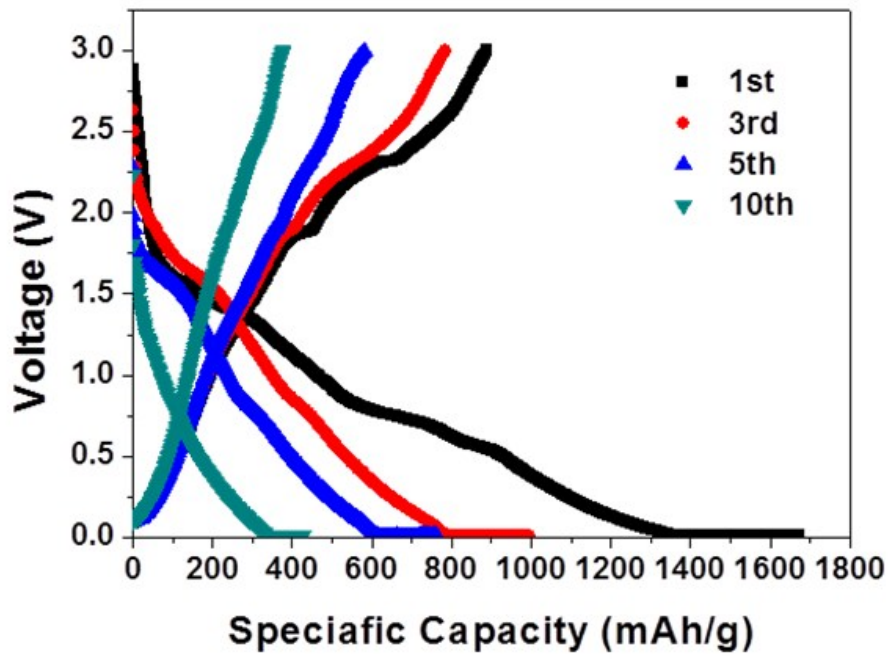


Figure S5 Galvanostatic charge-discharge profile of MDGC-D

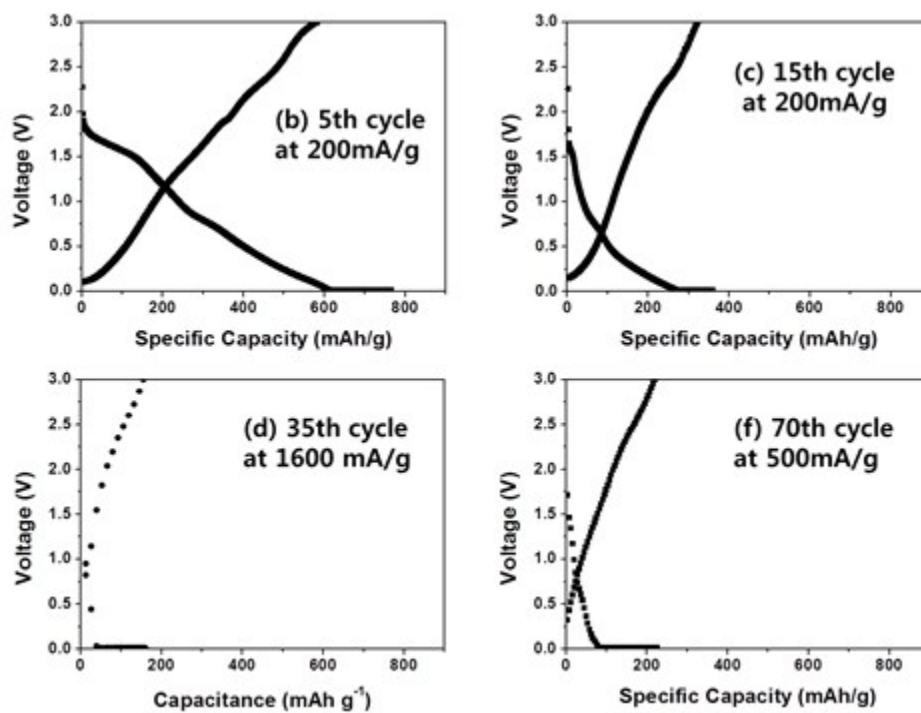
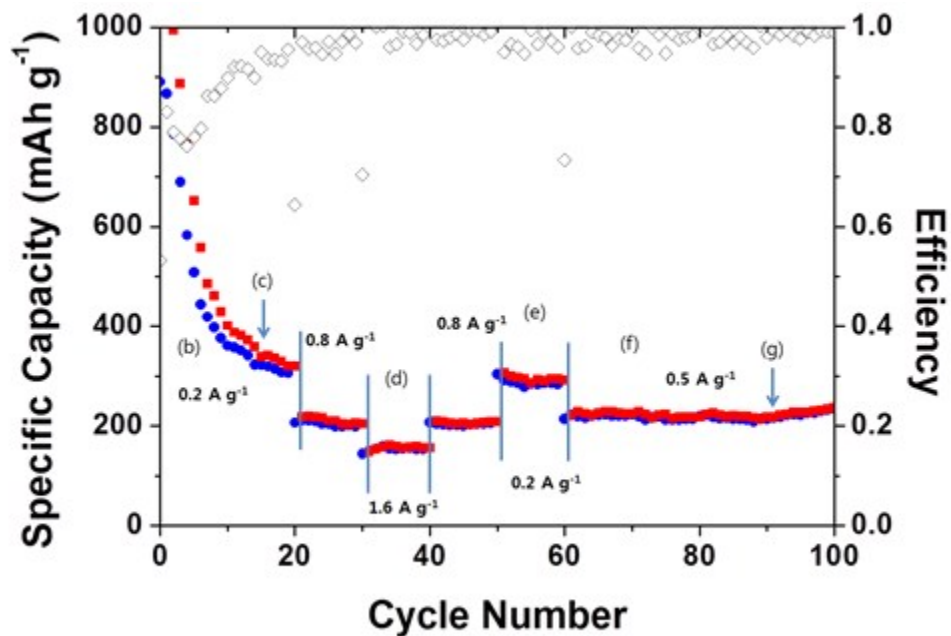


Figure S6 Cycling performance of MDGC-D and corresponding charge discharge curve with different rate condition.