

Electronic Supplementary Material

MoS₂/MnO₂ heterostructured nanodevices for electro-chemical energy storage

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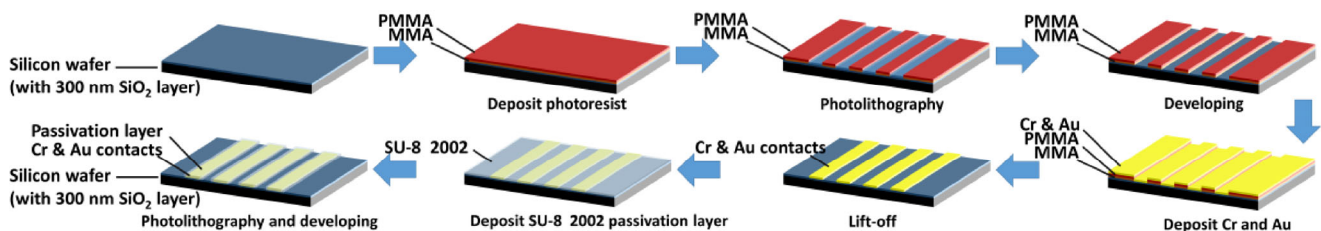


Figure S1 An illustration of the fabrication processes of the energy storage device based on individual heterostructure of MoS₂ nanosheet/MnO₂ nanowire. This process includes the following steps: Step 1, deposited MnO₂ nanowires via spin-coating method on the prepared silicon wafers (with 300 nm SiO₂ layer), and MoS₂ nanosheets obtained through mechanical exfoliation were transferred immediately to the silicon wafer that with MnO₂ nanowires. Then, MMA and PMMA photoresists were coated on silicon wafers with 300 nm SiO₂ layer; Step 2, patterned the metal contacts pad with EBL on silicon wafers; Step 3, developing and rinsing; Step 4, deposited 5 nm Cr and 150 nm Au on to the silicon wafers with thermal evaporation; Step 5, lift-off; Step 6, coated SU-8 2002 on silicon wafers; Step 7, patterned SU-8 2002 with EBL to form the passivation layer that covered the metal contacts, which avoided leakage current.

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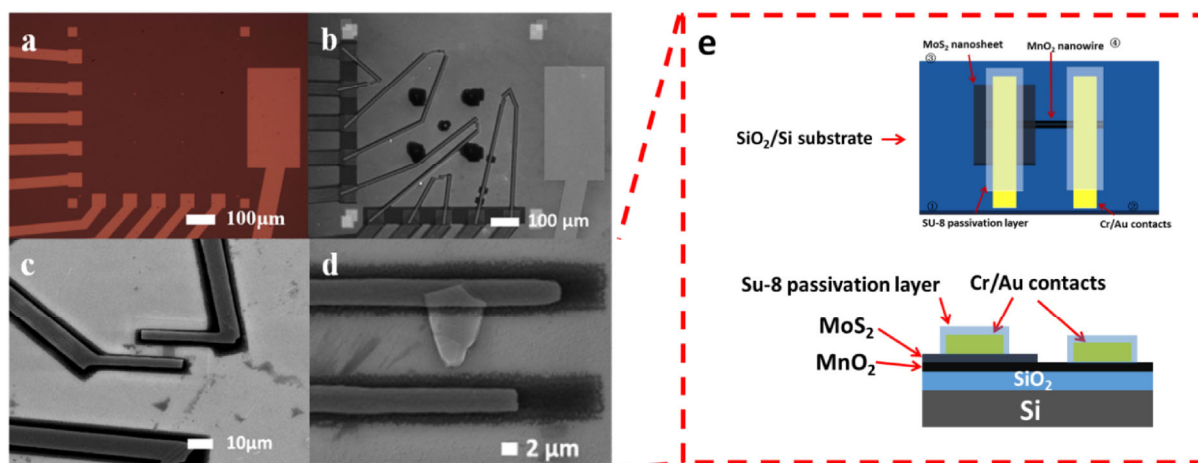


Figure S2 (a) The optical image of the outer electrodes; (b) SEM image of the entailed outer electrodes and inner electrodes. The inner electrodes are covered by SU-8 passivation layer to avoid the leakage current; (c) The selected multilayer MoS₂ nanosheet contacts with inner electrodes; (d) SEM image of MoS₂/MnO₂ heterostructure; (e) The illustration of the energy storage device based on individual heterostructure of MoS₂ nanosheet/MnO₂ nanowire in overhead view and front view.

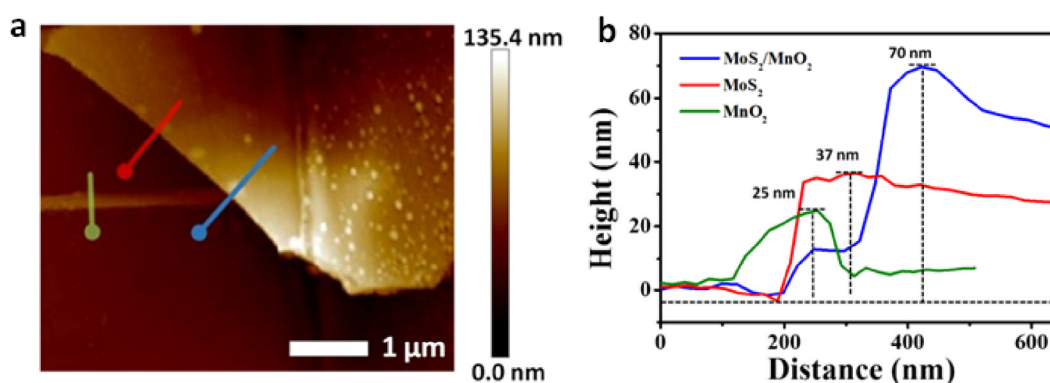


Figure S3 (a) AFM image of MoS₂/MnO₂ heterostructure; (b) The height profile of MoS₂/MnO₂ heterostructure, few-layer MoS₂, and MnO₂ nanowire.

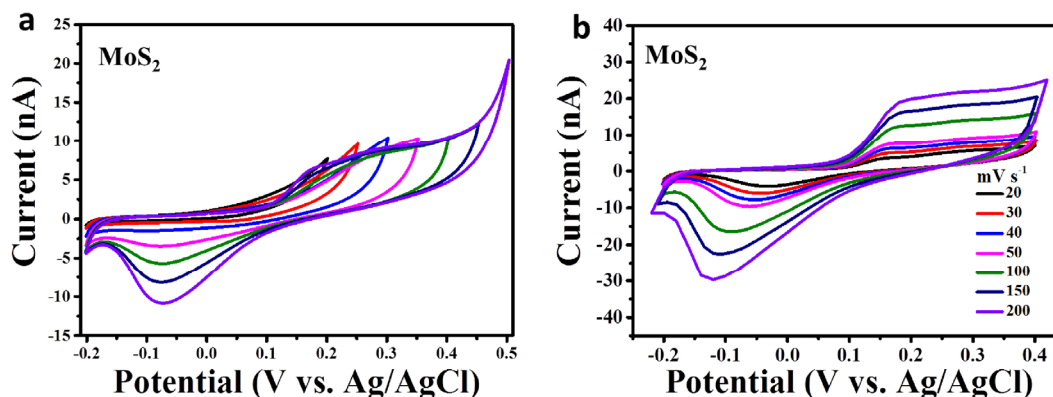


Figure S4 (a) CV curves of multilayer MoS₂ nanosheet at a scan rate of 50 mV s⁻¹. And the CV curves of the electrode materials are tested at different voltage range; (b) CV curves of multilayer MoS₂ nanosheet at different scan rates.

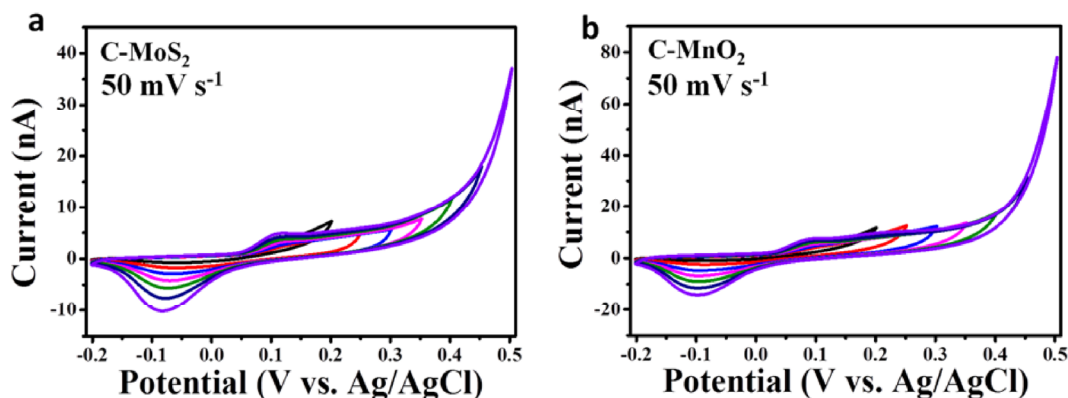


Figure S5 (a) CV curves of MoS₂/MnO₂ heterostructure under C-MoS₂ mode tested at different voltage ranges at a scan rate of 50 mV s⁻¹; (b) CV curves of MoS₂/MnO₂ heterostructure under C-MnO₂ mode tested at different voltages range at a scan rate of 50 mV s⁻¹.

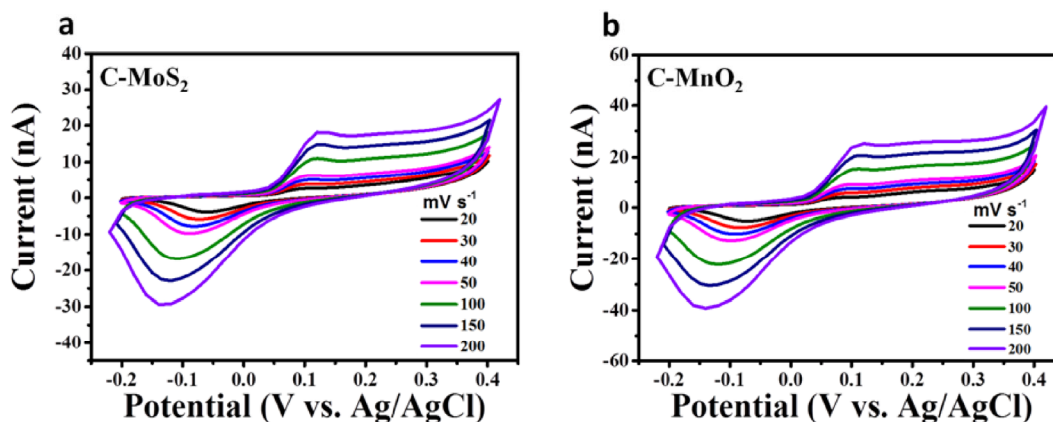


Figure S6 (a) CV curves of MoS₂/MnO₂ heterostructure under C-MoS₂ mode tested at different scan rates; (b) CV curves of MoS₂/MnO₂ heterostructure under C-MnO₂ mode tested at different scan rates.

The electrochemical performances of the energy storage devices were investigated by combining the probe station with the electrochemical workstation. The CV curves of the multilayer MoS₂ nanosheet and MoS₂/MnO₂ heterostructure energy storage devices with different scan rates and voltage window are shown in Figure S4-6 respectively. It can be found that the redox peaks of the MoS₂ appear at 0.15 and -0.04 V respectively. Meanwhile, there is obvious hydrogen evolution reaction in both MoS₂ nanosheet and MoS₂/MnO₂ heterostructure when expanding the voltage window from -0.2 – 0.4 V to -0.2 – 0.5 V. (Figure S4, S5) Thus, a suitable electrochemical voltage window of -0.2 – 0.4 V is chosen for the further measurement.

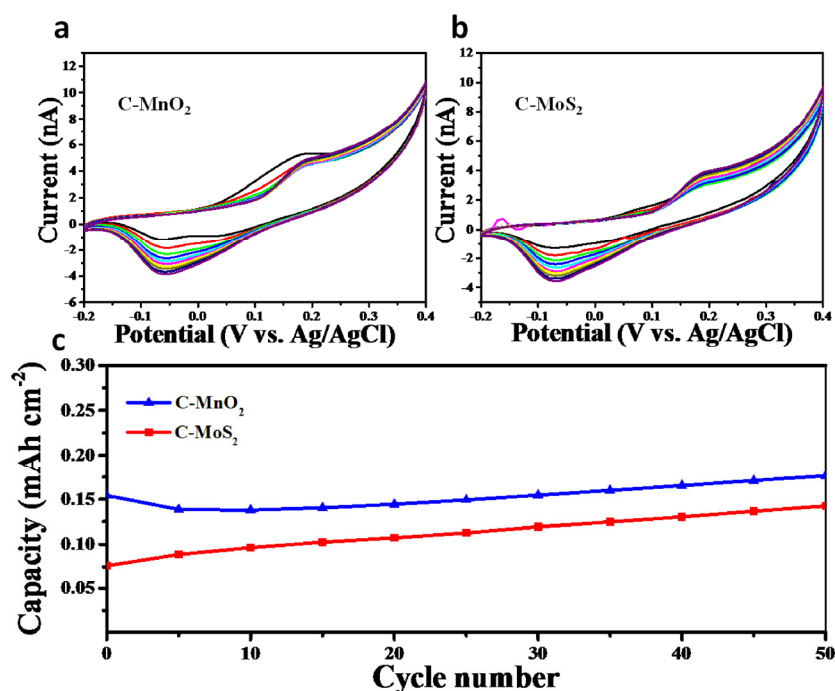


Figure S7 (a) CV curves of MoS₂/MnO₂ heterostructure under C-MnO₂ mode tested at a scan rate of 100 mV s⁻¹; (b) CV curves of MoS₂/MnO₂ heterostructure under C-MoS₂ mode tested at a scan rate of 100 mV s⁻¹; (c) Cycling performances of the MoS₂/MnO₂ heterostructure under C-MnO₂ and C-MoS₂ modes.

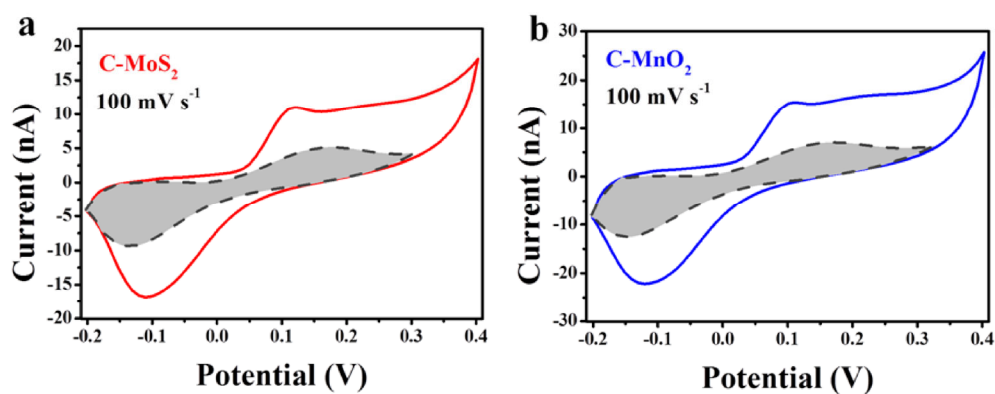


Figure S8 (a) The voltage profile for the surface capacity current (shaded region) is compared with the total measured current obtained at a scan rate of 100 mV s⁻¹ for the MoS₂/MnO₂ heterostructure under C-MoS₂ mode; (b) The voltage profile for the surface capacity current (shaded region) compared with total measured current obtained at a scan rate of 100 mV s⁻¹ for the MoS₂/MnO₂ heterostructure under C-MnO₂ mode.