VALUE ADDED ABSTRACT

Self-standing MoS2/CNT and MnO2/CNT one dimensional core shell heterostructures for asymmetric supercapacitors application

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well as highly conductive current collector. For positive electrode, MnO2 robust nature, with high coulombic efficiency of 92%. (manganese oxide) nanoparticles loaded CNTs have been fabricated, owing to the work function contrast between MoS2 and MnO2, in order to utilize the maximum potential window in our asymmetric design. Both the electrodes, developed using a unique combination of chemical

Recently, two-dimensional layered nanostructures, specially MoS₂ and physical deposition techniques, form hierarchical core-shell (molybdenum disulfide) has come out as the most investigated electrode heterostructure consisting of large number of edge-exposed catalytic sites material for batteries and supercapacitors, possessing well preserved in- available for electrode-electrolyte interaction. The electrodes were first plane covalent bonding, leading to extraordinary mechanical elasticity tested in a three-electrode configuration, found to have very high areal within the layers as well as outstanding firmness along the c-axis. We capacitance of 0.41 and 0.6 F/cm² and at a scan rate of 10 mV/sec in 1M have synthesized vertically aligned edge exposed molybdenum disulfide Na₂SO₄ system for MoS₂-CNT and MnO₂-CNT, respectively. MoS₂-CNT nanoflakes onto free standing carbon nanotubes sheet as an efficient electrode depicted purely electrostatic polarization in the voltage range of negative electrode material. Functionalization of carbon nanotubes with 0.6 to 0.2 V, whereas MnO2-CNT electrode also displayed non-faradaic metal oxides or chalcogenides is rather a complicated practice due to the charge storage in the range of 0 to 1 V. Next, the device was fabricated and superhydrophobic nature of these films. The inherit hydrophilicity and tested in a wide potential range of 0.8 to 1 V, with the calculated areal controlled chirality of carbon nanotubes makes them an ideal candidate capacitance and volumetric capacitance of 820 mF/cm² and 4.2 F/cm², for heterostructured electrode material synthesis. Moreover, their self- respectively, at the scan rate of 10 mV/s. Moreover, the device showed a standing nature allows them to function both as the active material, as capacitance retention of ~97.2 % in 2000 cycles, displaying extremely

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