

Supporting Information

Synthesis and Electrochemistry of Highly Pseudocapacitive Carbon Nano Onions aka Multilayer Fullerenes and its MnO₂ nanocomposite

Vedi kuyil Azagan M^a, Mukta V. Vaishampayan^a, Manjusha V. Shelke^{a,b,c*}

^aPhysical and Materials Chemistry Division, CSIR-National Chemical Laboratory, Pune-411008, MH, India.

^bCSIR-Network Institute for Solar Energy, CSIR-National Chemical Laboratory, Pune-411008, MH, India.

^cAcademy of Scientific and Innovative Research (AcSIR), Anusandhan Bhawan, 2 Rafi Marg, New Delhi-110 001

1. Comparative CV of CNOs, MnO₂/CNOs and carbon paper:

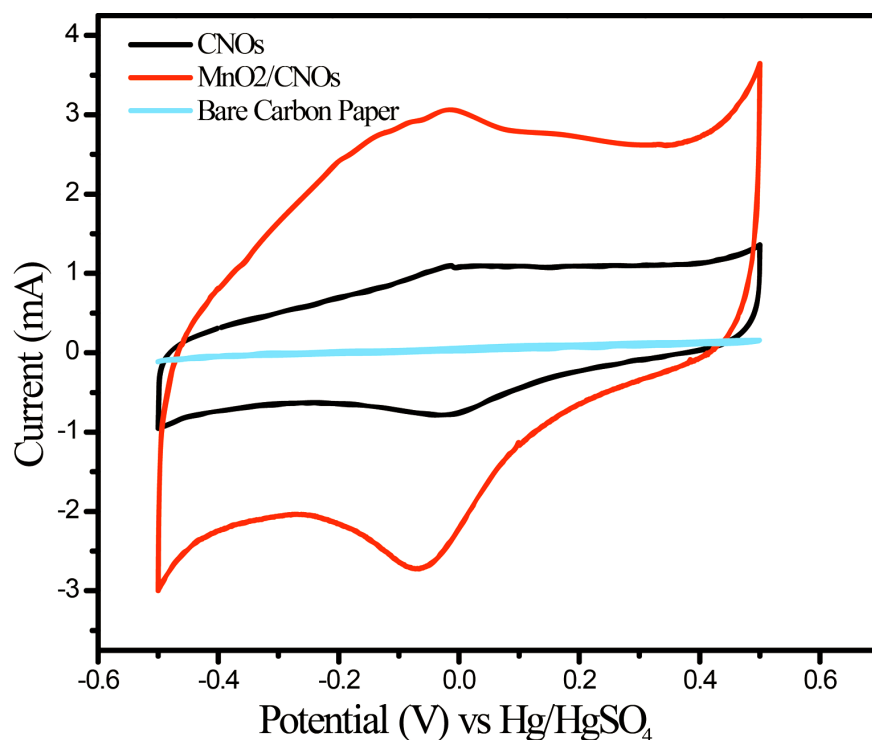


Figure S1. Comparative cyclic voltamogram shows the contribution of bare carbon paper which was used as the current collector at the scan rate of 5mV/s.

2. Surface area analysis:

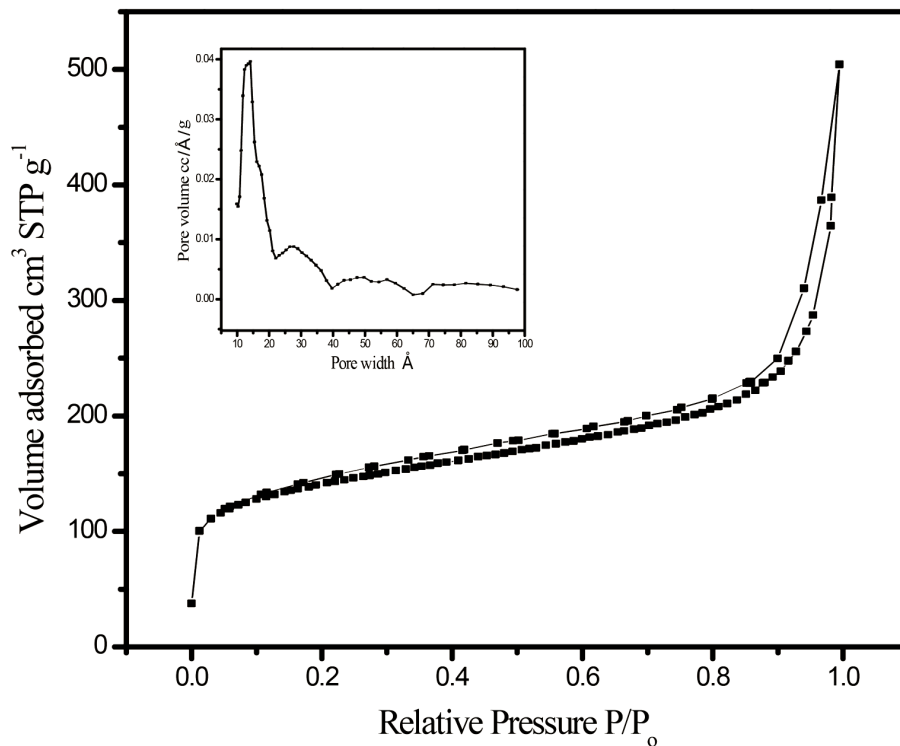


Figure S2. Nitrogen sorption isotherms at 77 K and the Inset shows pore size distributions of CNOs.

By using Brunauer-Emmett-Teller (BET) theory and Density Functional Theory (DFT) on the collected nitrogen-adsorption isotherms, pore size and the surface area distributions were calculated. For N₂-sorption measurements, Autosorb-iQ automatic volumetric instrument was performed at 77 with pressures in the range 0–760 Torr. The calculated average pore size of CNOs chain structure is ranging from 1.2-1.4 nm. CNOs exhibit VI isotherm and it has surface area of 486m²g⁻¹.

3. HRTEM images of CNOs:

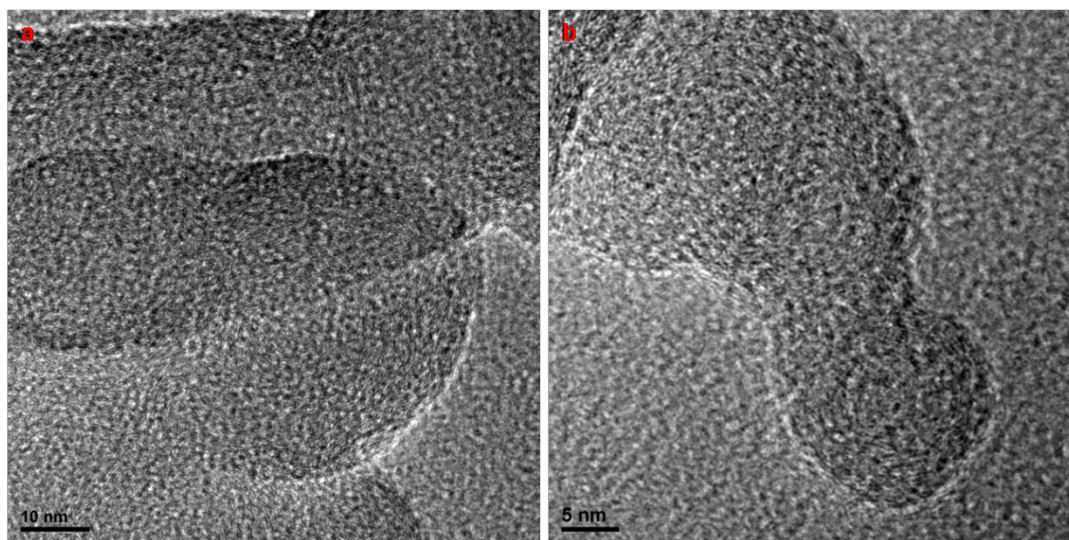


Figure S3. HRTEM images of CNOs showing (a) completed ring of multilayer fullerene and (b) fused fullerene rings

4. TEM images of thermal carbon black:

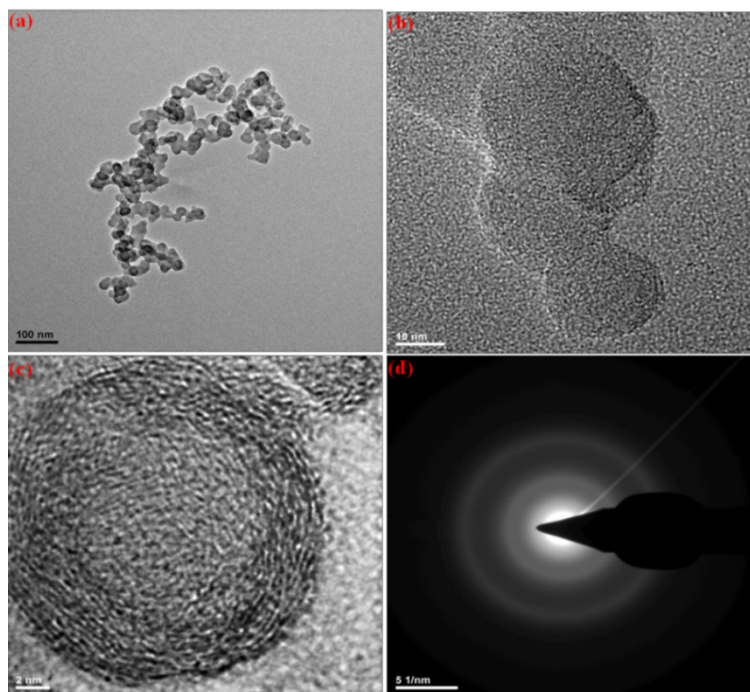


Figure S4. (a,b,c) Shows the HRTEM images of as collected thermal carbon black (d) shows the SAED pattern of the CNO, the inter atomic distance calculated is 0.34nm; corresponding to (002) plane of carbon.

5. Thermal gravimetric analysis (TGA):

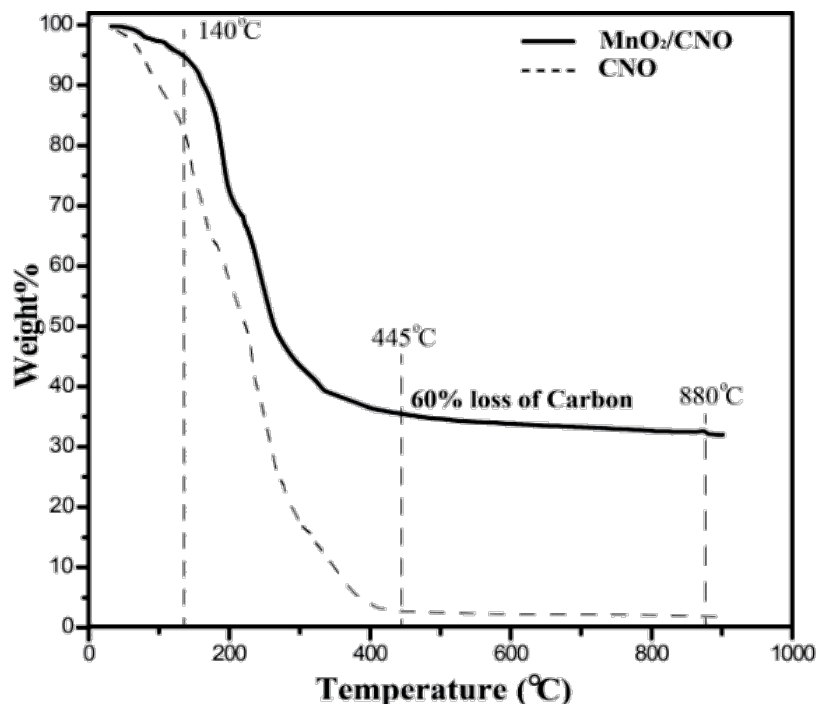


Figure S5. Thermogravimetric analysis (TGA) of CNOs and its composite MnO₂/CNOs before annealing at 800°C.

For MnO₂/CNOs composite 5% weight loss has been observed between 30 and 140 °C that is attributed to the liberation of adsorbed water molecules from the composites. Low onset potential shows pyrolysis of organic residue. Further weight loss of 60% up to 445 °C corresponds to the loss of organic residues in CNOs. In addition, we have observed a final weight loss of 1% after 880 °C is due to the conversion of MnO₂ to Mn₃O₄[Xiaofeng Xie, Lian Gao, 'Characterization of a manganese dioxide/carbon nanotubes composite fabricated using an in situ coating method', *Carbon*, 45, (2007) 2365–2373.]

6. XPS of MnO₂/CNOs composite:

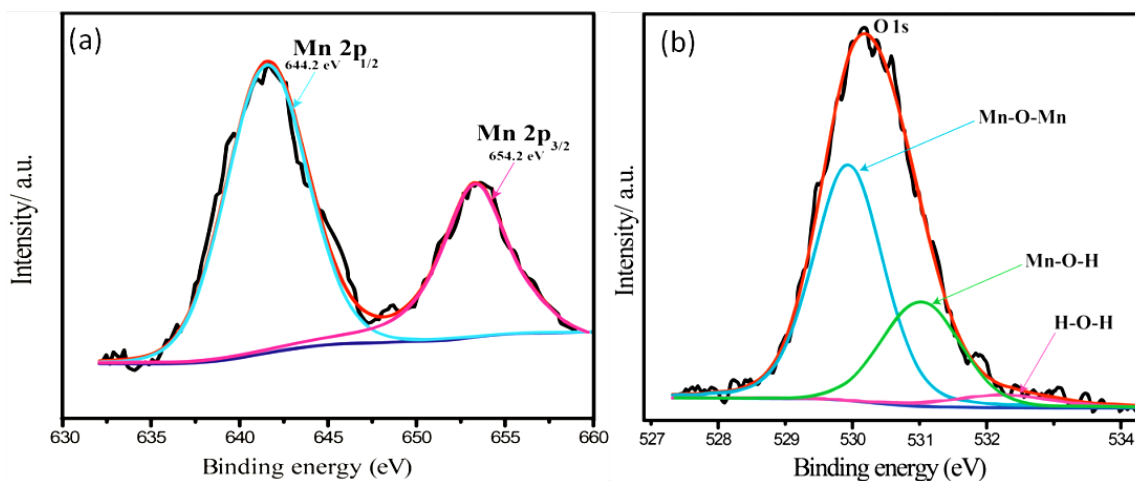


Figure S6. XPS spectra of MnO₂/CNOs composite.

The XPS measurements have been taken to study the components and oxidation states of CNOs and MnO₂/CNOs composite. In figure (a) the binding energies 644.2 eV and 654.2 eV corresponding to Mn 2p band for the binding energies 2p_{1/2} and 2p_{3/2} which proves that the Mn in the composite is having oxidation state IV. The XPS of O 1s has been deconvoluted into three components (fig b); the binding energy at 529.86 eV gives the evidence for Mn-O-Mn bonds for the tetravalent oxide which has highest intensity among others and the another binding energy at 531.19 eV shows that the hydroxide group has been bound to manganese Mn-OH; the binding energy at 532.48 eV corresponds to H-O-H bonds.

7. SEM images of precursor ghee, thermal carbon black and CNOs:

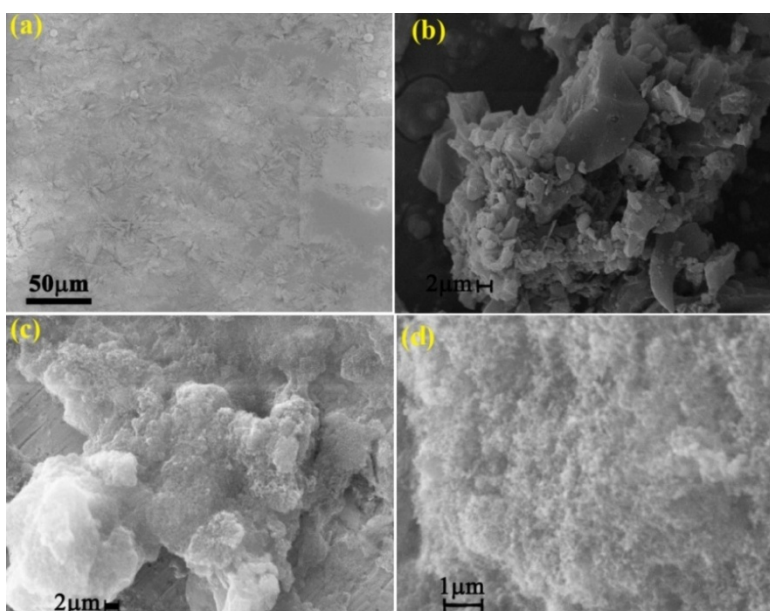


Figure S7. SEM images of (a) ghee (butter oil), (b) as collected thermal black carbon and (c), (d) CNOs after annealing at 800°C.