

Supporting information

Fabrication and Electrochemical Evaluation of Flexible Spinel CdMn₂O₄ Carbon Nanofibers for Advanced Supercapacitor Applications

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Electrochemical Performance Measurements

The electrochemical studies of the spinel CdMn_2O_4 CNFs were performed using an electrochemical workstation (CHI660E) in a 6M KOH and 1M H_2SO_4 aqueous electrolyte solution. The experiment involved utilizing a conventional three-electrode setup, with the CdMn_2O_4 CNFs serving as the working electrode, Ag/AgCl with 3.0 M KCl as the reference electrode, and platinum wire as the counter electrode. Cyclic Voltammetry (CV) was performed at different scan rates, galvanostatic charge-discharge (GCD) was performed at various current densities, and EIS was conducted in the frequency range of 100 kHz to 0.1 Hz with an AC amplitude. Two methods are employed to determine the specific capacitance in a three-electrode configuration: CV and GCD.

For CV, the specific capacitance (C_s) is computed using equation (S1):¹

$$C_s = \frac{1}{mv(E_2 - E_1)} \int_{E_1}^{E_2} i(E) dE \#(S1)$$

Where the difference between E_2 and E_1 represents the potential window. The variable m represents the mass of the active synthesized material, v denotes the scan rate, and $i(E)$ represents the current at each potential.

For GCD, the specific capacitance (C_s) is determined using equation (S2):

$$C_s = \frac{I \times \Delta t}{m \times \Delta V} \#(S2)$$

Where C_s represents the specific capacitance in ($\text{F} \cdot \text{g}^{-1}$), I stands for the current in (A), Δt denotes the time interval in seconds for discharge, ΔV and m represents the voltage window (V) and mass of active material in (g).

Two-electrode systems evaluated the properties of a symmetrical solid-state supercapacitor. The specific capacitances of a single electrode were calculated based on the GCD outcomes using the following equation (S3):²

$$C_s = \frac{4 \times I \times \Delta t}{m \times \Delta V} \#(S3)$$

The energy density (E , Wh kg^{-1}) and power density (P , W kg^{-1}) of the fabricated SSC device were calculated by the following formulas:

$$E = \frac{1}{2} C_s V^2 \times \frac{1}{3600 \times m} \#(S4)$$

$$P = \frac{E \times 3600}{t_{discharge}} \#(S5)$$

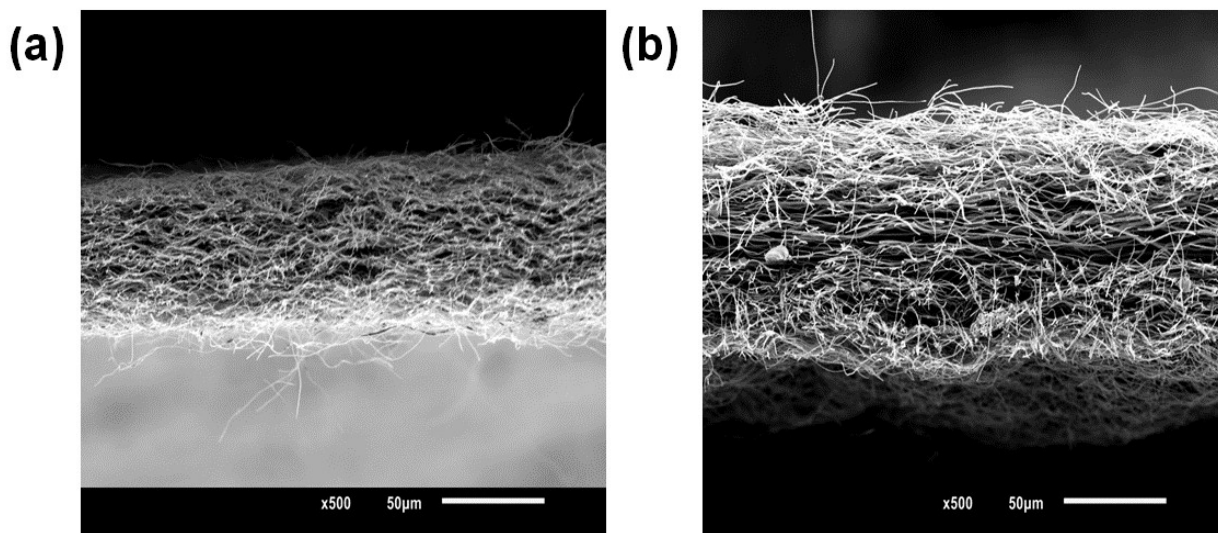


Figure S1. Cross-sectional SEM image of (a) CMO-1 and (b) CMO-2

The thickness of CMO-1 is approximately 67 μm, whereas CMO-2 has a thickness of around 88 μm.

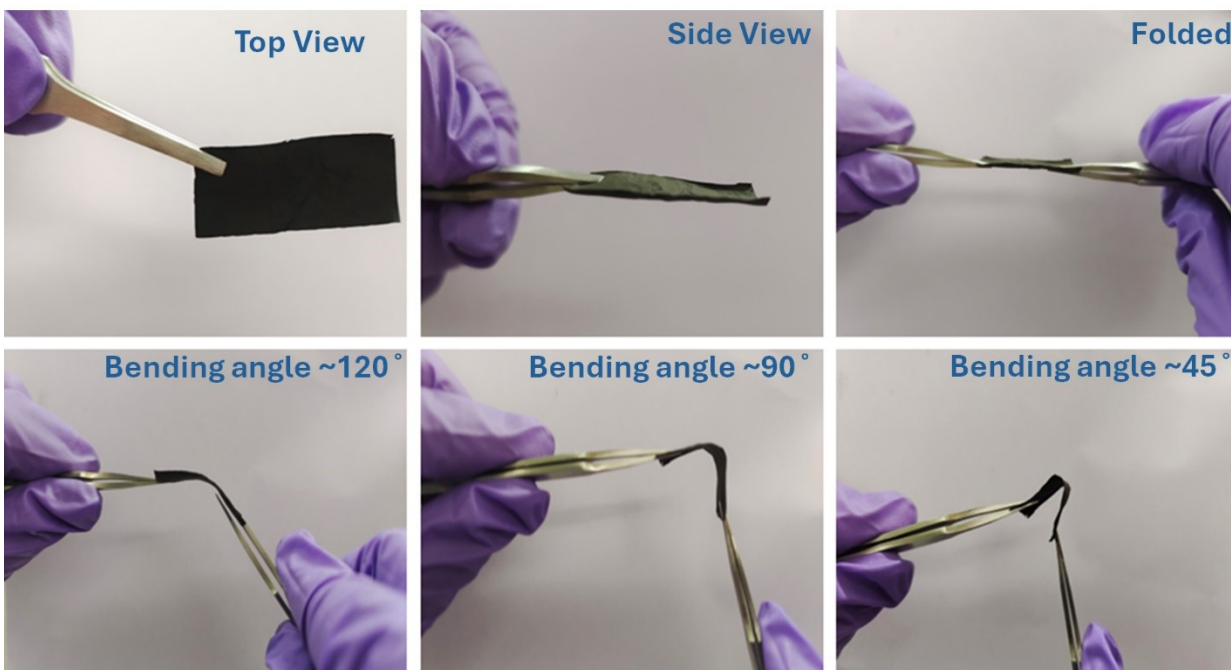
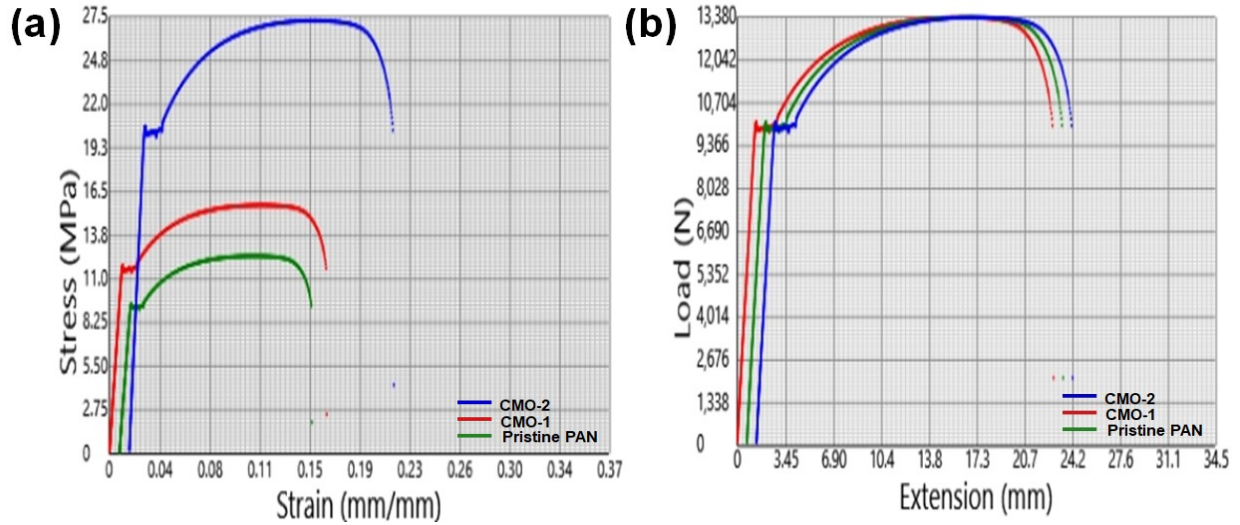


Figure S2. Photographic Images of spinel CdMn_2O_4 carbon nanofibers(CNFs)

Figure S2 shows images of the spinel CdMn_2O_4 CNFs at different angles, demonstrating the thickness and flexibility of the material. This visual representation provides valuable insights into the physical characteristics of the flexible material, showcasing its structural properties from



various perspectives.

Figure S3. Stress-strain and load-extension curves for Pristine PAN, CMO-1, and CMO-2 CNFs.

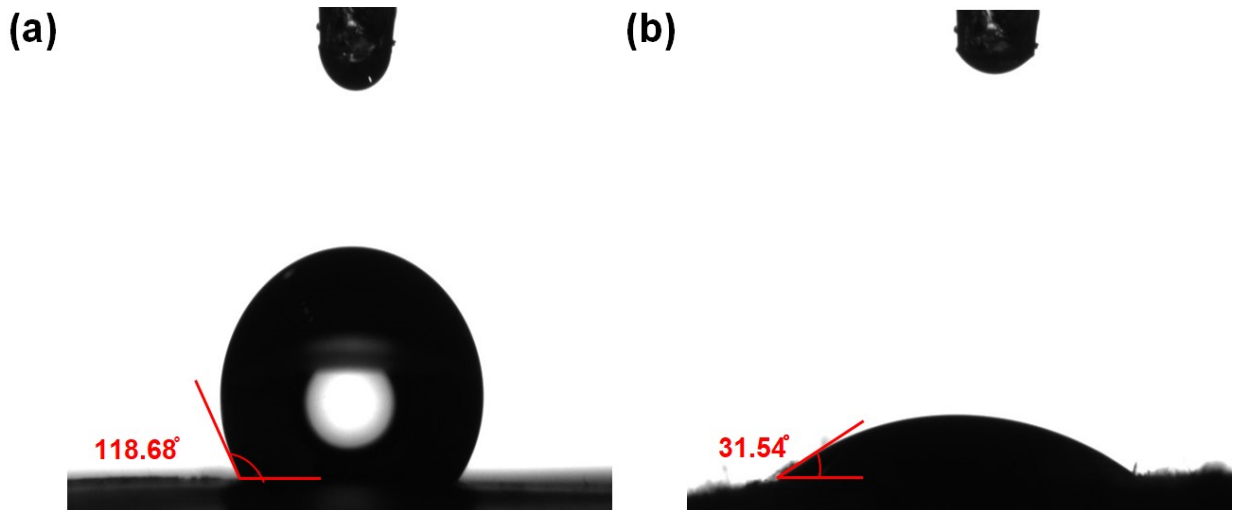


Figure S4. Contact angle of a water droplet on the surface of (a) Pristine PAN carbon nanofibers and (b) CdMn_2O_4 carbon nanofiber

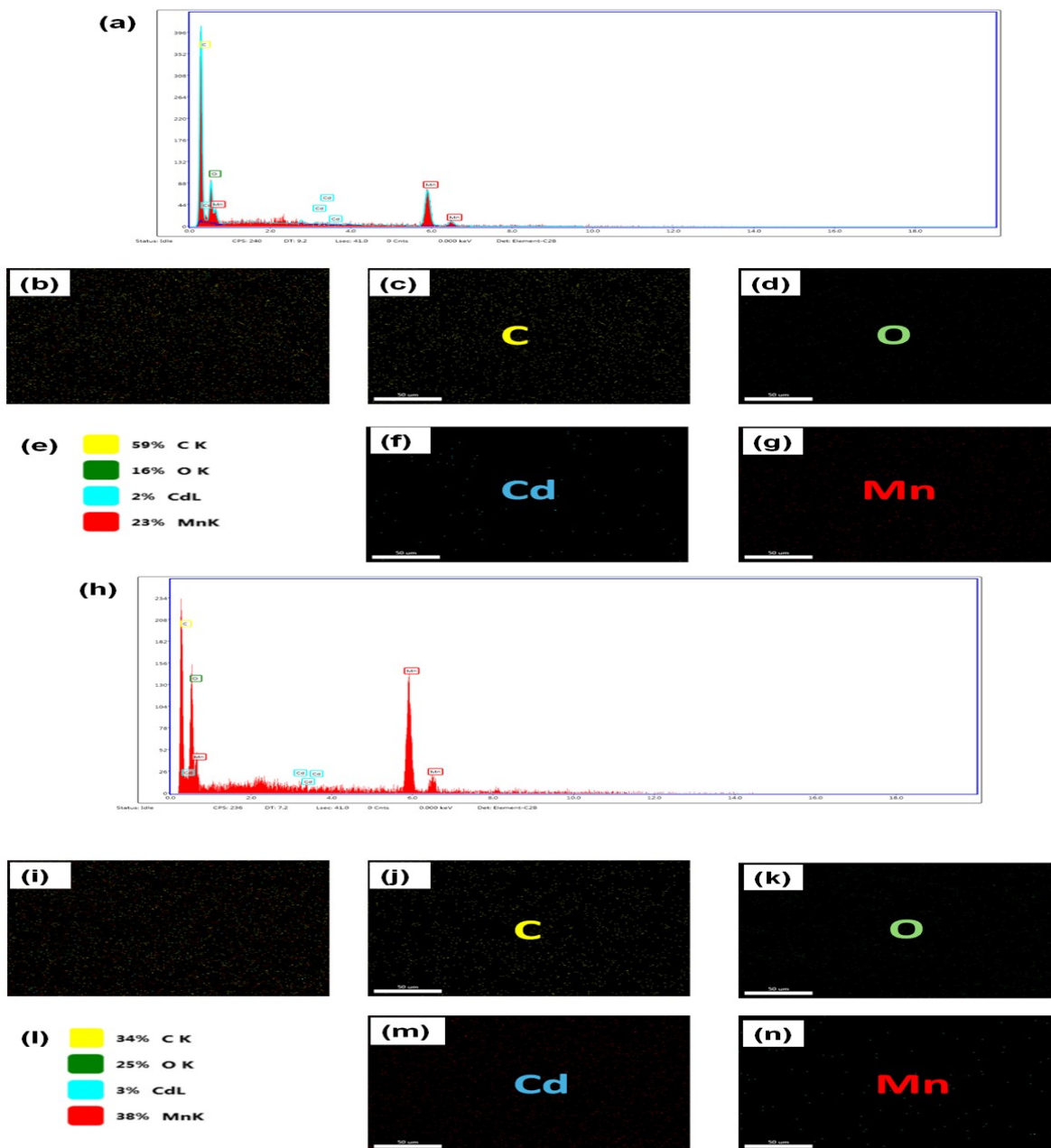


Figure S5. (a-g) presents the EDAX analysis for CMO-1, including (e) the EDAX spectrum showing the percentage of chemical elements present. (c-g) display the elemental mapping images for the individual distribution of C, O, Cd, and Mn, respectively. (h-n) presents the EDAX analysis for CMO-2, including (l) the EDAX spectrum showing the percentage of chemical elements present.

(j-n) display the elemental mapping images for the individual distribution of C, O, Cd, and Mn, respectively.

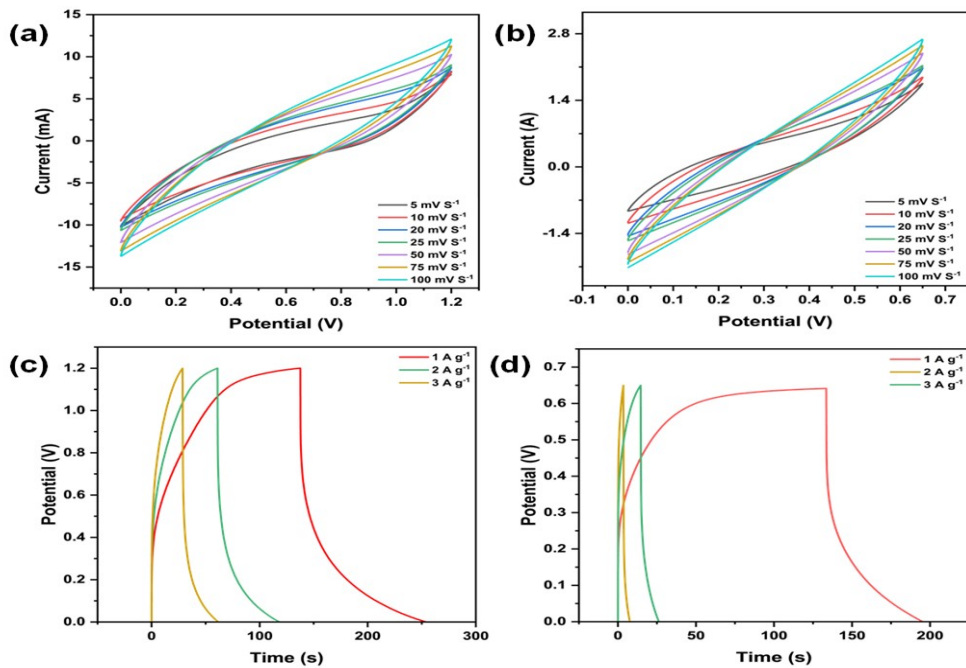


Figure S6. CV curves at varying scan rates (a, b) and GCD curves at different current densities (c, d) of Pure MnO₂

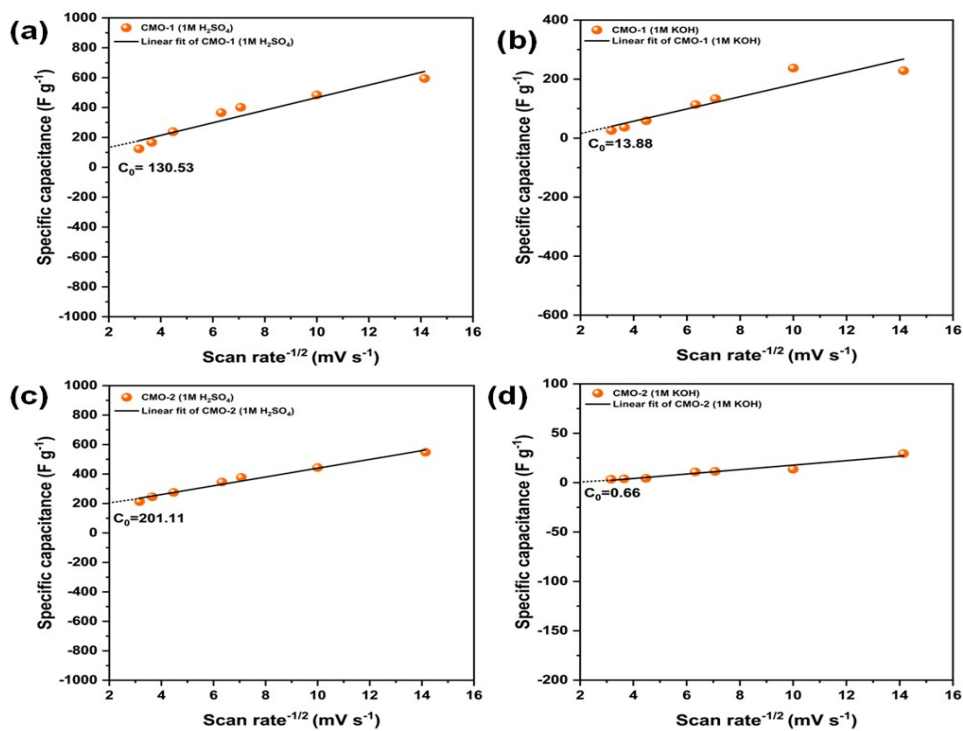


Figure S7. Specific capacitance vs. scan rate^{-1/2} for (a-b) CMO-1 and (c-d) CMO-2 in 1M H₂SO₄ and 1M KOH, with linear fits showing C₀ values

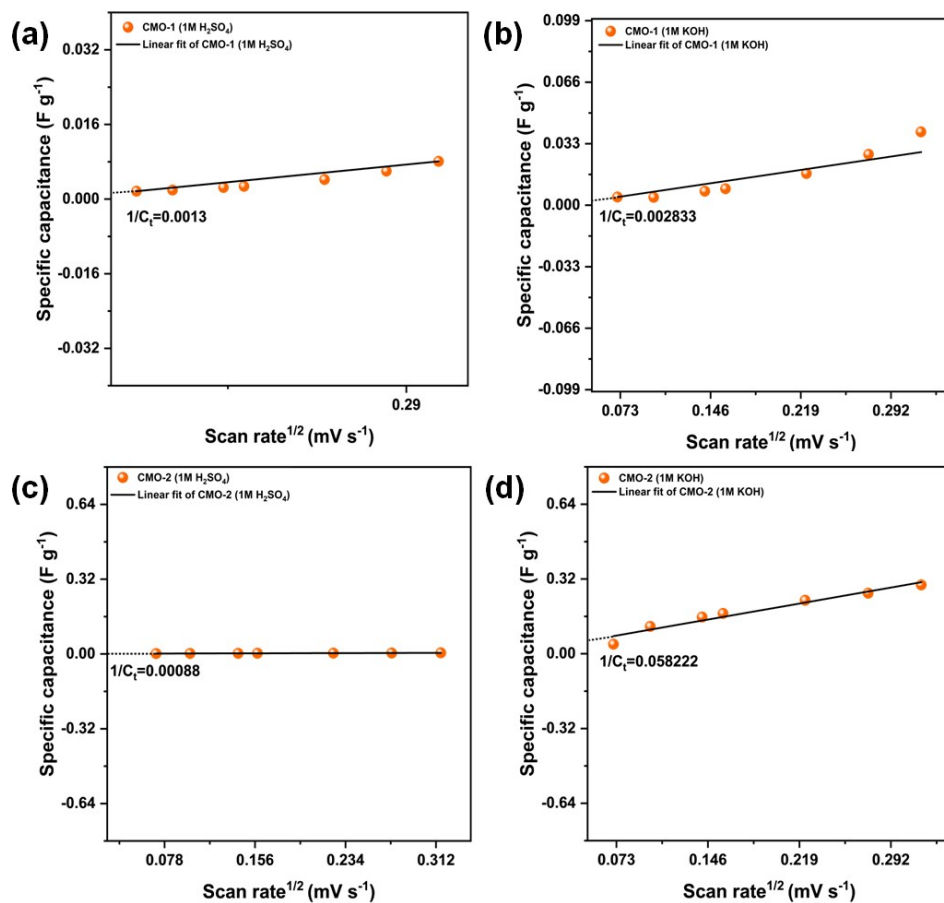


Figure S8. Specific capacitance vs. scan rate^{1/2} for (a-b) CMO-1 and (c-d) CMO-2 in 1M H₂SO₄ and 1M KOH, with linear fits showing C_t values.

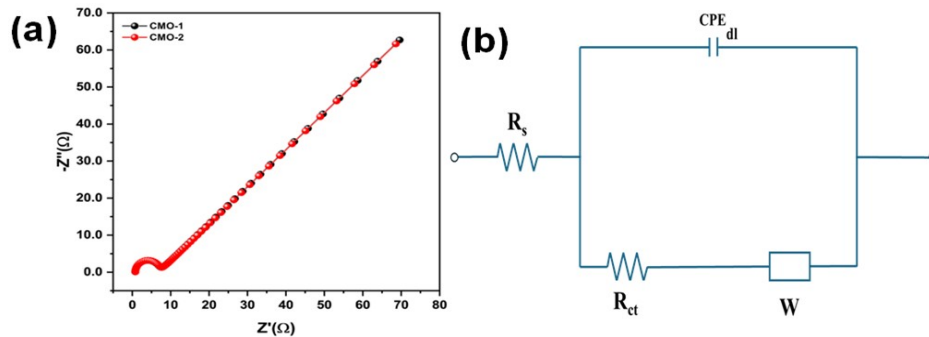


Figure S9. Nyquist plots of CMO-1 and CMO-2 in (a) 1M H₂SO₄ and (b) 1M KOH using a three-electrode setup, showing impedance behavior of the materials. (c) Fitted equivalent circuit

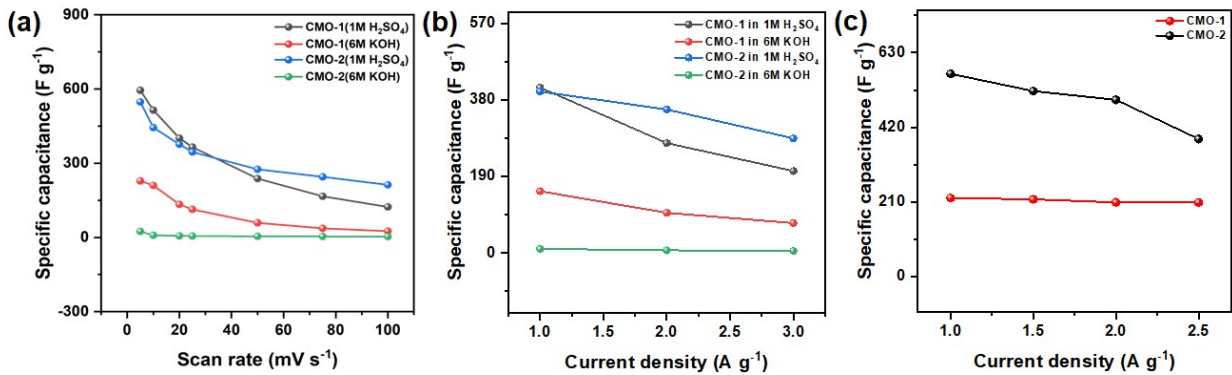
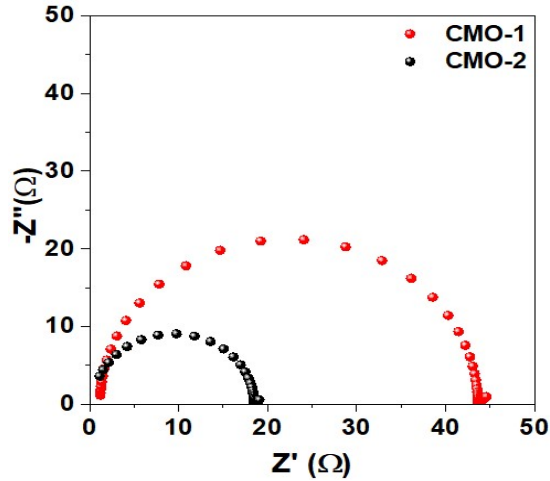


Figure S10. Nyquist plots of CMO-1 and CMO-2 CNFs

Figure S11. (a) Variation of calculated specific capacitance with scan rate, (b) Variation of calculated specific capacitance with current density for CMO-1 and CMO-2 in two different electrolytes and (c) Variation of calculated specific capacitance with current density for CMO-1 and CMO-2 of solid state symmetric supercapacitor

Figure S11(a) shows specific capacitance variation with scan rate for CMO-1 and CMO-2 in different electrolytes, providing insight into supercapacitor performance. Figure S11(b) depicts specific capacitance variation with current density for CMO-1 and CMO-2 in two electrolytes, offering information on behaviour at different charging rates. Figure S11(c) illustrates specific capacitance variation with current density for CMO-1 and CMO-2 in a solid-state symmetric supercapacitor, providing data on performance under different operating conditions.

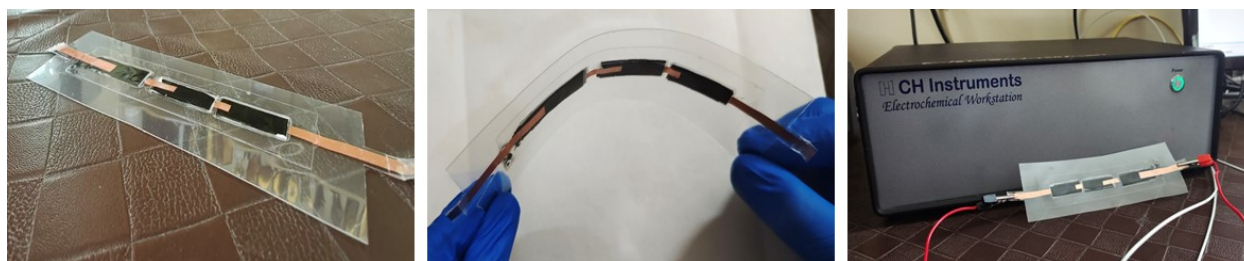


Figure S12. Photographic images of real-time flexible devices

Figure S12 presents photographic images of real-time flexible devices. The images likely depict the actual flexible supercapacitor devices based on the spinel CdMn_2O_4 carbon nanofibers. These images are crucial for demonstrating the supercapacitor devices in real-world application and flexibility. The devices flexibility is an essential characteristic that can contribute to their potential use in various flexible and portable electronic applications.

Table S1. ICP-OES elemental analysis of CMO-1 and CMO-2 showing the concentration of key elements (in ppm).

Element	CMO-1	CMO-2
Al	0.198	0.223
B	0.208	0.278
Cd	685.233	712.236
Co	0.022	0.020
Mn	528.413	1442.127
Ni	0.024	0.022
Se	0.197	0.370
V	0.005	0.011
Zn	0.080	0.097

Table S2. Comparison of three electrode data of reported MnO_2 -based electrode material and Spinel CdMn_2O_4 carbon nanofiber.

Electrode Material	Specific Capacitance	Scan Rate/Current Density	Voltage window	Electrolyte	Reference
$\alpha\text{-MnO}_2$ nanorod	166.22 F g^{-1}	10 mV S^{-1}	0 V- 1.0V	0.1 M Na_2SO_4	³

Carbon nanofiber/MnO ₂	151.1 F g ⁻¹	1 A g ⁻¹	-0.2 V – 0.8V	1.0 M Na ₂ SO ₄	⁴
MnO ₂ /carbon nanotubes–embedded carbon nanofibers	374 F g ⁻¹	2 mV S ⁻¹	0 V– 0.9V	0.1 M Na ₂ SO ₄	⁵
Mn ₂ O ₃ @MnO ₂ composite nanofibers	190 F g ⁻¹	1 A g ⁻¹	0 V– 1.0V	1.0 M Na ₂ SO ₄	⁶
MnO _x Nanofiber	214.7 F g ⁻¹	10 mV S ⁻¹	-0.2 V – 0.8V	0.5 M Na ₂ SO ₄	⁷
Spinel CdMn ₂ O ₄ nanofiber	210 F g ⁻¹	1 A g ⁻¹	0 V– 1.0V	1.0 M Na ₂ SO ₄	⁸
Spinel CdMn ₂ O ₄ carbon nanofiber	For CMO-1-594.7 F g ⁻¹ For CMO-2-547.64 F g ⁻¹	5 mV S ⁻¹	0 V -1.2 V	1.0 M H ₂ SO ₄	This Work

Table S3. Impedance parameters for CMO-1 and CMO-2 derived from electrochemical impedance spectroscopy (EIS) analysis in both three-electrode and symmetric solid-state supercapacitor configurations

Configuration	sample	Rs	Rct	C	W
3 Electrode	CMO-1	0.787	6.13	2.826E-6	0.01423
	CMO-2	0.8001	6.187	2.902E-6	0.01448
2 Electrode	CMO-1	1.981	224.9	1.209E-5	0.008733
	CMO-2	1.486	360.5	1.280E-5	0.00542

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