

Supporting Information

Pore Structure Improvement of Carbon Aerogel and Investigation of Supercapacitive Behavior of Co₃O₄ Nanoball /Carbon Aerogel Composite

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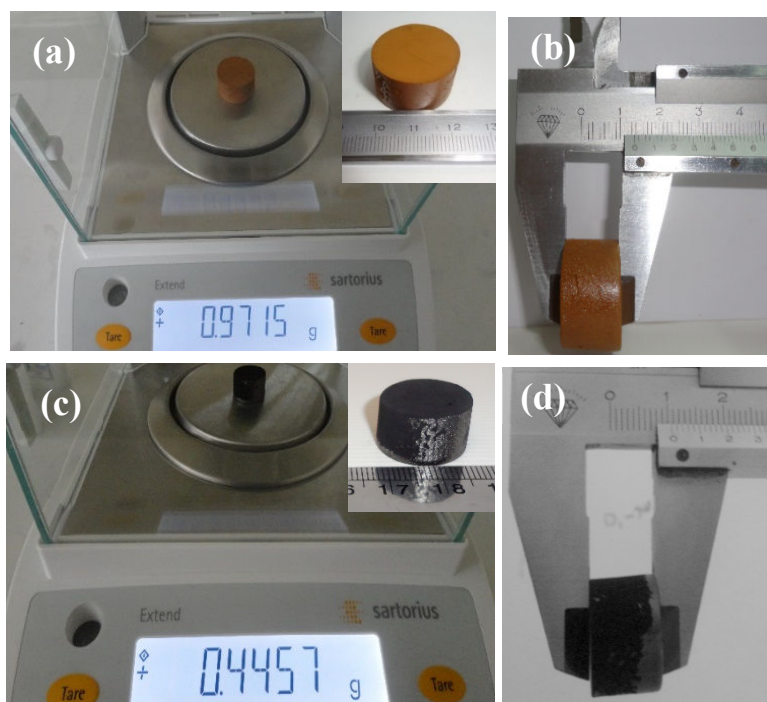


Fig. S1: digital images of OA1, before (a, b) and after carbonization (c, d), the weight measurement process shows the carbon aerogel weight after carbonization of 0.44 g with volume of 2.851 cm³ ($\rho=0.15$ g cm⁻³).

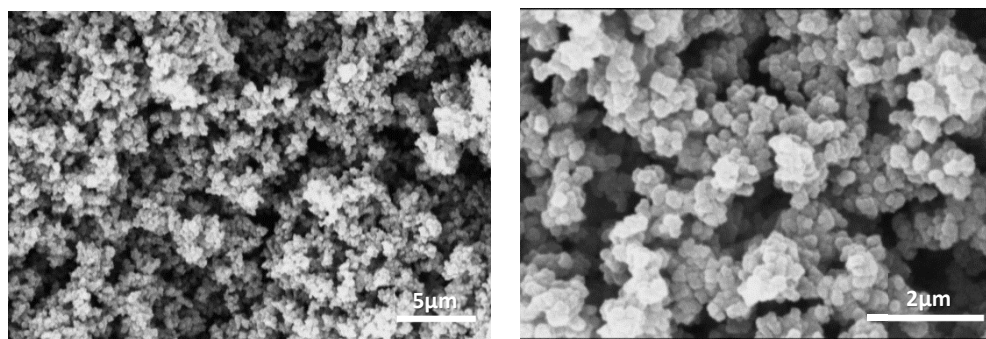


Fig. S2: FE-SEM images of CA2 sample.

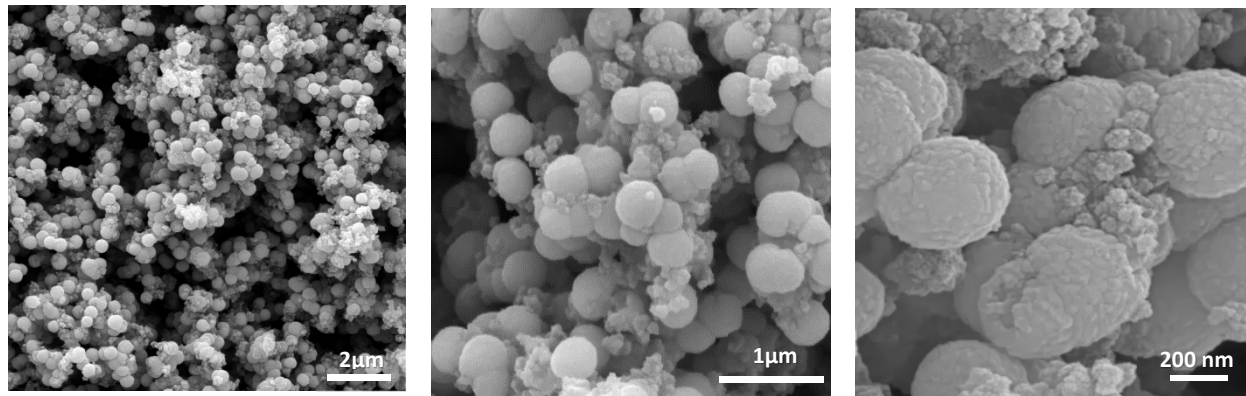


Fig. S3: Additional FE-SEM images of Co₃O₄ nanoballs anchored CA1.

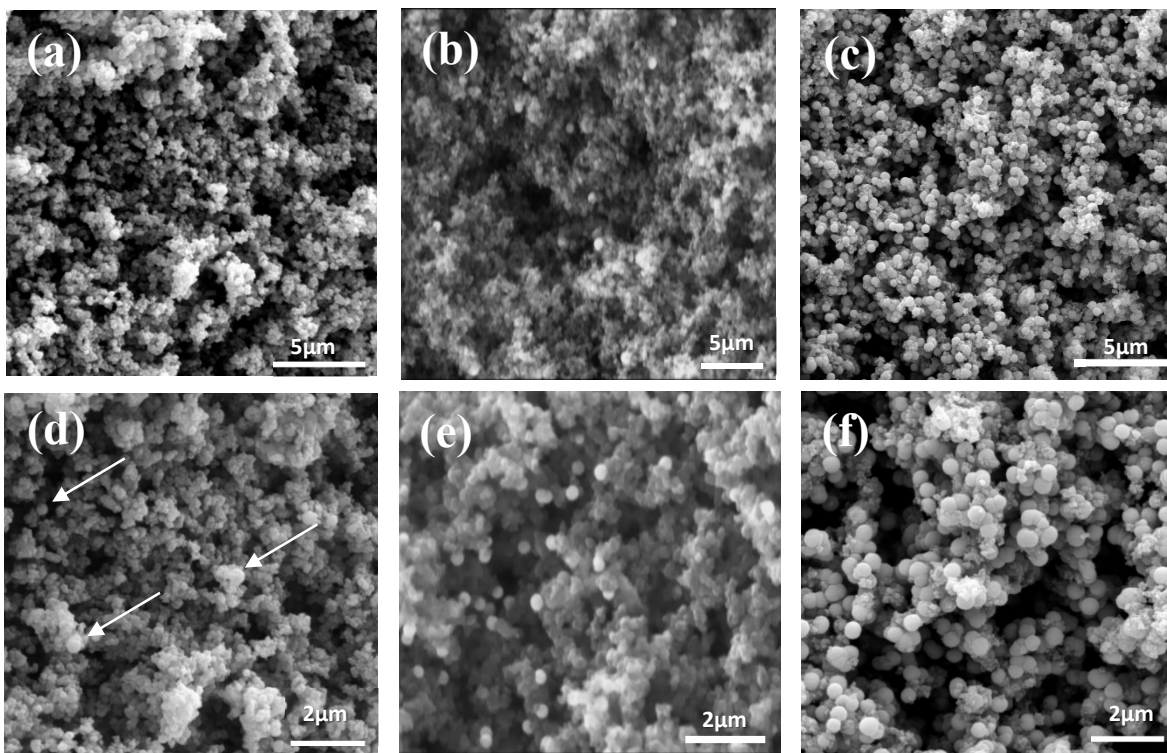


Fig. S4: FE-SEM images of Co₃O₄ nanoballs anchored CA2 (a, d-b, e) and CA1 (c, f), for comparison. Arrows shows Co₃O₄ nanoball.

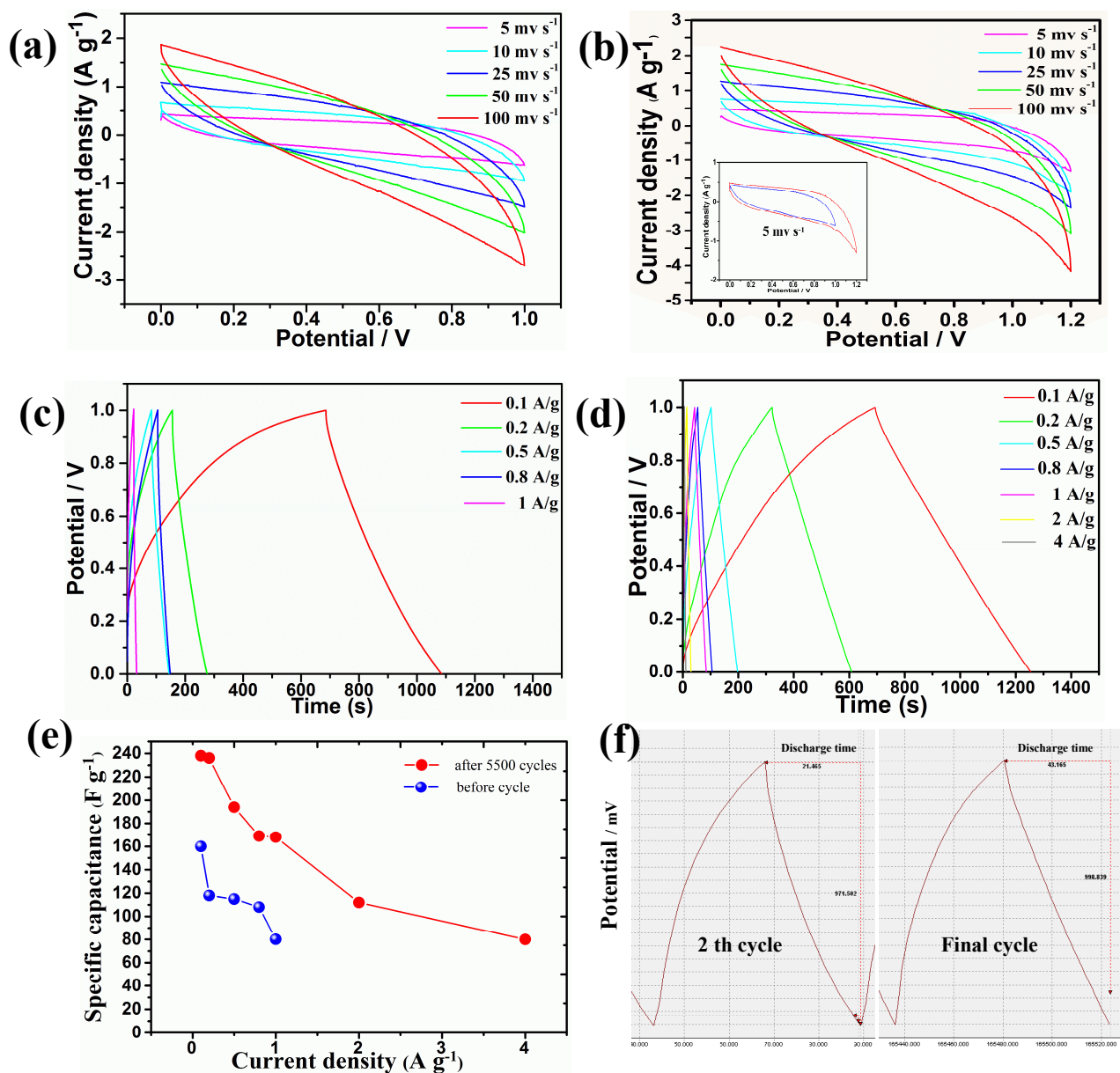


Fig. S5: Cyclic voltammograms of symmetric two-electrode supercapacitor at different scan rates (a, b) and collected at different voltage windows (b, inset). Galvanostatic charge/discharge curves of CAC2 at various current densities before cycling test (c). Galvanostatic charge/discharge curves of CAC2 at various current densities after 5500 cycles (d). The specific capacitance of CAC2 as a function of current density (e). Galvanostatic charge/discharge curves of CAC2 at a current density of 1 A g^{-1} in 2 th cycle and final cycle (6000 th cycle), indicates increase in the discharge time of final cycle (these graphs are retrieved from the software of *Analyzer* version 7.0.9) (f).

Table S1: Comparison of Pore structure properties of carbon aerogels.

Sample	$S_{BET} (\frac{m^2}{gr})$	$V_{total} (\frac{cm^3}{gr})$	$V_{mic} (\frac{cm^3}{gr})$	$V_{meso} (\frac{cm^3}{gr})$	Drying method	ref
CA1	907	6.26	0.35	4.86	Infrared dryer setup	This work
CA2	477	0.44	0.13	0.14	Microwave drying	This work
CA-K-A1000	2796	1.89	1.33	NC	Ambient pressure drying	
CA-A1000	2139	1.00	0.81	NC	Ambient pressure drying	[1]
CA-K-A900	1403	0.64	0.58	NC	Ambient pressure drying	
CA900	1005	1.2	0.54	0.66	Ambient pressure drying	
CA900-A	1539	1.7	0.71	0.99	Ambient pressure drying	[2]
CA-A3	752	NC	0.15	1.33	Hot supercritical drying	[3]
CA#1	669	5.6	0.33	1.9	Cold supercritical drying	[4]
CA-SC-A1000	2617	1.7	0.92	NC	Cold supercritical drying	[5]

Where S_{BET} , V_{mic} , V_{mes} , V_{total} are BET surface area, volume of micro and mesopores and total pore volume respectively.

K-A refers to chemical and thermal activation, respectively.

NC refers to “the data are not calculated

Table S2. Synthetic methods and capacitances of the previously reported Co₃O₄ / carbon-based supercapacitors and Co₃O₄ nanoball/CA composites in our work.

Material	Synthetic method	Current Density	Potential window (ΔV)	Capacitance (F g⁻¹)	Reference
CNT@1600-Co ₃ O ₄ /CC	ALD (atomic layer deposition)	1-10 (mA cm ⁻²)	0.7	350-225	[6]
Co ₃ O ₄ /C-800 °C	precipitation	1-20 (A g ⁻¹)	0.5	~200-195	[7]
PSAC/Co ₃ O ₄ Nps	Hydrothermal method	1-10 (A g ⁻¹)	0.45	~ 100-50	[8]
NCA/ Co ₃ O ₄ Nps	precipitation	1-10 (A g ⁻¹)	0.4	~ 600-450	[9]
RGOA/ Co ₃ O ₄ Nps	Solvothermal process	0.5-10 (A g ⁻¹)	0.5	~ 650-500	[10]
Mesoporous Co ₃ O ₄ /Carbon	Solvothermal process	0.5-5 (A g ⁻¹)	0.6	~ 400-175	[11]
needle-like Co ₃ O ₄ /GNS	Hydrothermal method	0.1-1 (A g ⁻¹)	0.6	~160-63	[12]
Nanoball Co ₃ O ₄ /CA	Hydrothermal method	0.8-10 (A g ⁻¹)	0.7	~350-185.7	This work

Table S3: Electrochemical characteristics of the Co₃O₄ nanoball /carbon aerogel symmetric two-electrode supercapacitor after cycling performance.

Current density (A g⁻¹)	Specific Capacitance (F g⁻¹)	Energy Density (W h Kg⁻¹)	Power Density (W Kg⁻¹)	Coulombic efficiency-η (%)
0.1	232.4	32.22	55.55	83.69
0.2	236.05	32.77	111.1	91.90
0.5	194.3	26.94	277.7	96.03
0.8	169.1	23.47	442.8	98.14
1	168.2	23.33	555.5	97.6
2	112.7	15.5	1107	100
4	80.2	11.11	2222	100

Reference

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