

Supporting Information

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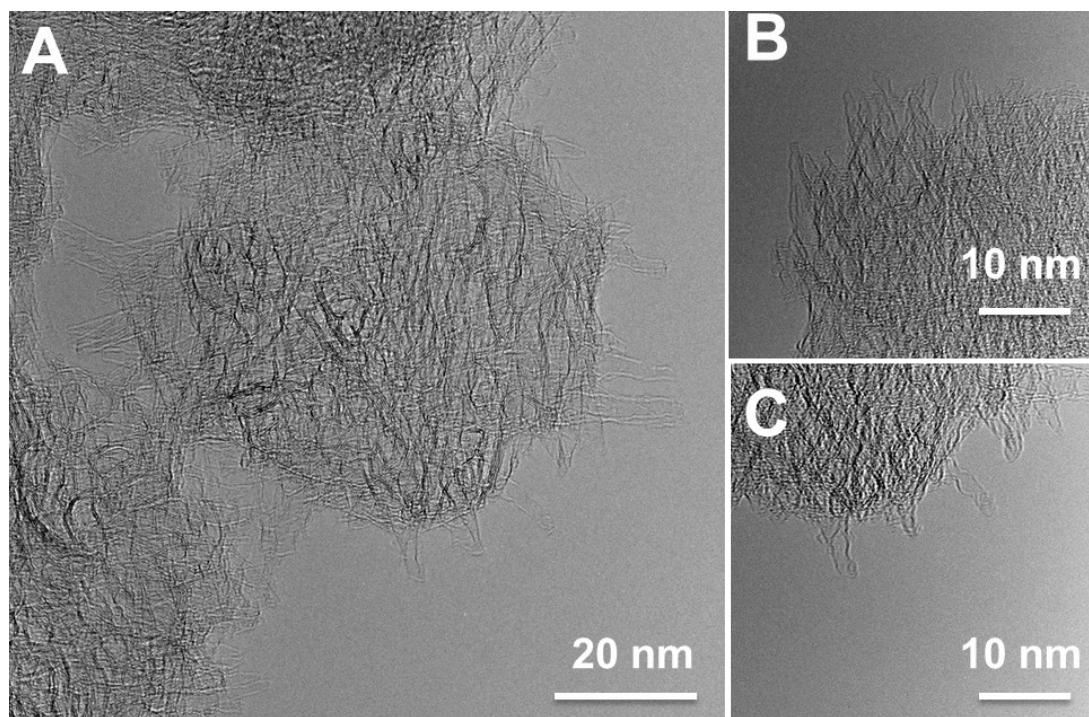


Figure S1. TEM analysis of the CNHs used in this study. (A) TEM of an individual CNH emphasizing the tortuous, defective nature of the carbon comprising the CNH. (B,C) Zoomed-in analysis of the edges of the CNH illustrating the conical carbon terminations characteristic of CNH materials.

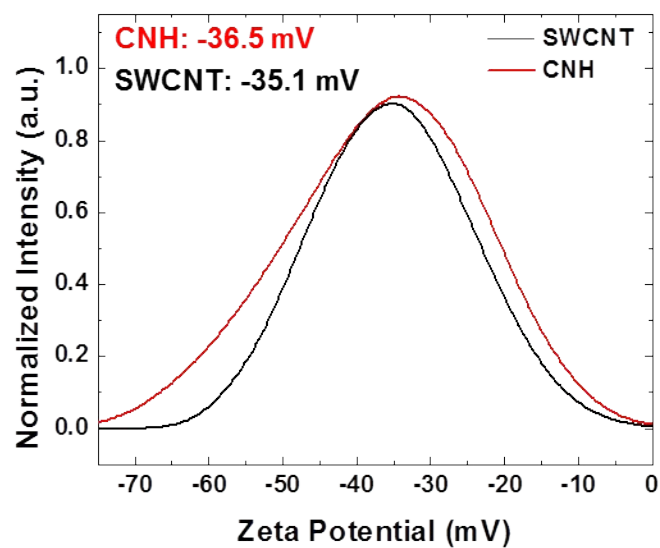


Figure S1. Zeta potential measurements of NMP solutions of the nanostructures used in this study. Measurements were performed on solutions with a concentration of 50 $\mu\text{g/ml}$.

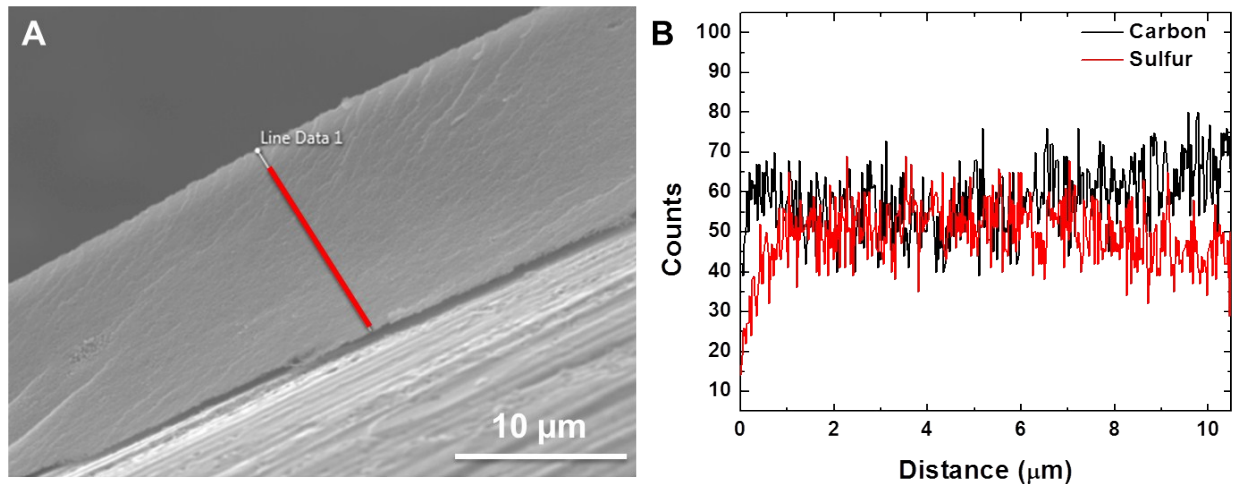


Figure S2. (A) SEM image of a 70% CNH cross-sectional sample infiltrated to 75 wt.% sulfur. The red line inset describes the location of the EDS elemental analysis performed to investigate the uniformity of the sulfur coating. (B) Corresponding EDS linescan emphasizing the uniformity of the sulfur content throughout the entirety of the carbon film.

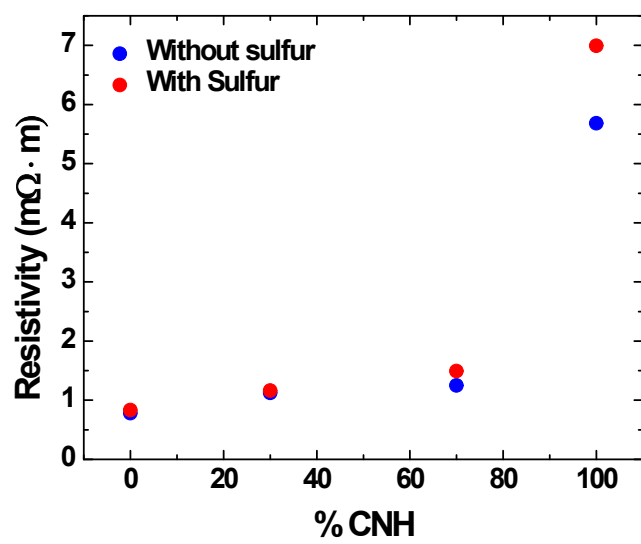


Figure S3. Resistivity measurements of the different hybrid materials before and after sulfur coating.

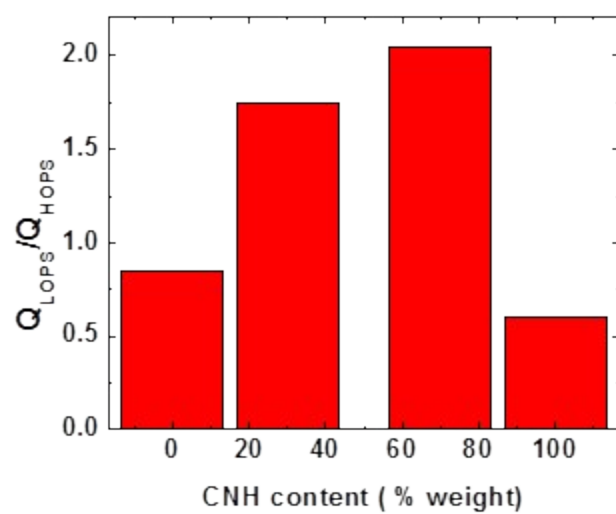


Figure S4. An analysis of the LOPS to HOPS for the rate study presented in Fig. 5A.

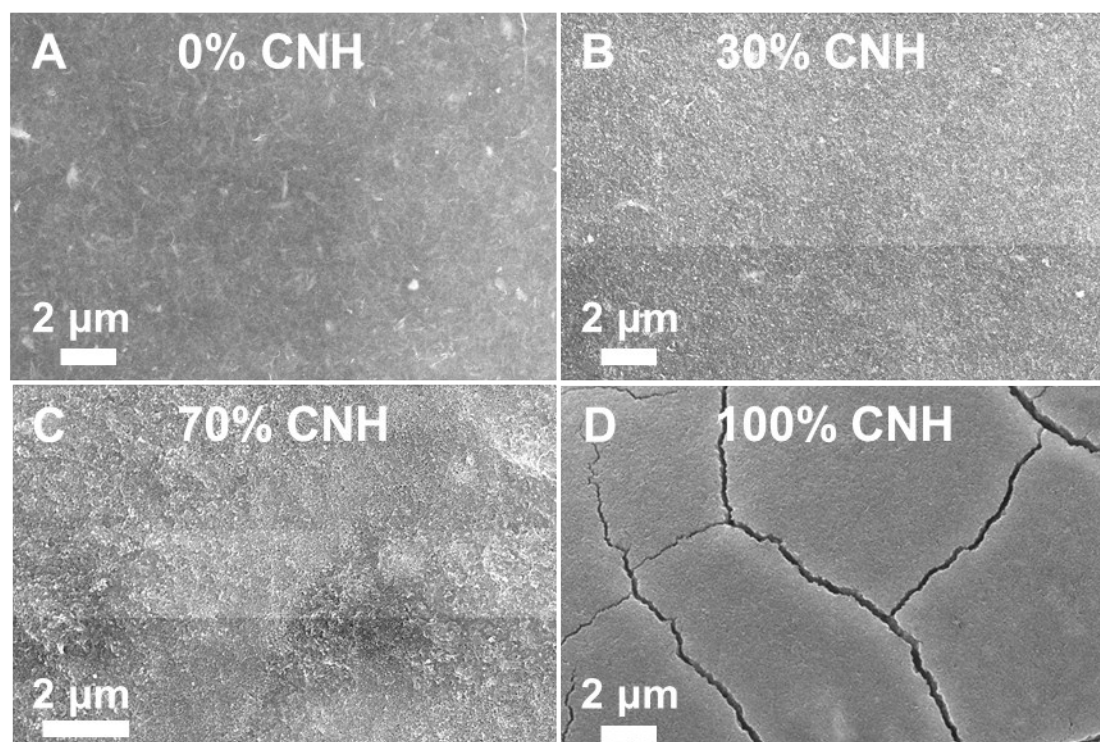


Figure S5. SEM characterization of the large-scale morphologies of the material systems used in this study. Insets indicate the amount of CNH loading present in each system. From these images, it is evident that the SWCNTs impart structural stability to the films and those lacking the presence of any SWCNTs have demonstrate cracking and delamination.

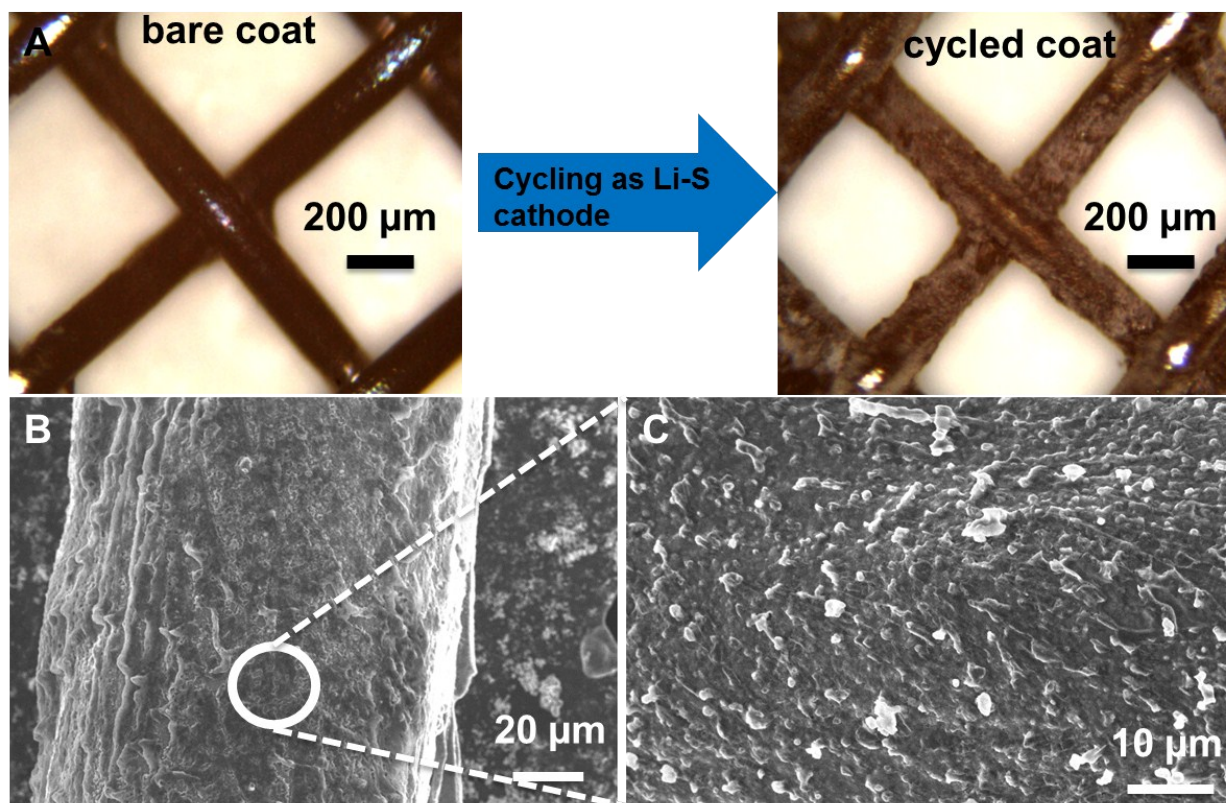


Figure S6. (A) Optical image of the coated mesh before and after cycling. The light colored residue on the cycled mesh electrode is due to dried electrolyte on the samples. (B) SEM characterization of the coated electrode after 10 charge-discharge cycles at a rate of 0.1 C emphasizing the stable adhesion of the carbon-sulfur composite on the mesh after cycling. (C) Higher magnification image of the sample in (B) demonstrating uniformity of the coating at small scales.

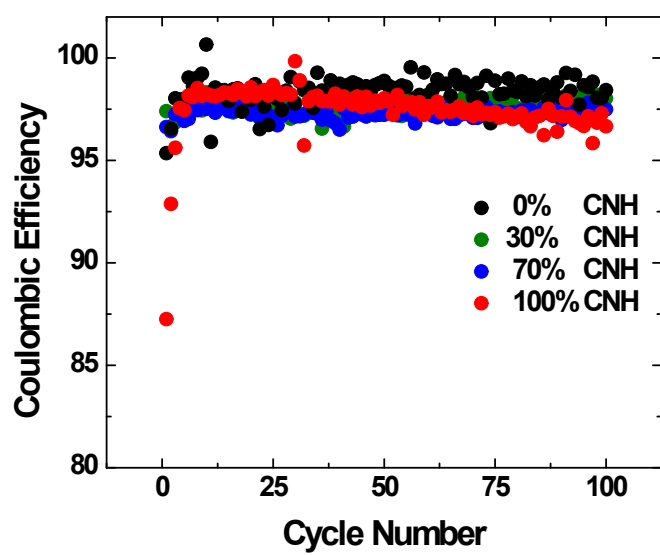


Figure S7. Coulombic efficiency for the devices cycled at 0.2 C.

ref.	capacity (mAh/g)	loading (%)	additives (S:CB:binder)	utilization (mAh/g)
1	1400	37	binder-free	518
2	1382	72	80:10:10	995.04
3	1368	52.5	70:10:20	718.2
4	1374	59	80:10:10	810.66
5	1219	53	binder-free	646.07
6	1633	68	80:10:10	1110.44
7	1120	56	75:10:15	627.2
8	1264	60	binder-free	758.4
9	1239	72	80:10:10	892.08
10	1070	56	binder-free	599.2
11	1400	35	70:20:10	490
12	1048	50	binder-free	524
13	1246	63.7	85:12:03	793.702
14	1346	62	binder-free	834.52
15	1340	62	binder-free	830.8
16	1021	57.12	80:10:10	583.1952
17	1260	58.4	80:15:10	735.84
18	911	53	binder-free	482.83
19	1317	48.3	70:20:10	636.111
20	1070	53.6	80:12:08	573.52
21	1310	63	binder-free	825.3
22	1620	65	binder-free	1053
23	800	52	binder-free	416
24	750	76.5	85:15:00	573.75
25	1010	62	90:10:00	626.2
26	1278	47.25	75:15:10	603.855
27	998	33.6	70:15:15	335.328

Table S1. Detailed description of the relevant parameters used to generate the plot in Figure 5. In order to calculate utilization, the sulfur loading was multiplied by the discharge capacity.

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