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Electronic Supplementary Information

MOF-Derived Conductive Carbon Nitrides for Separator-Modified

Li-S Batteries and Flexible Supercapacitors

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Fig. S1 XRD of the Ni-based MOF precursor.



Fig. S2 SEM image of the incomplete generated c-CN, showing that Ni serves as the catalytic center for the generation of c-CN.



Fig. S3 Digital photographs showing the preparation of c-CN with hydrophobic nature during and after the etching process.



Fig. S4 SEM image of $g-C_3N_4$ nanosheets prepared from the conventional thermal condensation of urea. The edges of nanosheets are indicated by red arrows.



Fig. S5 XPS high-resolution C 1s spectrum of (a) c-CN and (b) $g-C_3N_4$.



Fig. S6 TEM image of c-CN, demonstrating porosities and morphological defects.



Fig. S7 The pore size distribution of c-CN.



Fig. S8 Sheet resistance map of the $g-C_3N_4$ and MWCNTs films.



Fig. S9 Digital photograph showing the Li₂S₆ adsorption tests of g-C₃N₄, MWCNTs, and c-CN.



Fig. S10 XPS high-resolution S 2p spectrum of $g-C_3N_4$ after the adsorption test of Li_2S_6 for 3 h.



Fig. S11 Binding geometric configurations and binding energies of Li_2S_6 with g-C₃N₄, the strong interaction induces the distortion of C₃N₄ layer. The hydrogen, lithium, carbon, nitrogen, and sulfur atoms are marked with pink, brown, green, blue, and yellow, respectively.



Fig. S12 ^{13}C nuclear magnetic resonance (NMR) spectrum of g-C_3N_4 before and after Li_2S_6 adsorption.



Fig. S13 The geometrical configuration and binding energy of Li_2S_6 with defected c-CN. The lithium, carbon, and sulfur atoms are marked with brown, green, and yellow, respectively.



Fig. S14 Cross-sectional SEM and EDS characterization showing the one-side coating of c-CN on the separator after a Li_2S_6 permeation test.



Fig. S15 STEM image and corresponding elemental maps of c-CN after ${\rm Li}_2{\rm S}_6$ adsorption.



Fig. S16 TGA curve of the sulfur-graphene hybrid, showing a sulfur loss of 79.35%.



Fig. S17 CV profiles of (a) no interlayer, (b) $g-C_3N_4$ interlayer, and (c) c-CN interlayer-based cells at various scan rates from 0.1-0.5 mV s⁻¹. (d) Reaction kinetics with respect to the Li⁺ ion diffusion properties of peak ii at various scan rates.



Fig. S18 EIS of the bare PP, $g-C_3N_4$ PP and c-CN PP based cells.



Fig. S19 Galvanostatic charge–discharge curves of c-CN PP based cell at various current densities.



Fig. S20 Cyclic performance at 5.0 C of g-C₃N₄ PP and c-CN PP, respectively.



Fig. S21 SEM inspection of the c-CN PP a) before, and b) after 200 cycles at 2.0 C. c) The precipitation of solid short-chain polysulfides on the c-CN after battery cycling.



Fig. S22 EIS spectrum of c-CN-derived supercapacitor in a three-electrode configuration.



Fig. S23 Galvanostatic charge-discharge curves of c-CN at various current density of $1-10 \text{ A g}^{-1}$ in a three-electrode configuration.



Fig. S24 (a) CV profiles of c-CN-derived SSC at different scan rates of 10–100 mV s⁻¹. (b) Galvanostatic charge-discharge curves of SSC at different current densities of 1–5 A g⁻¹.

Table S1 Comparison of battery performances at high sulfur loading conditions based on variousmodified-separators between this work and other reported studies.

Coating materials	Host	Loading (mg cm ⁻²)	S conten t (wt%)	Rate (C)	Cycles	Initial capacity (mAh cm ⁻²)	Capacity decay (% per cycle)	Ref.
c-CN	Graphene	3.56	63	1.0	100	2.82	0.32	This work
		5.23	60	0.2	100	5.75	0.21	
		9.80	60	0.2	50	5.77	0.56	
MoS ₂ -PDDA	Ketjen Black	4.0	60	1.0	100	3.1	0.35	1
PAN@APP	Carbon powders	6.0	70	0.2	50	3.66	0.07	2
N-doped graphene	Graphene	3.8	61	0.3	30	4.45	1.06	3
Co ₉ S ₈	Super P	5.6	70	-	200	5.05	0.08	4
rGO@SL	Carbon powders	3.8	50	0.1	50	2.30	0.34	5
Ce- MOF/CNT	Ketjen Black	2.5	-	1.0	800	2.55	0.02	6
LDH/NG	CNTs	4.3	63	0.5	100	4.64	0.26	7
Ni ₃ (HITP) ₂ or ZIF-8	Carbon black	3.5	64	1.0	500	2.98	0.03	8

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