

Electronic Supplementary Information

Rational design of Co embedded N, S-codoped carbon nanoplates as anode materials for high performance lithium-ion batteries

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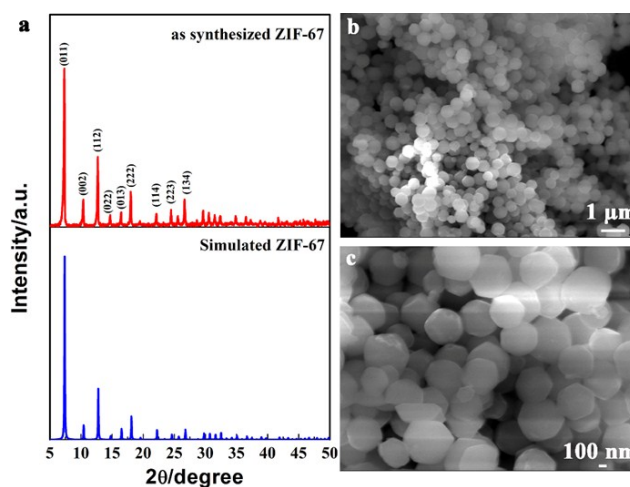


Fig. S1 (a) XRD patterns and (b, c) FE-SEM images of the ZIF-67.

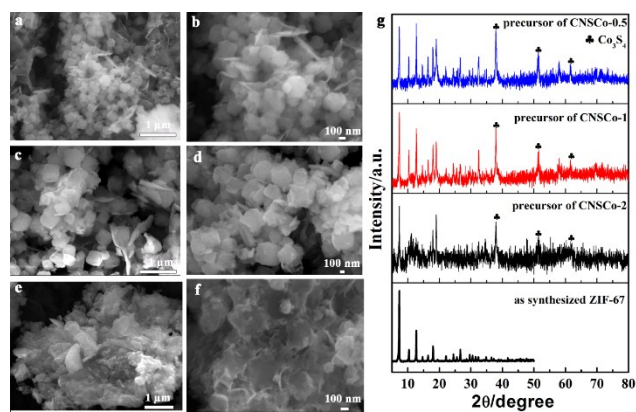


Fig. S2 FE-SEM images of the CNSCo-0.5 precursor (a, b), the CNSCo-1 precursor (c, d), the CNSCo-2 precursor (e, f) and XRD patterns of all the precursors.

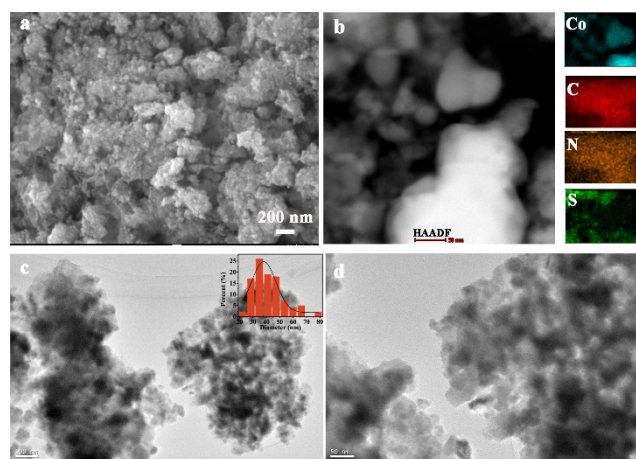


Fig. S3 (a) FE-SEM images, (b) HAADF-STEM images and (c, d) TEM images with corresponding elemental mappings of the CNSCo-2.

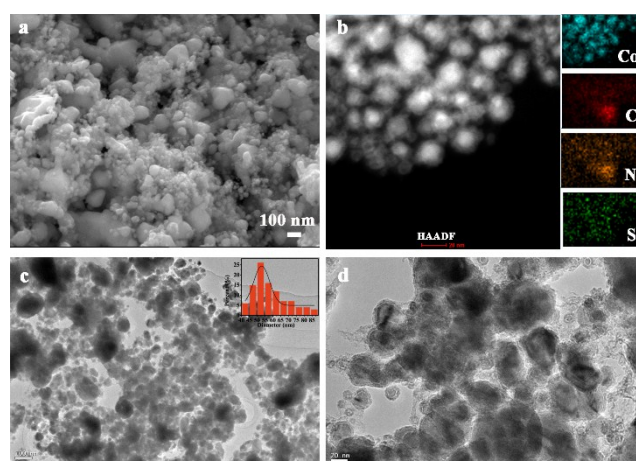


Fig. S4 (a) FE-SEM images, (b) HAADF-STEM images and (c, d) TEM images with corresponding elemental mappings of the CNSCo-1.

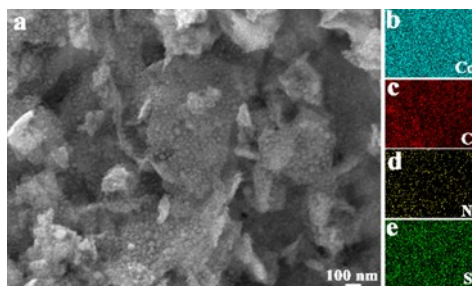


Fig. S5 (a) FE-SEM images and (b, c, d, e) corresponding elemental mappings of the CNSCo-0.5.

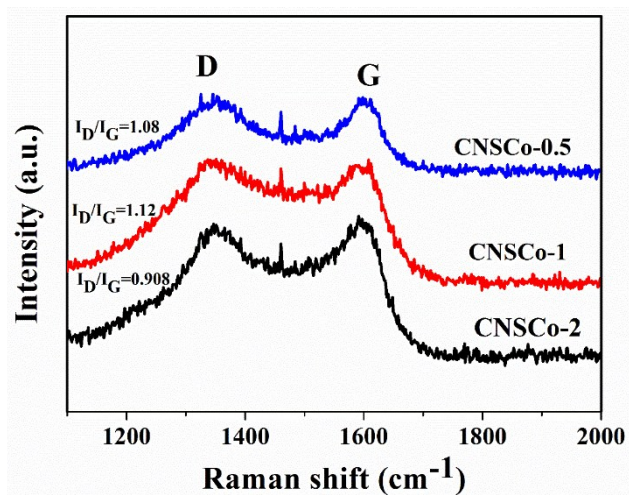


Fig. S6 Raman spectra for the CNSCo-2, CNSCo-1 and CNSCo-0.5 samples.

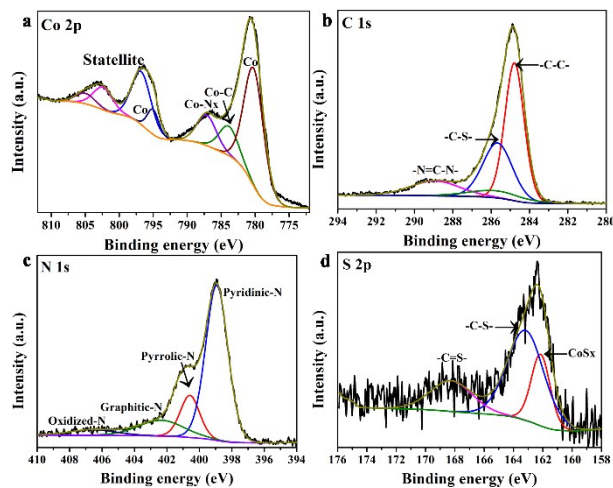


Fig. S7 High-resolution XPS spectra and fitted curves of (a) Co 2p, (b) C 1s, (c) N 1s, (d) S 2p of the CNSCo-1.

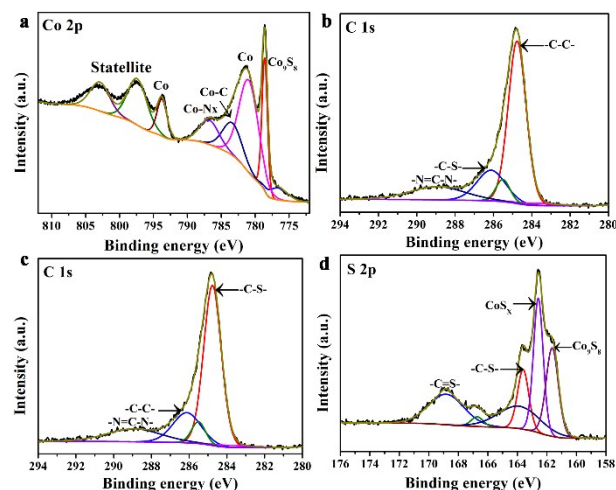


Fig. S8 High-resolution XPS spectra and fitted curves of (a) Co 2p, (b) C 1s, (c) N 1s, (d) S 2p of the CNSCo-2.

Table S1. Elemental contents of the CNSCo-2, CNSCo-1, CNSCo-0.5 from XPS.

Sample	C (at. %)	N (at. %)	S (at. %)	Co (at. %)
CNSCo-2	59.08	4.03	17.84	19.05
CNSCo-1	72.05	9.01	1.36	17.58
CNSCo-0.5	76.80	3.82	1.31	18.07

N₂ sorption measurement at 77 K was performed to analysis the specific surface area and porosity of the CNSCo-x samples (Fig. S9). The CNSCo-0.5 exhibits the highest Brunauer–Emmett–Teller (BET) specific surface area (113.13 m² g⁻¹) among the obtained samples, while the CNSCo-2 and CNSCo-1 have BET surface areas of 62.56 and 29.66 m² g⁻¹, respectively. The high BET surface area of the CNSCo-0.5 can provide a large interface to facilitate the easy uptake and release of lithium ions in electrolyte. The three obtained CNSCo-x samples all possess two-size mesoporous pores, 4.0 nm, 19.0 nm of the CNSCo-2, 2.2 nm, 13.1 nm of the CNSCo-1, 3.8 nm, 33.0 nm of the CNSCo-0.5. The mesopores of the CNSCo-2 and CNSCo-1 are from the space among non-uniform nanoparticles. The smaller mesopores originate from the space of nanoparticles while the larger mesopores may come from the space with in the carbon networks.

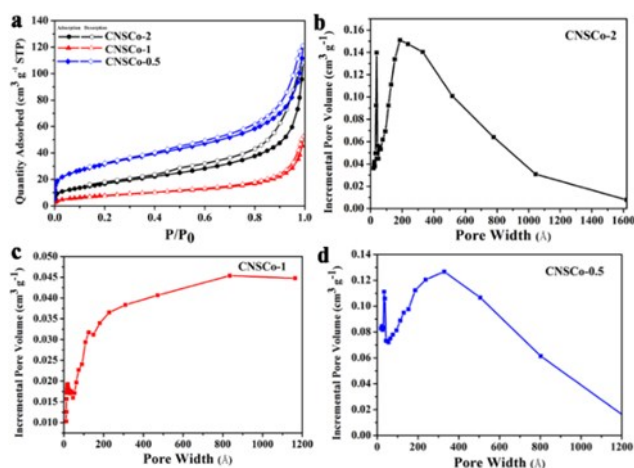


Fig. S9 (a) N₂-sorption isotherms of the CNSCo-2, CNSCo-1, CNSCo-0.5, and pore size distribution curves for (b) the CNSCo-2, (c) the CNSCo-1, and (d) the CNSCo-0.5.

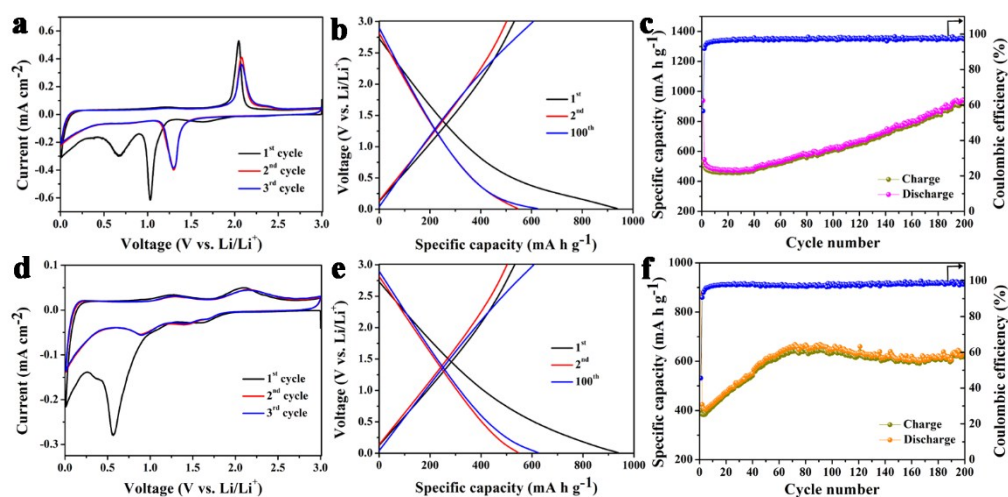


Fig. S10 (a) CV curves at a scan rate of 0.1 mV s⁻¹ in the range of 0.01-3.0 V, (b) charge-discharge profiles of 1st, 2nd, 100th cycle at current density of 200 mA g⁻¹ of the CNSCo-2, (c) cycling performance and coulombic efficiency at current density of 200 mA g⁻¹, and (d) CV curves at a scan rate of 0.1 mV s⁻¹ in the range of 0.01-3.0 V, (e) charge-discharge profiles of 1st, 2nd, 100th cycle at current density of 200 mA g⁻¹ of the CNSCo-1, (f) cycling performance and coulombic efficiency at current density of 200 mA g⁻¹.

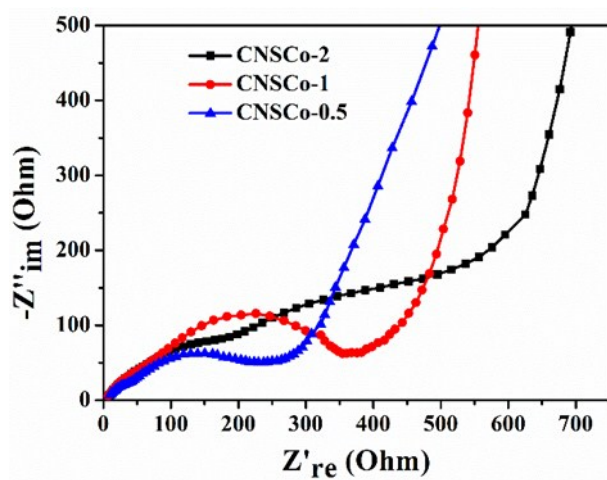


Fig. S11 EIS of the CNSCo-2, CNSCo-1, CNSCo-0.5.

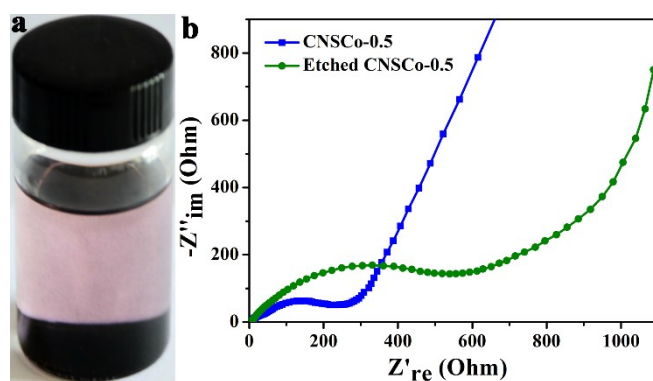


Fig. S12 (a) The photo of pink solution and (b) EIS of the CNSCo-0.5 etched by 1.0 M HCl.

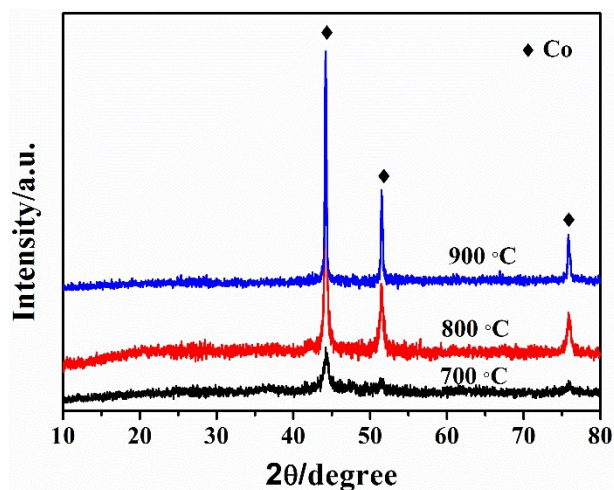


Fig. S13 XRD patterns of the CNSCo-0.5 synthesized at 700 °C, 800 °C and 900 °C.

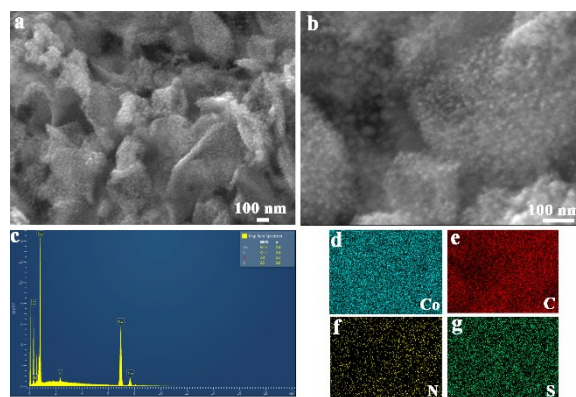


Fig. S14 (a, b) FE-SEM images, (c) EDS image and (d, e, f, g) corresponding elemental mappings of the CNSCo-0.5 synthesized at 700 °C.

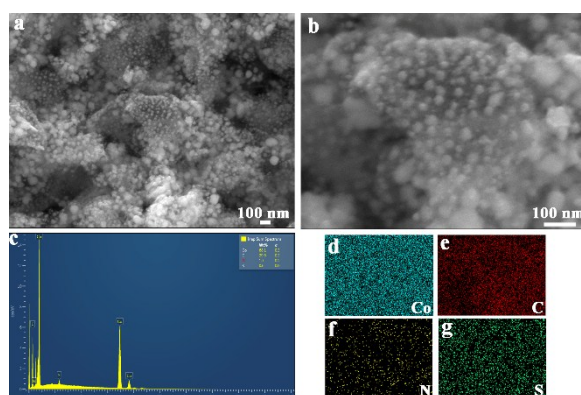


Fig. S15 (a, b) FE-SEM images, (c) EDS image and (d, e, f, g) corresponding elemental mappings of the CNSCo-0.5 synthesized at 800 °C.

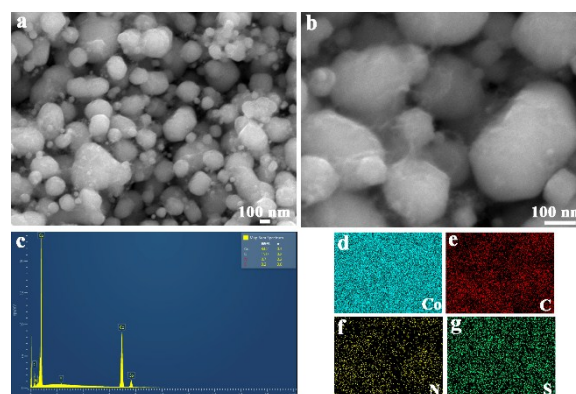


Fig. S16 (a, b) FE-SEM images, (c) EDS image and (d, e, f, g) corresponding elemental mappings of the CNSCo-0.5 synthesized at 900 °C.

Table S2. Elemental contents of the CNSCo-0.5 synthesized at 700 °C, 800 °C and 900 °C from EDS.

Sample	C (at. %)	N (at. %)	S (at. %)	Co (at. %)
700 °C	70.32	4.10	0.38	25.20
800 °C	65.46	3.22	0.51	30.81
900 °C	45.68	1.83	0.22	52.27

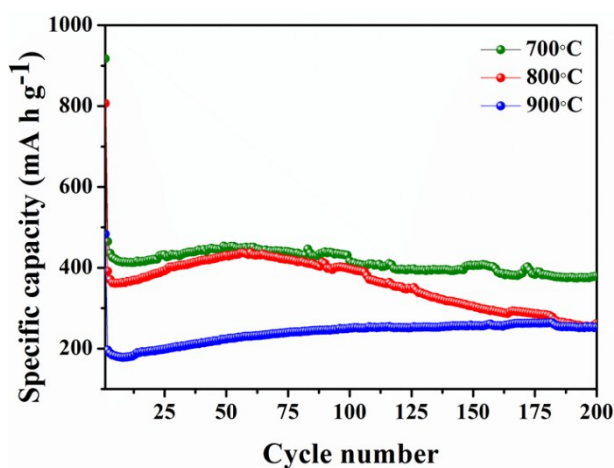


Fig. S17 Discharge capacities of the CNSCo-0.5 synthesized at 700 °C, 800 °C and 900 °C.

Table S3. The electrochemical performance of heteroatom-doped carbon materials as anode materials for LIBs.

Sample	Current density (mA g ⁻¹)	Cycle number	Capacity (mA h g ⁻¹)	Ref.
CNSCo-0.5	200	200	1360.4	Our work
N-C octahedral particles	100	100	890	1
NS-C films	200	2000	357.2	2
NSDPC	100	50	864	3
NFCs	100	300	645	4
NPCMs	100	400	655.1	5
C60@N-MPC	1000	400	1077	6
3D NS-GSs	100	80	1117	7
N-doped graphene	50	30	827	8

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