## Supporting information

## High-Rate Lithium-Sulfur Battery Assisted with Ultrafine La<sub>2</sub>O<sub>3</sub>

## Nanoparticles Decorated and Nitrogen-Enriched Mesoporous Carbons

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Figure S9. EIS before 1st cycle of nitrogen-free MC/La<sub>2</sub>O<sub>3</sub>/S composites.

	Element analysis			XPS				<b>c</b> [d]	
Samples	Ν	N/C	Ν	N/C	pyridinic N <sup>[a]</sup>	pyrrolic N <sup>[b]</sup>	graphitic N <sup>[c]</sup>	$S_{\text{BET}}$	$V_{T}^{lej}$
	wt.%	at./at.	at.%	at./at.	%	%	%	$m^2 g^{-1}$	$\mathrm{cm}^3 \mathrm{g}^{-1}$
MCs	-	-	-	-	-	-	-	847	2.6
NMCs	8.1	0.08	20.4	0.31	56.2	31.7	12.1	731	2.6

Table S1. Chemical nature of the NMCs and MCs

[a] the relative concentration of pyridinic N (398.5 $\pm$ 0.3 eV); [b] the relative concentration of pyrrolic N (400.5 $\pm$ 0.3 eV); [c] the relative concentration of graphitic N (401.6 $\pm$ 0.3 eV); [d] BET specific surface area; [e] total pore volume (P/P<sub>0</sub>= 0.985).

In our synthesis approach, the incorporation of high-nitrogen-content melamine into phenolic precursors can transfer nitrogen atoms into carbon framework under pyrolysis conditions. The N/C mole ratio of NMCs is 0.31 determined from high-resolution XPS, three times higher than the value obtained from elemental analysis. The variation of the values can be explained by the surface specificity of XPS measurements, suggesting the N atoms are apt to gather in the surface rather than the bulk of carbon framework.

Sample	${S_{\scriptscriptstyle BET}}^{[a]} {m^2 g^{-1}}$	$V_{T}^{[b]}$ cm <sup>3</sup> g <sup>-1</sup>
5La-MC	755	2.2
10La-MC	754	2.0
15La-MC	752	1.9
5La-NMC	674	2.4
10La-NMC	670	2.3
15La-NMC	668	2.1

Table S2. Pore structure parameters of the MC/La<sub>2</sub>O<sub>3</sub> composites and NMC/La<sub>2</sub>O<sub>3</sub> composites

[a] BET specific surface area; [b] total pore volume (P/P0= 0.985).

After incorporating  $La_2O_3$  onto the carbon framework, the porosity and the specific surface area show the expected decrease due to the introduction of heavier  $La_2O_3$  nanoparticles. Similar pore volume and  $S_{BET}$  have achieved at the same  $La_2O_3$  loading, which help us avoid the complication of the different pore structures while focus on the nitrogen doping determined electrochemical performance afterwards.



*Figure S1.* (a) XPS survey spectra of NMCs and MCs; (b) high-resolution XPS  $N_{1s}$  spectra of NMCs. A strong  $N_{1s}$  peak is observed for NMCs, while no obvious peak corresponding to nitrogen is found for MCs. The  $N_{1s}$  spectra were curve-fitted into three peaks with binding energies of 398.5, 400.5 and 401.6 eV corresponding to pyridinic N, pyrrolic N, and graphitic N, respectively.



*Figure S2*. TEM image of 10La-MC composites. Some obvious  $La_2O_3$  aggregates could be observed in the nitrogen-free 10La-MC composites due to the Ostwald ripening and/or migration-coalescence of  $La_2O_3$  particles easily occur during the nucleation and thermal-decomposition process.

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*Figure* **S3.** Coulombic efficiency of NMC/La<sub>2</sub>O<sub>3</sub>/S composites at 0.2 C (a), 1 C (b), 3 C (c) and 5 C (d). With 10 wt.% La<sub>2</sub>O<sub>3</sub> decoration, the ternary 10La-MC/S composites exhibits coulombic efficiency of 94.9 %, 96.2 %, 97.1 % and 97.2 %, respectively at 0.2 C, 1 C, 3 C and 5 C after 100 cycles.



*Figure S4.* EIS before 1st cycle of NMC/La<sub>2</sub>O<sub>3</sub>/S composites. The semicircle diameters in the high frequency region of Nyquist plots indicate that the charge-transfer resistances are highest in 15La-NMC/S composites. The lowest electronic conductivity should be due to the excessive insulated  $La_2O_3$  decoration.



*Figure* **S5.** Thermogravimetric analysis (TGA) curves of MC/La<sub>2</sub>O<sub>3</sub>/S composites. The sulfur contents of the MC/La<sub>2</sub>O<sub>3</sub>/S composites are also carefully controlled at the same of 60 wt. % as that of the NMC/La<sub>2</sub>O<sub>3</sub>/S composites. This could thus minimize the influence of the sulfur content, which could prompt us to focus on the effect of the La<sub>2</sub>O<sub>3</sub> decorating and the nitrogen doping on the electrochemical performances of the resulting composites.



*Figure S6.* Typical cyclic voltammogram of nitrogen-free MC/La<sub>2</sub>O<sub>3</sub>/S composites at 0.2 mV s<sup>-1</sup> (a) and 2 mV s<sup>-1</sup> (b). Similar with the nitrogen doped NMC/La<sub>2</sub>O<sub>3</sub>/S composites, the sulfur reduction onset potentials gradually increase with the increase of La<sub>2</sub>O<sub>3</sub> loading, especially at a high scanning rate of 2 mV s<sup>-1</sup>. The higher reduction potential indicates improved kinetics by the La<sub>2</sub>O<sub>3</sub> decoration.



*Figure* **S7.** Initial charge-discharge curves of nitrogen-free MC/La<sub>2</sub>O<sub>3</sub>/S composites at 0.2 C (a), 1 C (b), 3 C (c) and 5 C (d). At a low rate of 0.2 C, the initial discharge capacity could be increased from 1041 mAh g<sup>-1</sup> (based on the mass of sulfur) for MC/S to 1149 mAh g<sup>-1</sup> by La<sub>2</sub>O<sub>3</sub> decoration. At a high rate of 5C (8.4 A g<sup>-1</sup>), the high initial discharge capacity of 514 mAh g<sup>-1</sup> could be achieved for 10La-MC/S, much improved compared to the capacity of 334 mAh g<sup>-1</sup> for MC/S.

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*Figure S8.* Cycle stability of nitrogen-free MC/La<sub>2</sub>O<sub>3</sub>/S composites at 0.2 C (a), 1 C (b), 3 C (c) and 5 C (d). With 10 wt.% La<sub>2</sub>O<sub>3</sub> decoration, the ternary 10La-MC/S composites deliver reversible capacities of 676, 541, 416 and 367 mAh g<sup>-1</sup> respectively at 0.2 C, 1 C, 3 C and 5 C after 100 cycles.



**Figure S9.** EIS before 1st cycle of nitrogen-free MC/La<sub>2</sub>O<sub>3</sub>/S composites. Similar with the nitrogen doped 15La-NMC/S composite, the 15La-MC/S composite also exhibits the highest charge-transfer resistances.