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Encapsulating sulfur into hybrid porous carbon/CNTs substrate as cathode for lithium-sulfur batteries

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Fig. S1 (a)TG curves of the A-MCxy carbon substrate recorded under an argon atmosphere with the heating rate of 10 $^{\circ}$ C min⁻¹; (b) FTIR spectra of the A-MCxy carbon substrates.

Table S1 Weight loss of the as-prepare	d carbon substrates and	l the sulfur/carbon com	posites
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sample	A-MC12	S/A-MC12	A-MC11	S/A-	A-MC21	S/A-MC21
				MC11		
weight loss/ wt %	2.6	80.5	10.5	81.7	5.0	80.2



Fig. S2 SEM images: (a) the A-MC12 carbon substrate and (b) the S/A-MC12 composite; (c) the A-MC21 carbon substrate and (d) the S/A-MC21composite; (e) commercial carbon black. (The white scale bar is 100 nm.)



Fig. S3 Low and high-magnification TEM observations of the A-MC11 carbon substrate.



Fig. S4. Cycle performance of the as-prepared S/A-MCxy composites with the sulfur loading of $1.2-1.5 \text{ mg cm}^{-2}$ at the current density of 160 mA g⁻¹ (composite).



Fig. S5. Cycle performance of the S/A-MC11 composite with a higher sulfur loading at the current density of 160 mA g⁻¹ (composite) after first two cycles at 80 mA g⁻¹ (composite).



Fig S6. The discharge curves of the S/A-MC11composite at different current densities.

Cycle number	$\operatorname{Ret}\left(\Omega\right)$	$Zw\left(\Omega ight)$
Before	78 2	212.9
discharge	70.2	212.)
1 st	26.4	35.63
5 th	31.8	81.95
20 th	40.2	140.2
50 th	49.7	162.1
100 th	66.3	191.7

Table S2 The simulated data from EIS spectra of the cathode material at full charged state in the different cycles.