Electronic Supplementary Information for

Polyacrylonitrile/Graphene composite as a precursor of sulfur-based cathode material for high-rate rechargeable Li-S batteries

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Electronic Supplementary Material (ESI) for Energy & Environmental Science This journal is $\ensuremath{\mathbb{O}}$ The Royal Society of Chemistry 2012



Fig. S1. Photographs of aqueous dispersions of GO (a), AN (b), PAN (c), PAN/GO (b) and PAN/GNS (e).

Table S1. GO or GNS content in the products:

Samples	GO or GNS content in ca. wt%			
PAN/GO ^a	20	10	5	
PAN/GNS ^b	16	8	4	
pPAN-S/GNS ^c	8	4	2	

^aAs-prepared PAN/GO

^bAfter reduction (GNS contents are less than these of GO due to the loss of functional groups in the process of reduction.)

^cAfter thermal treatment with S (GNS contents decreased due to the S was embedded in composite after thermal treatment.)

Table S2. Component distributions of the final products in ca. wt%:

Samples	S	pPAN	GNS
pPAN-S		53	0
pPAN-S/GNS 2%	47	51	2
pPAN-S/GNS 4%		49	4
pPAN-S/GNS 8%		45	8

 Table S3. BET measurement results of different samples:

Samples	PAN	pPAN-S	PAN/GNS	pPAN/GNS	pPAN-S/GNS
BET total surface area (m^2g^{-1})	19.41	10.76	37.25	35.85	25.41



Fig. S2. SEM images of PAN/GNS (a) and pPAN/GNS (b).



Fig. S3. TEM image and elemental mapping of pPAN-S/GNS 4% composite. (a) TEM image of pPAN-S/GNS composite; Elemental mapping of (b) sulfur and (c) EDS spectrum captured for the region shown in (a).



Fig. S5. FTIR spectra of GO, PAN and PAN/GO composite.



Fig. S6. TEM images of PAN/GNS (a) and pPAN-S/GNS (b) composite with GNS content ca. 8%



Fig. S7. First discharge/charge profiles of pPAN-S/GNS composite electrodes with 8% and 4% GNS.



Fig. S8. Electronic conductivities of the pPAN-S/GNS composites with different GNS contents.