Electronic Supplementary Material

Rational design of metal oxide hollow nanostructures decorated carbon nanosheets for superior lithium storage

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Fig.S1 Photos of $Fe(NO_3)_3$ (0.05 g) and GO (0.01 g) mixed solution (20 mL) with (a) or without (b) PVP (0.01 g).



Fig.S2 SEM (a), TEM (b) images and XRD pattern (c) of Fe_2O_3/G .



Fig.S3 XRD patterns of GO and Fe(NO₃)₃@PVP@GO.



Fig.S4 (a) Low magnification SEM image of $Fe(NO_3)_3$ @PVP@GO, suggesting the dense and bulk particles. (b) SEM image of a bulk $Fe(NO_3)_3$ @PVP@GO particle, indicating the thick and multilayered nanostructures.



Fig.S5 Thermogravimetric analysis of GO, PVP, iron nitrate and Fe(NO₃)₃@PVP@GO in nitrogen atmosphere at 10 °C min⁻¹ heating rate.



Fig.S6 Photo image (left) of 500 mg Fe(NO₃)₃@PVP@GO powder in a glass container, after carbonation process, the volume of the resulting 80 mg FeO_x@C@G exceeds that of Fe(NO₃)₃@PVP@GO (right).



Fig.S7 HRTEM images of h-Fe₂O₃@C@G with 84 wt.% Fe₂O₃(a), 78 wt.% Fe₂O₃(b) and 70 wt.% Fe₂O₃(c).



Fig.S8 (a) Elemental analysis of h-Fe₂O₃@C@G hybrids (with 0.3 g PVP dose), in which element C, N, Fe and O uniformly disperse in the observed zone; SEM image (b) and TEM image (e) of h-Fe₂O₃@C@G with 0.1 g PVP dose (corresponding to 84 wt.% Fe₂O₃); SEM image (c) and TEM image (f) of h-Fe₂O₃@C@G with 0.5 g PVP dosage (corresponding to 70 wt.% Fe₂O₃); (d) TGA analysis of h-Fe₂O₃@C@G hybrids with different Fe₂O₃ contenting, which are recorded under air atmosphere at 10 °C min⁻¹.



Fig.S9 (a) XRD patterns of $CoO_x@C@G$ and h- $Co_3O_4@C@G$ hybrids; (b) XRD patterns of NiO_x@C@G and h-NiO_x@C@G hybrids; (c) TEM image of h-FeNiO_x@C@G; (d) XRD pattern and element analysis (inset) of h-FeNiO_x@C@G hybrid.



Fig.S10 (a) TGA analysis of FeNiO_x@C@G hybrid in air at the heating rate of 10 °C min⁻¹; (b) TEM image of h-FeNiO_x@C@G hybrid synthesized by air oxidation at 310 °C, in which many pores created by over oxidation of carbon could be found.



Fig.S11 Discharge/charge voltage profiles of G anode at 1A g⁻¹.



Fig.S12 Cycling stability of h-Fe₂O₃@C@G anodes with 84 wt.% Fe₂O₃ content and with 70 wt.% Fe₂O₃ content at 1 A g⁻¹.



Fig.S13 Nyquist plots of Fe₂O₃ NPs, Fe₂O₃/G and h-Fe₂O₃@C@G anodes.



Fig.S14 TEM image of cycled h-Fe₂O₃@C@G anode.



Fig.S15 (a) Discharge/charge voltage profiles of h-NiO_x@C@G anode, and (b) TEM image of h-NiO_x@C@G hybrid, in which few solid sphere or big particles could be found.



Fig.S16 Cycling stability (a) at 0.5 A g^{-1} and rate capability (b) of the h-FeNiO_x@C@G anode.

Active materials	Electrochemical properties	Reference
h-Fe ₂ O ₃ @C@G	430-1163 mAh g ⁻¹ at 0.5-15 A g ⁻¹ ,	This work
	724 mAh g ⁻¹ for 500 cycles at 5 A g ⁻¹	
3D Fe ₂ O ₃ /N-doped	420-1140 mAh g ⁻¹ at 0.2-6.0 A g ⁻¹ ,	Ref [1]
graphene	11200 mAh g ⁻¹ for 500 cycles at 0.50 A g ⁻¹	
Bubble-nanorod	491-913 mAh g ⁻¹ at 0.5-5 A g ⁻¹ ,	Ref [2]
structured Fe ₂ O ₃ /C	824 mAh g^{-1} for 300 cycles at 1 A g^{-1}	
nanofibers		
Yolk-shell FeO _x @C	370-843 mAh g ⁻¹ at 0.2-4 A g ⁻¹	Ref [3]
structure		
Iron oxide-rGO	~500-1050 mAh g ⁻¹ at 0.1-1.6 A g ⁻¹	Ref [4]
Porous iron oxide	615 mAh g ⁻¹ at 1.86 A g ⁻¹	Ref [5]
ribbons -graphene	1046 mAh g ⁻¹ for 130 cycles at 0.074 A g ⁻¹	
α -Fe ₂ O ₃ /graphene	615 mAh g ⁻¹ at 5 A g ⁻¹	Ref [6]
	1046 mAh g ⁻¹ for 130 cycles at 0.074 A g ⁻¹	
Fe ₃ O ₄ /CN _x /rGO	450-1110 mAh g ⁻¹ at 0.1-10 A g ⁻¹ ,	Ref [7]
	590 mAh g^{-1} for 500 cycles at 5 A g^{-1}	
Hollow	420-900 mAh g ⁻¹ at 0.2- 2 A g ⁻¹ ,	Ref [8]
Fe ₃ O ₄ @graphene	940 mAh g ⁻¹ for 50 cycles at 0.2 A g ⁻¹	
Fe ₃ O ₄ /CNTs	150-850 mAh g ⁻¹ at 0.15-2.4 A g ⁻¹ ,	Ref [9]
@graphene	680 mAh g ⁻¹ for 100 cycles at 0.2 A g ⁻¹	
2D G@Fe ₃ O ₄ @C	550-900 mAh g ⁻¹ at 0.2-0.5 A g ⁻¹	Ref [10]
Fe ₃ O ₄ /graphene	629-913 mAh g ⁻¹ at 0.1-2 A g ⁻¹	Ref [11]
Fe ₃ O ₄ @graphene	410-740 mAh g ⁻¹ at 0.3-1.0 A g g ⁻¹ ,	Ref [12]
	1048 mAh g ⁻¹ for 90 cycles at 0.1 A g ⁻¹	

Table S1 A comparison of the electrochemical properties between $h-Fe_2O_3@C@G$ anodes and other iron oxide-graphene electrodes

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