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Remarkable Enhancement of the Electrochemical Properties of Co₃O₄ Nanowire

Array by In Situ Surface Derivation of an Amorphous Phosphate Shell Wenhui Li,¹ Mingyue Chen,¹ Wenhao Ma, Pengcheng Qi, Wanjun Yang, Shiyu Wang, Yu Lu and Yiwen Tang*

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Fig S1. (a-b) Low and high magnification TEM images of Co₃O₄@Co-Pi, (c) EDS spectrum of Co₃O₄@Co-Pi.



Fig S2. (a) XPS spectra of Co 2p for Co_3O_4 . (b) FT-IR spectra of Co_3O_4 and $Co_3O_4@Co-Pi$.



Fig S3. Illustration of the proposed morphology evolution mechanism of Co₃O₄@Co-

Pi.



Fig S4. N_2 adsorption-desorption isotherms with the corresponding pore size distribution (a) and (b): Co_3O_4 ; (c) and (d): Co_3O_4 @Co-Pi.



Fig S5. (a) XRD patterns of $Co_3O_4@Co_2P_4O_{12}$ -400, $Co_3O_4@Co_2P_4O_{12}$ -500, and $Co_3O_4@Co_2P_4O_{12}$ -600 powders removed from the nickel foam. (b-d) SEM images of $Co_3O_4@Co_2P_4O_{12}$ -400, $Co_3O_4@Co_2P_4O_{12}$ -500 and $Co_3O_4@Co_2P_4O_{12}$ -600.



Fig S6. (a) CV curves of $Co_3O_4@Co_2P_4O_{12}$ electrode with different annealing temperature at 10 mV s⁻¹. (b-c) TEM image, HRTEM image $Co_3O_4@Co_2P_4O_{12}$ -500. (d) Elemental mapping images of $Co_3O_4@Co_2P_4O_{12}$ -500.



Fig S7. (a) Specific capacitance versus various current densities of Co_3O_4 , Co_3O_4 @Co-Pi, and Co_3O_4 @Co_P4O_12. (b) CV curves of Co_3O_4 @Co-Pi electrode collected at different scan rate. (c) GCD curves of Co_3O_4 @Co-Pi electrode collected at different current density. (d) The voltage drop of three materials at a current density of 10 A g⁻¹.



Fig S8. Schematic illustration of the phosphating process with different temperature and time for fabricating core-shell Co₃O₄@Co-Pi nanowires array.



Fig S9. SEM images of Co_3O_4 @Co-Pi with different phosphating time for 250 °C, (a) 1 h, (b) 2 h, (c) 3 h. (d) XRD patterns of Co_3O_4 @Co-Pi prepared under 200 and 300°C, respectively, for 2 h.



Fig S10. SEM images of Co_3O_4 @Co-Pi with different phosphating temperature for 2 h: (a) 200°C; (b) 250°C; (c) 300°C. (d) XRD patterns of Co_3O_4 @Co-Pi prepared under 200 and 300°C, respectively, for 2 h.



Fig S11. Electrochemical characterization of Co₃O₄@Co-Pi: (a) CV curves of different phosphating time; (b) GCD plots; (c) CV curves of different phosphating temperature; (d) GCD plots.



Fig S12. (a) CV curves of activated carbon. (b) CV curves at 50 mV/s of $Co_3O_4@Co-$

Pi//AC supercapacitor.

Type of electrode	Specific (areal)	Rate	Capacitance retention	Ref.
	capacitance	capability		
Co ₃ O ₄ @Co-Pi core-	$1692 \text{ F g}^{-1} \text{ at}$	56.1% from 1	86% after 6,000 cycles	This
shell hybrid	of 1 A g ⁻¹	to		work
nanostructure		10 A g^{-1}		
RuO ₂ /Co ₃ O ₄ nanosheets	905 F g ⁻¹ at 1 A	78% from 1 to	96% after 5,000 cycles	[1]
	g ⁻¹	40 A g^{-1}		
layered Co ₃ O ₄	265 F g ⁻¹ at 1 A	64.6% from 1	89.6% after 1,000	[2]
	g-1	to 6 A g^{-1}	cycles	
Co ₃ O ₄ nanowires	746 F g ⁻¹ at 0.6	No data	86% after 500 cycles	[3]
	A g ⁻¹			
α -Co(OH) ₂ /Co ₃ O ₄	583 F g ⁻¹ at 1 A	No data	No obvious specific	[4]
nanorods	g ⁻¹		capacitance loss after	
			2000 cycles	
Co ₃ O ₄ @MnO ₂	560 F g ⁻¹ at 0.2	54.5% from	No negligible	[5]
	A g ⁻¹	0.2 to 10 A g^{-1}		
Co ₃ O ₄ nanoparticles	370 F g ⁻¹ at 0.5	No data	No negligible	[6]
	A g ⁻¹			
plate-like Co ₃ O ₄	393.6 F g ⁻¹ at	No data	96.5% after 500 cycles	[7]
	1 A g ⁻¹			
ultralayered Co ₃ O ₄	548 F g ⁻¹ at	59.4% from 4	98.5% after 2000 cycles	[8]
	4 A g ⁻¹	to 32 A g^{-1}		
Hollow Co ₃ O ₄ nanowire	599 F g ⁻¹ at	73.3% from 2	No negligible	[9]
arrays	2 A g ⁻¹	to 40 A g^{-1}		
Co ₃ O ₄ /GO	157.7 F g ⁻¹ at	51.8% from	70% after 4000 cycles	[10]
	0.1 A g ⁻¹	0.1 to 2 A g^{-1}		
flower-like NiO-Co ₃ O ₄	1190 F g ⁻¹ at	No data	99% after 5000 cycles	[11]
	4 A g ⁻¹			
PCO NWAs	1716 F g ⁻¹ at 5	No negligible	85% after 1,0000 cycles	[12]
	$mV s^{-1}$			

Table S1. Electrochemical performance obtained from Co₃O₄ based electrodes.

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