Supporting Information

Rationally Designed Hierarchical MnO₂@NiO Nanostructures for Improved Lithium Ion Storage

Nana Wang^a, YanjunZhai^a, Xiaojian Ma^a, and Yitai Qian*^{a,b}

^aKey Laboratory of Colloid and Interface Chemistry, Ministry of Education, School of Chemistry and Chemical Engineering, Shandong University, Jinan 250100, China. Email: <u>qianyt@sdu.edu.cn</u>.

^bHefei National Laboratory for Physical Science at Microscale and Department of

Chemistry, University of Science and Technology of China, Hefei, 230026, China

Experimental section:

Synthesis of MnO_2: MnO₂ nanorods have been prepared by annealing obtained MnOOH nanorods under air at 400 °C for 2 h.

Synthesis of NiO: Ni(NO₃) $_2$ ·6H₂O (1.5 mmol) and urea (3 mmol) were dissolved into mixture of 20 mL deionized water and 20 mL ethanol. Then, the solution was transferred into Teflon autoclave and heated to 120 °C for 6h. The substrate was collected, washed with deionized water and dried at 60 °C overnight. The dry composite was then heated to 400 °C for 2 h in air to obtain NiO.



Figure S1. SEM image of MnOOH nanorods.

	Elements	Weight%	Atom%
0	С	0.47	1.26
	0	33.72	63.30
	Mn	51.09	27.91
	Ni	14.72	7.53
Ni	Mn	Ni ANi	
0 1 2 3 4	5 6	7 9	9 10

Figure S2. EDX spectra of MnO₂@NiO.



Figure S3. (a) XRD patterns and (b-d) TEM images of MnO₂ nanorods.



Figure S4. Nitrogen adsorption/desorption isotherms of MnO_2 nanorods.



Figure S5. (a) XRD patterns and (b-d) TEM images of NiO flowers.



Figure S6. Nitrogen adsorption/desorption isotherms and pore size distribution (inset) of NiO.



Figure S7. CV curves of the (a) MnO_2 and (b) NiO electrodes at a scanning rate of 0.1 mV s⁻¹ in the range of 0.01 - 3 V.