

Supporting information

Molten salt strategy and plasma technology induced MnO₂ with oxygen vacancy for high performance Zn-ions battery

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Calculation

According to the reported literature^{1, 2}, the current (i) and scan rate (v) in CV curves have relationships with equation:

$$i = av^b \quad (1)$$

$$\lg(i) = b\lg(v) + \lg(a) \quad (2)$$

where i is peak current, a correspond to constant, v is the scan rate and b is equaled to the slope of the $\lg(v)$ - $\lg(i)$ plots. If b approximately equals to 0.5, suggests the diffusion process is dominant, while a capacitive behavior will dominate when b is about 1³.

The contribution ratios of the two processes at scan rates from 0.1 to 0.5 mV s⁻¹ can be calculated by equations (3):

$$i(v) = k_1v + k_2v^{1/2} \quad (3)$$

where k_1v and $k_2v^{1/2}$ corresponds to the current contributions originating from the surface capacitive effects and the diffusion-controlled insertion processes, respectively. By determining the values of k_1 and k_2 , we can further analysis contribution ratios between diffusion-controlled process and capacitive effects at different scan rates. The Eq. (3) can be changed as follows:

$$i(v)/v^{1/2} = k_1v^{1/2} + k_2 \quad (4)$$

The values of k_1 and k_2 can be obtained from the slope and y intercept of the linear fit of the plot of $i(v)/v^{1/2}$ versus $v^{1/2}$, respectively. The capacitive and diffusion-controlled currents assistance to the total charge was separately calculated by inserting the values of k_1 and k_2 into the Eq. (4).

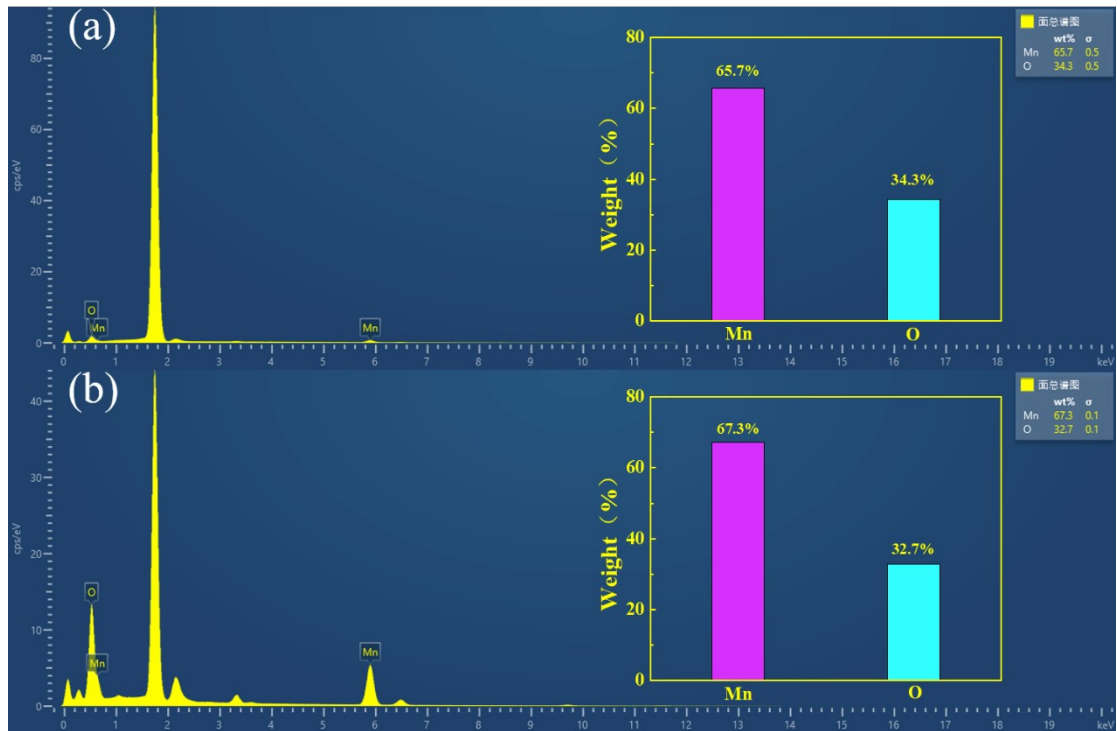


Figure S1. Elemental composition analysis of (a) MnO_2 and (b) MnO_{2-3} .

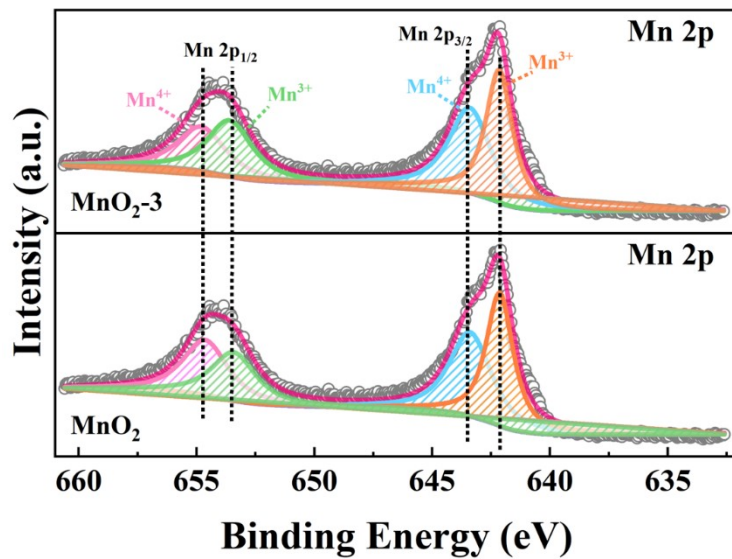


Figure S2. Mn 2p XPS spectra comparison of MnO_2 and MnO_{2-3} .

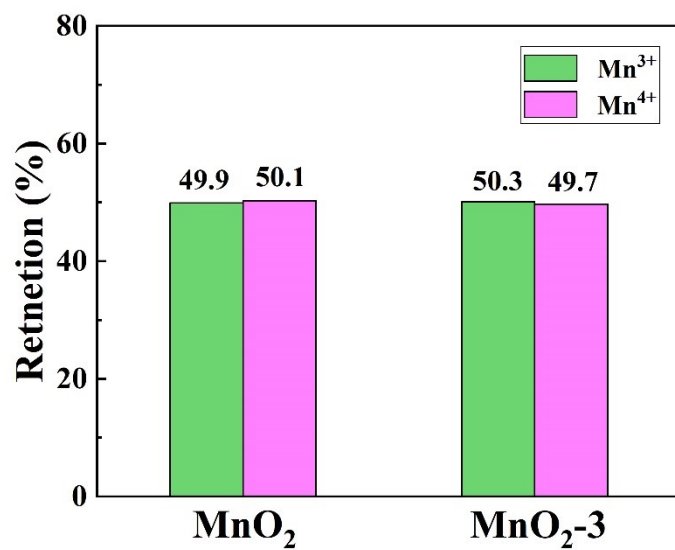


Figure S3. quantitatively statistics of the percentage of Mn³⁺ and Mn⁴⁺ in the samples.

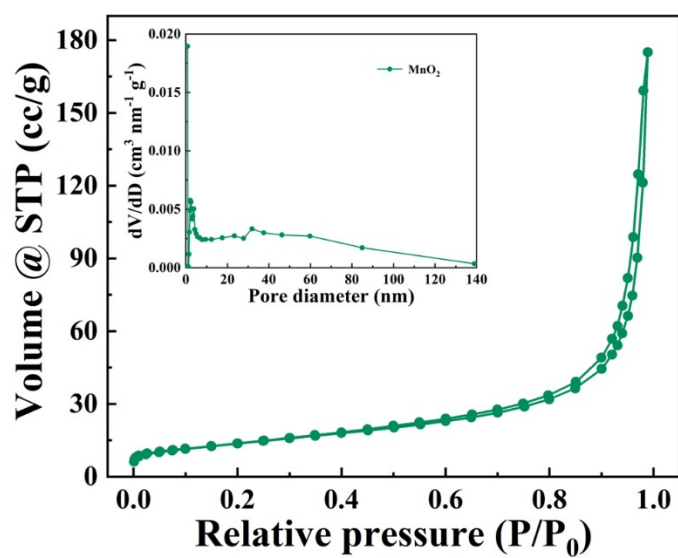


Figure S4. Nitrogen adsorption-desorption isotherm and pore distribution of MnO₂.

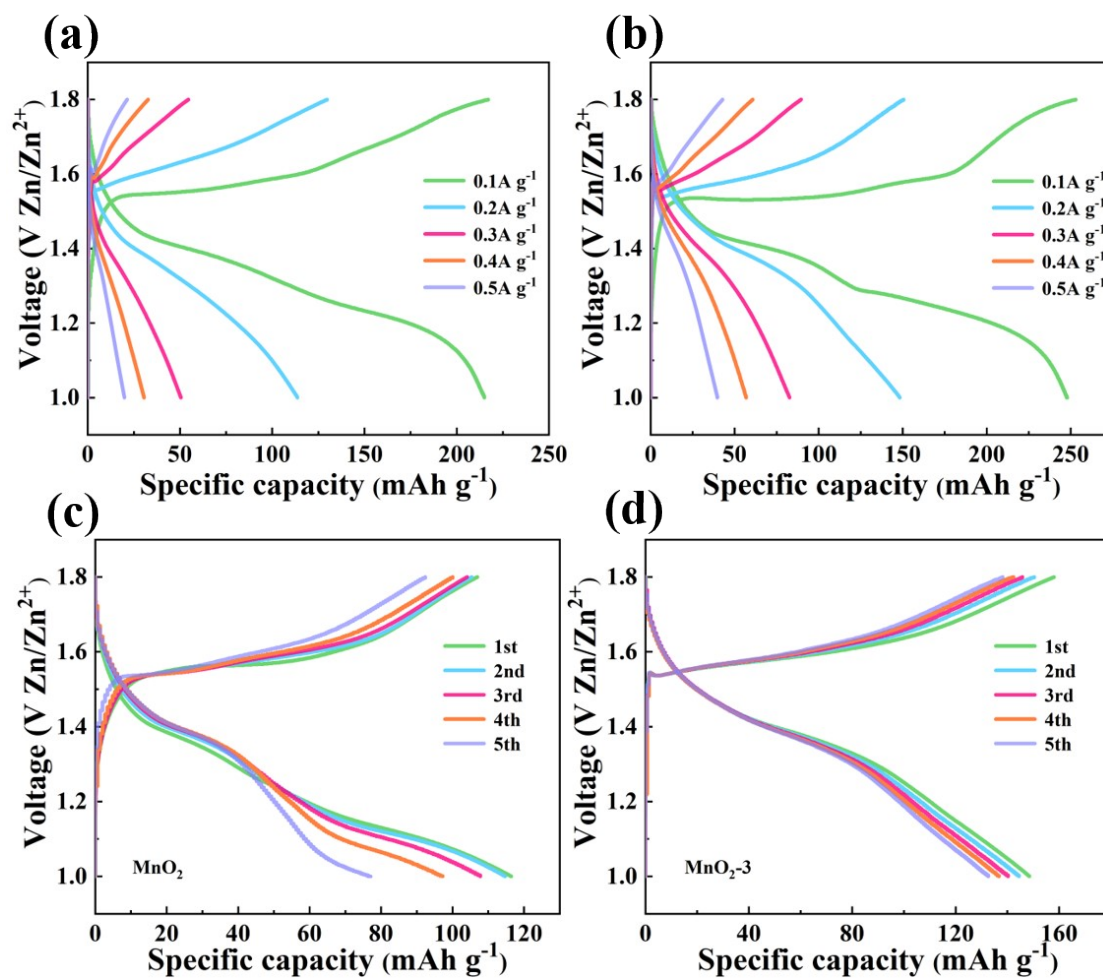


Figure S5. GCD curves at current densities from 0.1 to 0.5 A g⁻¹ of (a) MnO₂ and (b) MnO₂-3.

GCD profiles in the initial 5 cycles of (c) MnO₂ and (d) MnO₂-3.

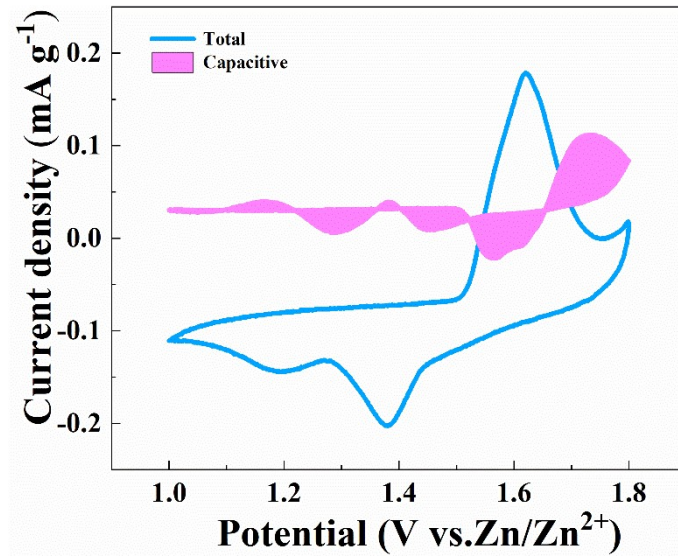


Figure S6. Capacitive and diffusion-controlled contribution analysis of the CV curves of MnO₂-3.

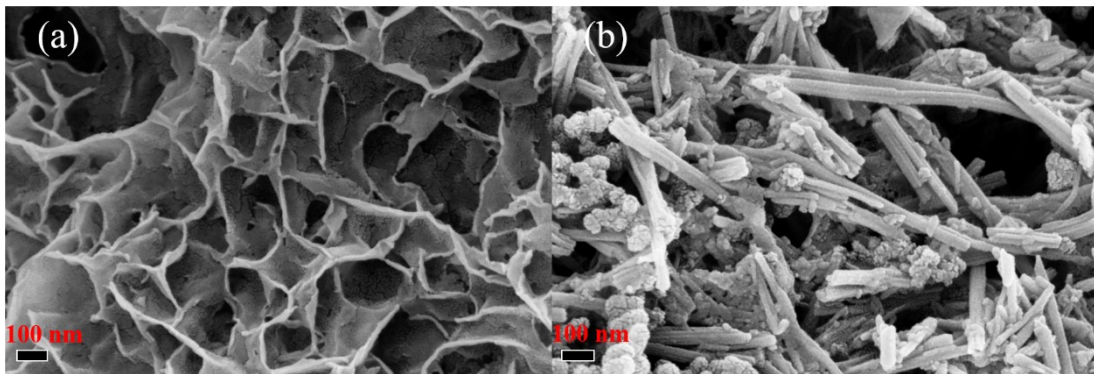


Figure S7. SEM images of (a) MnO₂ and (b) MnO₂-3 after 10 cycles.

Table S1. Comparison of electrochemical performance of other materials

Materials	Specific capacity (mA h g⁻¹)	Cycle number	Capacity retention	Referenc e
δ -MnO ₂	170 (0.1A g ⁻¹)	100 (0.1 A g ⁻¹)	83%	4
α -MnO ₂ @PPy	137 (0.1A g ⁻¹)	100 (0.1 A g ⁻¹)	58%	5
α -MnO ₂	180 (42 mA g ⁻¹)	30 (42 mA g ⁻¹)	76%	6
δ -MnO ₂	252 (82 mA g ⁻¹)	100 (83 mA g ⁻¹)	44%	7
δ -MnO ₂	108 (12.3 mA g ⁻¹)	125 (12.3 mA g ⁻¹)	63%	8
UCT-1-250	222 (0.1A g ⁻¹)	200 (0.1A g ⁻¹)	57%	9
γ - MnO ₂	250 (0.5 mA cm ⁻²)	40 (0.5 mA cm ⁻²)	63%	10
This work	252 (0.1A g⁻¹)	100 (0.2A g⁻¹)	81%	

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