

Supplementary Material (ESI) for Physical Chemistry Chemical Physics
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Supporting Information

**Enhancement of electrocapacitive performance of manganese dioxide by
introduction of microporous carbon sphere network**

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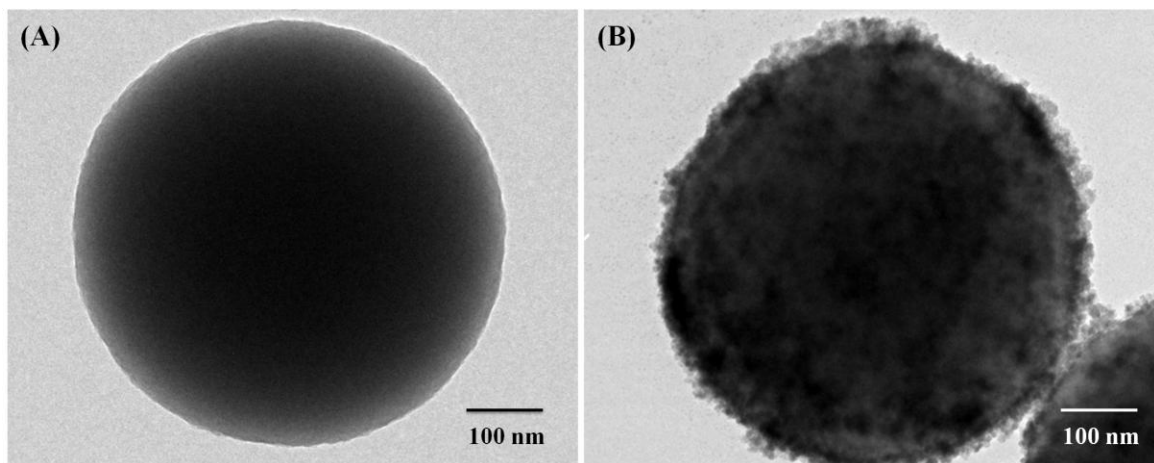


Fig. S1 TEM images of (A) MCS and (B) α -MnO₂·nH₂O/25 wt% MCS.

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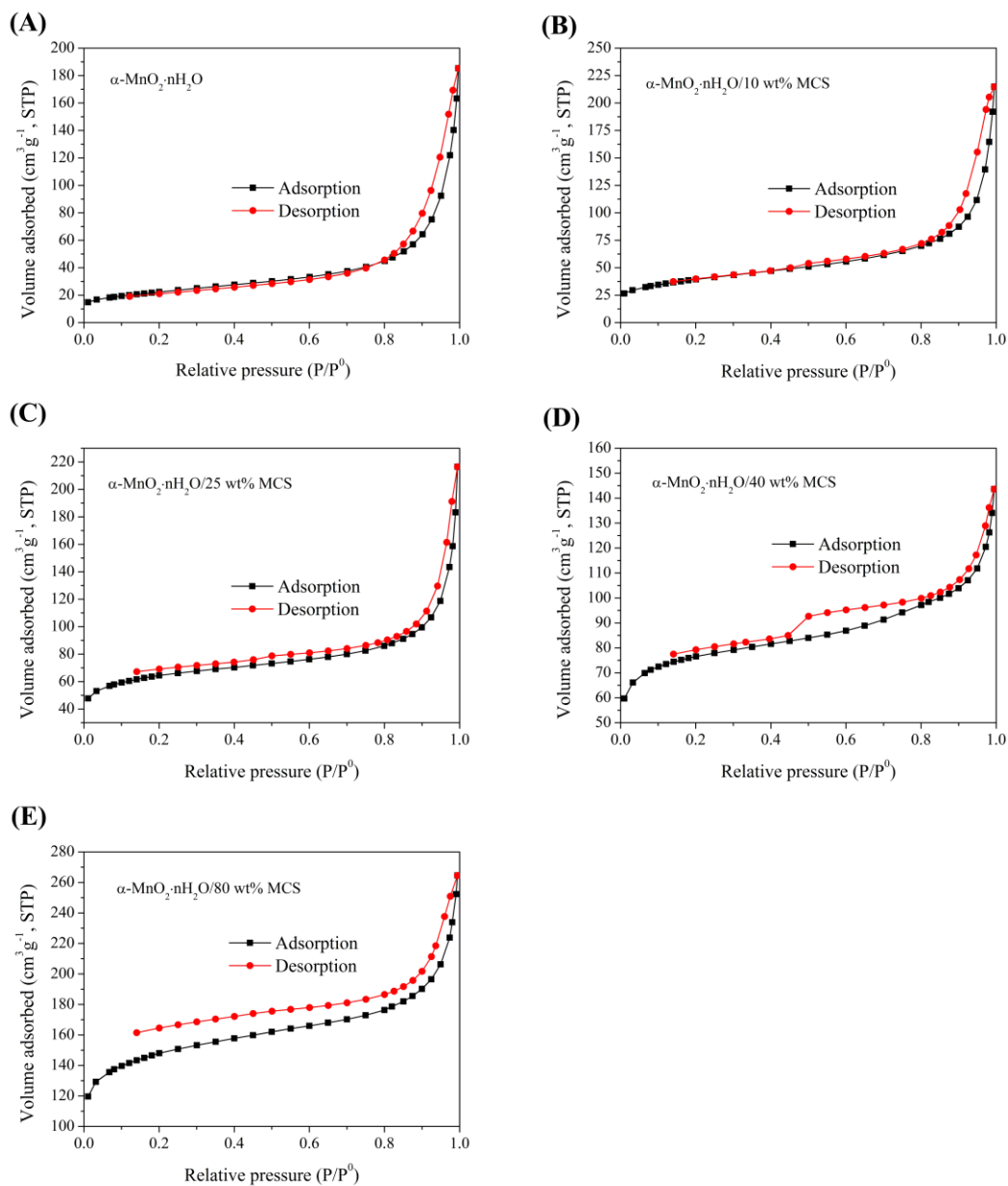


Fig. S2 Nitrogen adsorption/desorption isotherms of (A) $\alpha\text{-MnO}_2 \cdot n\text{H}_2\text{O}$, (B) $\alpha\text{-MnO}_2 \cdot n\text{H}_2\text{O}/10 \text{ wt}\% \text{ MCS}$, (C) $\alpha\text{-MnO}_2 \cdot n\text{H}_2\text{O}/25 \text{ wt}\% \text{ MCS}$, (D) $\alpha\text{-MnO}_2 \cdot n\text{H}_2\text{O}/40 \text{ wt}\% \text{ MCS}$, and (E) $\alpha\text{-MnO}_2 \cdot n\text{H}_2\text{O}/80 \text{ wt}\% \text{ MCS}$.

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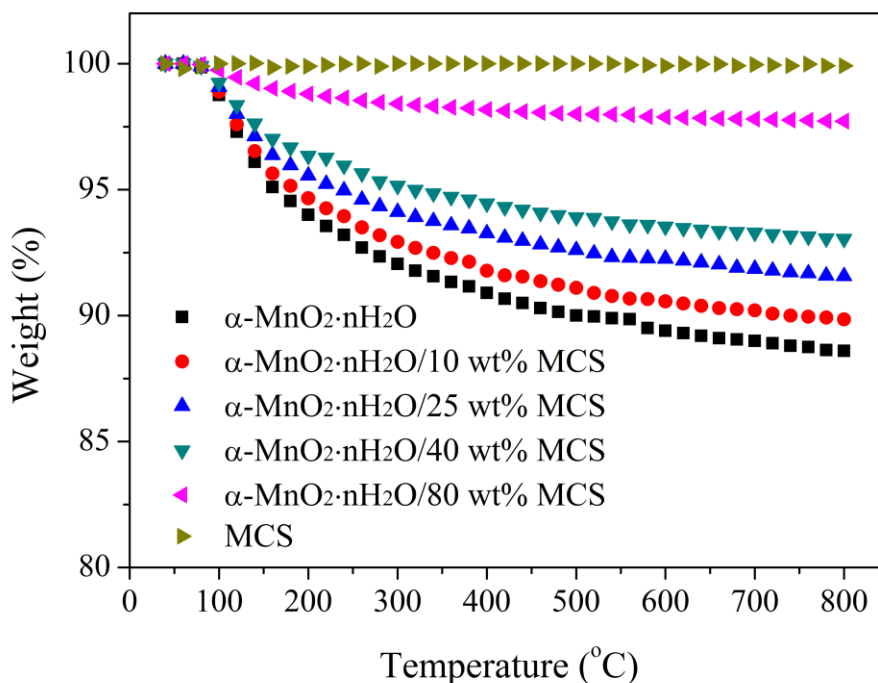


Fig. S3 TGA results for all the samples.

Fig. S3 shows the TGA results for all the samples. For MCS, because it is the final carbon product of the precursor after heat treated at 1200 °C for 5 h in the protection of nitrogen, so the weight of MCS is stable and no weight loss can be observed in the TGA test under nitrogen atmosphere. For α -MnO₂·nH₂O, the most weight loss occurs between 100 °C to 450 °C, and is about 10 wt %, which is attributed to the loss of the water. Since water evolution is continuous and mostly above 100 °C, that indicates that it is not adsorbed water, but water in the oxide structure. For the composite, the weight loss is due to the dehydration of the crystal water of α -MnO₂·nH₂O component. The sample containing more α -MnO₂·nH₂O component exhibits more weight loss. The temperature

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of the initial weight loss for all the composite samples is same. The above result demonstrates that the as-prepared MnO_2 is hydrated.

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Table S1 Summary of the BET surface area for the prepared samples

	Samples				
	α -MnO ₂ ·nH ₂ O	α -MnO ₂ ·nH ₂ O/10	α -MnO ₂ ·nH ₂ O/25	α -MnO ₂ ·nH ₂ O/40	α -MnO ₂ ·nH ₂ O/80
		wt% MCS	wt% MCS	wt% MCS	wt% MCS
BET surface area (m ² g ⁻¹)	79.6	135.9	223.5	261.2	505.5