

Supplementary Information

Facile hydrothermal synthesis and electrochemical properties of orthorhombic LiMnO_2 cathode materials for rechargeable lithium batteries

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Figure captions

Fig. S1. XRD patterns of products synthesized in 0.1 mol L⁻¹ MnCl₂, 0.6 mol L⁻¹ LiOH, 0.1 mol/L NaClO with (a) and without (b) EDTA (0.1 mol L⁻¹) at 180 °C for 24 h.

Fig. S2. XRD patterns of the synthesized pure-phased and aluminum-doped *o*-LiMnO₂: (a) M₀, (b) M₃, (c) M₆, (d) M₉, and (e) M₁₂.

Fig. S3. Particle size distributions of M₀ (a), M_m (b), M₃ (c), M₆ (d), M₉ (e) and M₁₂ (f).

Fig. S4. TEM images of M₀ (a), M₃ (b), M₆ (c), M₉ (d), M₁₂ (e).

Fig. S5. XRD patterns of products synthesized under different conditions: (a) 0.1 mol L⁻¹ MnCl₂, 0.6 mol L⁻¹ LiOH, 0.1 mol L⁻¹ NaClO, 0.1 mol L⁻¹ EDTA, 180 °C, 24 h; (b) heat treatment of (a) in air at 300 °C for 2 h; (c) heat treatment of (a) in air at 600 °C for 2 h.

Fig. S6. Discharge specific capacities vs. cycle number with current density of 100 mA g⁻¹ for M₀, M_m, and M₆₀₀ after heat-treatment at 110 °C for 12 h.

1. The effect of EDTA on product composition

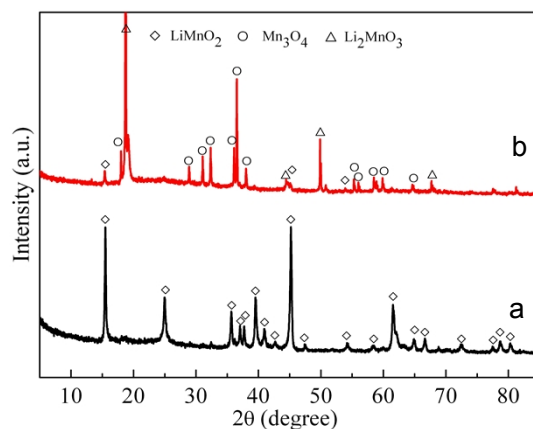


Fig. S1. XRD patterns of products synthesized in 0.1 mol L^{-1} MnCl_2 , 0.6 mol L^{-1} LiOH , 0.1 mol/L NaClO with (a) and without (b) EDTA (0.1 mol L^{-1}) at 180 °C for 24 h.

2. The preparation of Al doped o -LiMnO₂

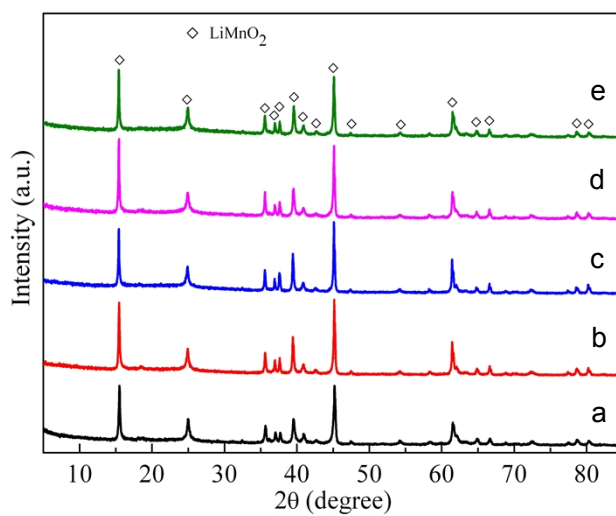


Fig. S2. XRD patterns of the synthesized pure-phased and aluminum-doped o -LiMnO₂: (a) M₀, (b) M₃, (c) M₆, (d) M₉, and (e) M₁₂.

3. The particle size distributions of the as-obtained samples

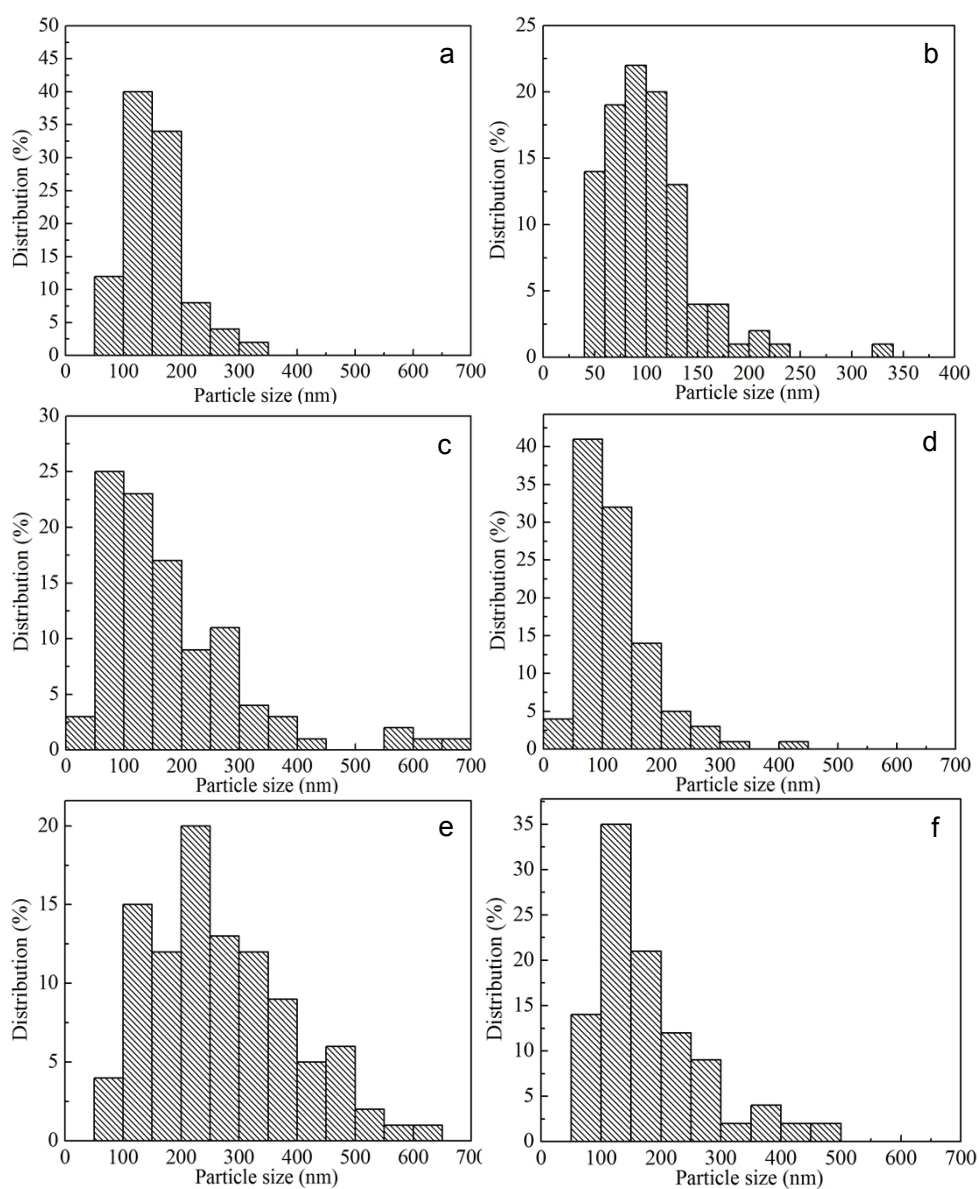


Fig. S3. Particle size distributions of M₀ (a), M_m (b), M₃ (c), M₆ (d), M₉ (e) and M₁₂ (f).

4. The micromorphology and particle size of pure and Al doped *o*-LiMnO₂

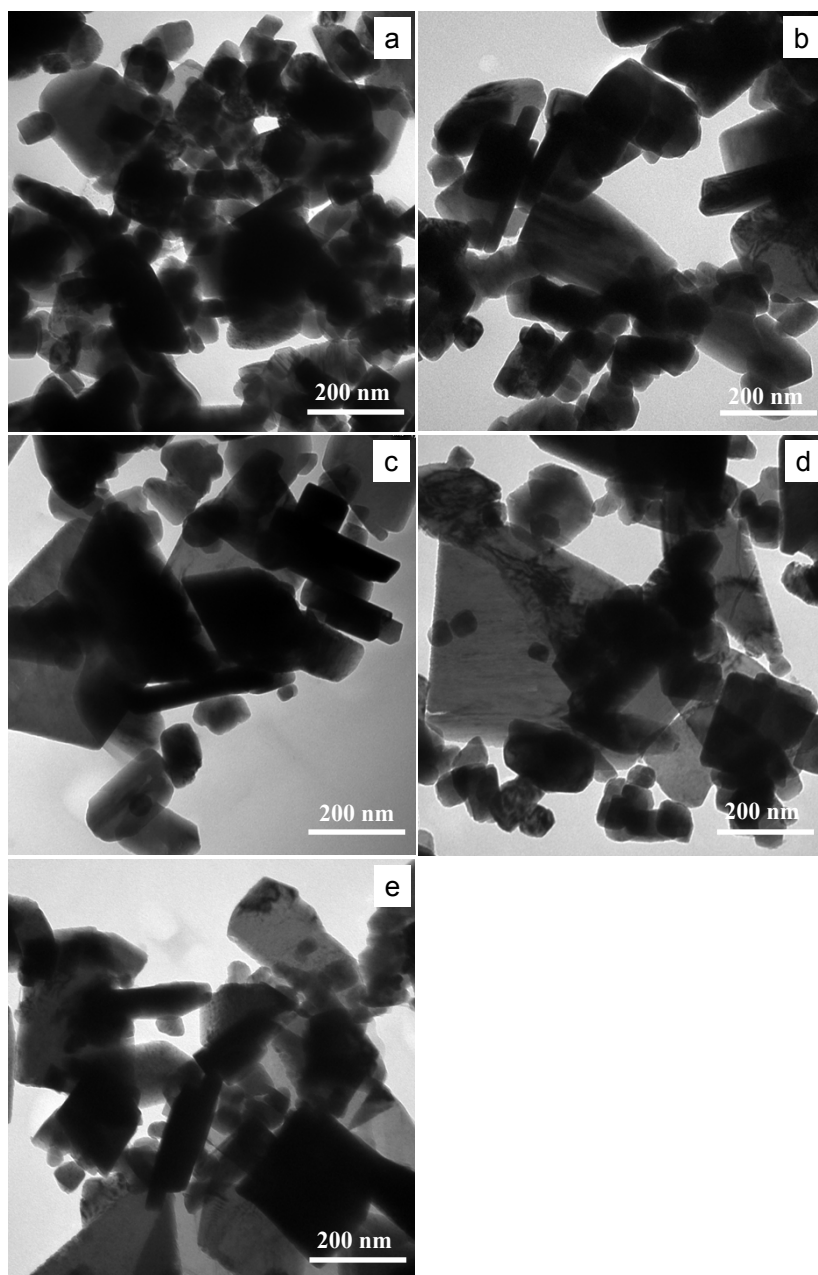


Fig. S4. TEM images of M_0 (a), M_3 (b), M_6 (c), M_9 (d), M_{12} (e).

5. The heat treatment of as-obtained α -LiMnO₂ in air

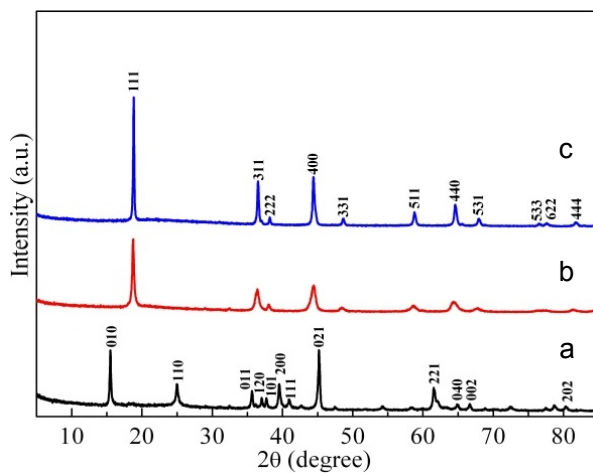


Fig. S5. XRD patterns of products synthesized under different conditions: (a) 0.1 mol L⁻¹ MnCl₂, 0.6 mol L⁻¹ LiOH, 0.1 mol L⁻¹ NaClO, 0.1 mol L⁻¹ EDTA, 180 °C, 24 h; (b) heat treatment of (a) in air at 300 °C for 2 h; (c) heat treatment of (a) in air at 600 °C for 2 h.

6. The electrochemical performance of the as-obtained ρ -LiMnO₂ and LiMn₂O₄

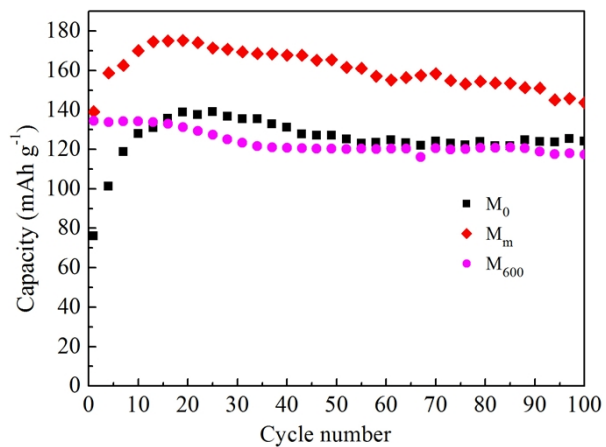


Fig. S6. Discharge specific capacities vs. cycle number with current density of 100 mA g⁻¹ for M_0 , M_m , and M_{600} after heat-treatment at 110 °C for 12 h.