Single Nanorod Devices for Battery Diagnostics: A Case Study on LiMn$_2$O$_4$

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ABSTRACT

This paper presents single nanostructure devices as a powerful new diagnostic tool for batteries with LiMn$_2$O$_4$ nanorod materials as an example. LiMn$_2$O$_4$ and Al-doped LiMn$_2$O$_4$ nanorods were synthesized by a two-step method that combines hydrothermal synthesis of $\beta$-MnO$_2$ nanorods and a solid state reaction to convert them to LiMn$_2$O$_4$ nanorods. $\lambda$-MnO$_2$ nanorods were also prepared by acid treatment of LiMn$_2$O$_4$ nanorods. The effect of electrolyte etching on these LiMn$_2$O$_4$-related nanorods is investigated by both SEM and single-nanorod transport measurement, and this is the first time that the transport properties of this material have been studied at the level of an individual single-crystalline particle. Experiments show that Al dopants reduce the dissolution of Mn$^{3+}$ ions significantly and make the LiAl$_{0.1}$Mn$_{1.9}$O$_4$ nanorods much more stable than LiMn$_2$O$_4$ against electrolyte etching, which is reflected by the magnification of both size shrinkage and conductance decrease. These results correlate well with the better cycling performance of Al-doped LiMn$_2$O$_4$ in our Li-ion battery tests: LiAl$_{0.1}$Mn$_{1.9}$O$_4$ nanorods achieve 96% capacity retention after 100 cycles at 1C rate at room temperature, and 80% at 60 °C, whereas LiMn$_2$O$_4$ shows worse retention of 91% at room temperature, and 69% at 60 °C. Moreover, temperature-dependent I–V measurements indicate that the sharp electronic resistance increase due to charge ordering transition at 290 K does not appear in our LiMn$_2$O$_4$ nanorod samples, suggesting good battery performance at low temperature.

Rechargeable batteries such as lithium ion batteries are important energy storage devices for portable electronic devices, power tools, and electrical vehicles. Anodes and cathodes in the existing lithium ion batteries consist of particles with various sizes and shapes, conductive carbon, and polymer binders. During battery charge/discharge, electrons and ions are simultaneously inserted or extracted from battery electrodes, which is accompanied by a series of other complex processes such as structure and phase transformation, volume change, materials dissolution, and side chemical reaction with electrolyte. Various technologies have been utilized for battery diagnostics, including electrochemical impedance spectroscopy, X-ray diffraction, Raman spectroscopy, and electron microscopy inspection of ensemble electrodes. However, the heterogeneous nature of ensemble electrodes averages all information and can not provide a direct correlation of electrochemical properties with the local morphology, structure, and chemical composition. Few reports focused on single particle diagnostics, as it is hard to handle single particles with size of 0.1~10 μm. Here we exploit single nanorod devices to demonstrate a novel powerful diagnostic tool, which allows for the direct correlation of the electrochemical property with the structure on the same nanoscale particle. Single nanostructure devices fabricated with lithographic techniques have been well developed in the field of nano science but rarely applied to battery researches. This work represents the first example of single nanostructure device battery diagnostics. We use single LiMn$_2$O$_4$ nanorod devices as an example in this study although this new methodology can generally be applied to a broad range of battery electrode materials. In the future, the capability of combining single nanostructure device diagnostics with in situ electron microscopy techniques can lead to a much deeper understanding of battery operational processes.

We choose spinel LiMn$_2$O$_4$ in this study since it has the advantages of low-cost, environmental friendliness, and safety. However, the fast capacity decay at high temperature impedes its practical application. The mechanism is that Mn$^{3+}$ ions in LiMn$_2$O$_4$ can disproportionate due to the generation of HF in an LiPF$_6$-based electrolyte and be...
dissolved in the electrolyte resulting in cathode capacity loss and the attack of carbon anodes by Mn ions.22–25 Significant efforts have been made to improve the cycling performance of LiMn2O4, such as coating36–38 and doping.29–32 Moreover, various methods have been utilized to analyze the mechanism and the process of Mn dissolution, including X-ray diffraction matter and impedance spectroscopy,5,6 which can provide useful information but in an indirect way. Nevertheless, there is no study that directly track the effect of the electrolyte on the morphology or properties of a single LiMn2O4 particle. The difficulties resulting from the ensemble nature of electrodes include how to distinguish a particle in the mixed film-like electrode and how to make probes, such as contacting metal electrodes, onto such small particles (typically less than 20 µm for battery materials).

In comparison, one-dimensional nanorods and nanowires can offer the following advantages: (1) In contrast to microsized particles, it is feasible to put multiple electrodes onto them to study their transport properties.19,33 (2) The single-nanostructure devices can be characterized by a variety of electron microscopy and even in situ techniques.15,16 (3) They are single-crystalline,34,35 which provides well-defined nanoscale domains for understanding their intrinsic properties. LiMn2O4 nanorods and nanowires have been synthesized by us36 and several other groups,37,38 but their transport properties have not been studied at the single nanostructure level. In this work, single-crystalline LiMn2O4 and Al-doped LiMn2O4 nanorods were synthesized by a two-step method and λ-MnO2 nanorods were also prepared by acid delithiation of LiMn2O4 nanorods. These three kinds of nanorods offer the possibility to study the electrochemical and transport properties of LiMn2O4, such as the effect of dopants and lithium concentration. We studied how electrolyte etching affects their electrical properties and correlated them with the electrochemical performance of these materials.

The synthesis of LiAl0.1Mn1.9O4 (0 ≤ x ≤ 0.2) nanorods is a two-step approach. First, β-MnO2 nanorods were synthesized as a template by hydrothermal methods.19 Typically 8 mmol of MnSO4·H2O and 8 mmol of (NH4)2SO4 were dissolved in 10 mL of deionized water separately to form clear solutions. They were mixed together and transferred to a 45 mL Teflon-lined stainless steel vessel (Parr Co.). The vessel was sealed and heated at 150 °C for 12 h to obtain β-MnO2 nanorods. After this step, as-synthesized MnO2 nanorods were mixed and ground with lithium acetate (Sigma Aldrich) with a molar ratio of 2:1. A total of 1 mL of methanol was added to make a uniform slurry mixture. The mixture was sintered at 700 °C for 10 h under air. For Al-doped LiMn2O4 samples, a stoichiometric amount of Al(NO3)3·9H2O was added into the slurry mixture as the Al source. A series of samples were prepared with x = 0, 0.05, 0.1, and 0.2. To prepare λ-MnO2 nanorods, a method reported previously was used with a slight modification. Generally 20 mg of LiMn2O4 nanorods were delithiated in 10 mL of 0.1 M HCl for 30 min under stirring, and then the delithiated nanorods were dried at 200 °C in air for 2 h. It is believed that hydrogen is not stable inside λ-MnO2 and the valence of Mn is close to 4 according to magnetic measurement and X-ray absorption spectra.41

The composition of as-synthesized LiAl0.1Mn1.9O4 nanorods was characterized by X-ray diffraction (XRD, PANalytical X’Pert diffractometer with Cu Kα-radiation) and energy dispersive X-ray spectroscopy (EDS). The morphologies of the nanostructures were observed by a field emission scanning electron microscope (FEI XL30 Sirion SEM) and a high resolution transmission electron microscope (HR-TEM, 200 kV Philips CM20).

The electrodes for electrochemical studies were prepared by making slurry of 78 wt % active material of LiAl0.1Mn1.9O4, 15 wt % conducting carbon black, and 7 wt % polyvinylidene fluoride (PVdF) binder in N-methyl-2-pyrrolidone (NMP) as the solvent. The slurry was coated onto an aluminum foil by an applicator and then dried at 120 °C overnight. The electrochemical performance of LiMn2O4 was investigated inside a coffee bag (pouch) cell assembled in an argon-filled glovebox (oxygen and water contents below 1 and 0.1 ppm, respectively). Lithium metal foil (Alfa Aesar) was used as the anode. The typical cathode loading was 3–4 mg/cm². A 1 M solution of LiPF6 in ethylene carbonate/diethyl carbonate (EC/DEC, 1:1 v/v) (Ferro Corporation) was used as the electrolyte with a Celgard 2321 triple-layer polypropylene-based membrane as the separator. The charge/discharge cycles were performed at different rates between 3.5–4.3 V at room temperature and 60 °C using Bio-Logic VMP3 battery testers.

Devices for single-nanorod transport measurement were fabricated by electron beam lithography (EBL) on Si/SiN4 (300 nm) chips. The metal electrode consisted of 10 nm Cr/190 nm Au. The electrolyte etching experiments were performed by immersing Si/SiN4 chips with nanorod devices into 2 mL of the electrolyte mentioned above and heating to 60 °C for different amounts of time. Then the electrolyte was washed away by DEC and acetone in sequence. SEM and transport (I–V) measurements were used to characterize the effect of the electrolyte on nanorod devices.

Figure 1, panels a and b, shows SEM images of LiMn2O4 and LiAl0.1Mn1.9O4 nanorods, respectively. The length of both LiMn2O4 and LiAl0.1Mn1.9O4 nanorods is around 0.7–1.2 µm and the diameter is 100–200 nm; these nanorods are shorter in length and larger in diameter than β-MnO2 nanorods (Supporting Information, Figure S1) as a result of high temperature sintering. Figure 1c shows an SEM image of delithiated λ-MnO2 nanorods, which look similar to the LiMn2O4 nanorods. This indicates that the delithiation process does not destroy the nanorod morphology. TEM characterization of these nanorods is also presented in Figure 1. The lattice fringes of LiMn2O4 and LiAl0.1Mn1.9O4 are illustrated in Figure 1, panels d and e, respectively. The inset is the corresponding diffraction pattern. The clear lattice structure and the sharp diffraction spots reveal the single crystalline nature of these samples. EDS results of LiAl0.1Mn1.9O4 samples give a calculated Al/Mn ratio of 4.8 at. % (Supporting Information, Figure S2). However, as represented in Figure 1f, inset, the diffraction spots of λ-MnO2 nanorods are elongated, which implies strain and slight lattice distortion after acid delithiation. An XRD study
was also performed to investigate the purity of samples and the effect of Al dopant on the fine structure of these nanorods (Supporting Information, Figure S3). No impurity peaks were detected in the XRD pattern, and a systematic shift of peaks toward higher angle was observed in doped samples, suggesting a smaller lattice parameter as a consequence of Al substitution.

Since the dissolution of LiMn$_2$O$_4$ in the electrolyte is an important issue which has impeded the application of this material, we studied the effect of the electrolyte on the morphology and transport properties of these LiMn$_2$O$_4$-based nanorods. To directly observe the dissolution of material and its effect on transport properties, two metal electrodes were deposited at the two ends of the nanorods, and $I$–$V$ measurements were used to track the effect of the electrolyte on a particularly identified nanorod. A typical device is shown in Figure S4 in the Supporting Information. Furthermore, these results could be well correlated with their electrochemical performance, which will be discussed later. Figure 2a–c shows SEM images of LiMn$_2$O$_4$ nanorods immersed in the electrolyte of 1 M LiPF$_6$ in EC/DEC at 60 °C after 0, 3, and 9 h, respectively, whereas Figure 2, panels d–f and g–i, represents images of LiAl$_{0.1}$Mn$_{1.9}$O$_4$ and λ-MnO$_2$ immersed in electrolyte after the same time, respectively. It is obvious that the middle part of the LiMn$_2$O$_4$ nanorod, which was not protected by the metal electrodes, was etched. In contrast, no change occurred in LiAl$_{0.1}$Mn$_{1.9}$O$_4$ samples. Results of the λ-MnO$_2$ samples are interesting. The nanorod becomes covered by electrolyte residue or solid electrolyte interphase (SEI) layer after a period of time, which is difficult to remove. This might be a consequence of λ-MnO$_2$ electrolyte decomposition on the surface of λ-MnO$_2$, since Mn (IV) has high oxidation power. The corresponding redox potential of λ-MnO$_2$ is ∼4.2–4.3 V vs Li/Li$^+$, which is close to the potential of electrolyte oxidation.42,43 These results are based on observation of at least 5 nanorod devices for each kind of material.

In addition to the evolution of their morphology, the conductance of these nanorods also changes with time in the electrolyte. Figure 3a–c shows the evolution of $I$–$V$ curves of the same LiMn$_2$O$_4$, LiAl$_{0.1}$Mn$_{1.9}$O$_4$, and λ-MnO$_2$ nanorod devices shown in Figure 2, immersed in the electrolyte. The nonlinear behavior of the normalized conductance of nanorods in the electrolyte.

Figure 1. SEM pictures of (a) LiMn$_2$O$_4$ nanorods, (b) LiAl$_{0.1}$Mn$_{1.9}$O$_4$ nanorods, and (c) λ-MnO$_2$ nanorods. High resolution TEM images of (d) LiMn$_2$O$_4$, (e) LiAl$_{0.1}$Mn$_{1.9}$O$_4$, and (f) λ-MnO$_2$. Inset is the corresponding diffraction pattern.

Figure 2. SEM images of nanorod devices in 1 M LiPF$_6$ in the EC/DEC electrolyte: LiMn$_2$O$_4$ (a) 0, (b) 3, and (c) 9 h; LiAl$_{0.1}$Mn$_{1.9}$O$_4$ (d) 0, (e) 3, and (f) 9 h; λ-MnO$_2$ (g) 0, (h) 3, and (i) 9 h. All scale bars are 200 nm.

Figure 3. $I$–$V$ curves of nanorods in the electrolyte for different times: (a) LiMn$_2$O$_4$, (b) LiAl$_{0.1}$Mn$_{1.9}$O$_4$, and (c) λ-MnO$_2$. (d) The evolution of the normalized conductance of nanorods in the electrolyte.
The conductance decreases gradually after immersion in the electrolyte. Results are summarized as the change in conductance versus time immersed in electrolyte at 60 °C, as illustrated in Figure 3d. The conductance is normalized to its value before immersion. The conductance of LiAl0.1Mn1.9O4 decreases slightly (~20%) while that of LiMn2O4 drops dramatically (>60%). This indicates that Al-doped samples are more resistive to electrolyte etching. The reason is that Al could suppress the dissolution of Mn3⁺ in the electrolyte since the concentration of Mn3⁺ becomes smaller. As a result, LiAl0.1Mn1.9O4 should show better capacity retention than LiMn2O4 in battery charge/discharge cycling. The conductivity of λ-MnO2 decreases a lot due to the reduction by the electrolyte and thus the dissolution of Mn. On a few occasions, broken λ-MnO2 nanorods were observed and the conductance was zero. The change in conductivity is less likely to come from the effect of the electrolyte on the metal/nanorod interface, since the interface is protected by the thick metal layer itself. Moreover, if the damage at the interface is the main factor in the conductivity change, it is hard to explain the quite different behavior of these three similar materials. It is worthwhile to point out that the total mass of nanorods immersed in the electrolyte is at most on the order of tens of micrograms. As a result, the ratio of the volume of organic electrolyte to the mass of active material is at least 3 orders of magnitude larger than that in practical batteries, so the etching rate is highly magnified in our experiments, though the trend should be the same.

As indicated by the I–V measurement above, Al-doped samples should give better capacity cycling retention. To confirm this, LiAl0.1Mn2.0O4 nanorods were incorporated into a composite ensemble electrode, as described in the experimental section. Typical discharge curves of samples with \( x = 0 \) and 0.1 at the cycling rate of C/5 (1C = 148 mA/g) are plotted in Figure 4a. The sample without doping exhibits the well-known two plateau behavior, but the doped samples show vague plateaus as a result of doping.
Figure 5. (a) Current–voltage curves of LiMn$_2$O$_4$, Li$_{0.1}$Mn$_{1.9}$O$_4$, and λ-MnO$_2$ nanorods. (b) The dependence of the conductivity of nanorods on temperature from 260 to 350 K.

Table 1. Activation Energy of Nanorods Based on Different Fitting Models

<table>
<thead>
<tr>
<th>Material</th>
<th>$\exp(-E_A/kT)/eV$</th>
<th>$T^{-1}\exp(-E_A/kT)/eV$</th>
<th>$T^{-3/2}\exp(-E_A/kT)/eV$</th>
</tr>
</thead>
<tbody>
<tr>
<td>LiMn$_2$O$_4$</td>
<td>0.28</td>
<td>0.31</td>
<td>0.32</td>
</tr>
<tr>
<td>Li$<em>{0.1}$Mn$</em>{1.9}$O$_4$</td>
<td>0.27</td>
<td>0.29</td>
<td>0.29</td>
</tr>
<tr>
<td>λ-MnO$_2$</td>
<td>0.15</td>
<td>0.18</td>
<td>0.19</td>
</tr>
</tbody>
</table>

might be a result of a closer Mn–Mn distance in λ-MnO$_2$ compared to LiMn$_2$O$_4$ and Li$_{0.1}$Mn$_{1.9}$O$_4$. Though a two-electrode configuration introduces contact resistance to the experiment, the measured conductivity of LiMn$_2$O$_4$ is still larger than previous results, implying that the contact resistance is not important in the measurement and the nanorods measured have good crystallinity. Ionic conductivity is considered not to be dominant (<15%) in the conductivity of LiMn$_2$O$_4$, so the as-measured $I$–$V$ data counts for electronic conductivity of these samples. Figure 5b indicates the temperature-dependent conductivity of these three kinds of nanorods. They all behave like semiconductors from 260 to 350 K. The sharp drop in conductivity at 290 K induced by the charge-ordering transition in LiMn$_2$O$_4$ was not observed in our devices, suggesting that our materials would have good battery performance at low temperatures from the perspective of electronic conductivity by the data of the temperature-dependent conductivity, the activation energy of these materials was calculated. Three models were used: simply $\sigma \propto \exp(-E_A/kT)$, adiabatic Arrhenius relationship $\sigma \propto T^{-1} \exp(-E_A/kT)$, and nonadiabatic Arrhenius relationship $\sigma \propto T^{-3/2} \exp(-E_A/kT)$, and the result is represented in Table 1. The activation energy of LiMn$_2$O$_4$ is slightly higher than that of Li$_{0.1}$Mn$_{1.9}$O$_4$ but almost twice that of λ-MnO$_2$. This result is consistent with the hopping model of manganese oxide-based materials, which predicts that smaller Mn–Mn distance leads to a lower activation barrier. No matter which model is used, the values in this table are smaller than previous results. This also suggests that our LiAl$_{0.1}$Mn$_{1.9}$O$_4$ nanorods have good crystallinity and the single nanorod measurement is free of interparticle hopping.

In conclusion, LiMn$_2$O$_4$ and Al-doped LiMn$_2$O$_4$ nanorods were synthesized by a method combining hydrothermal synthesis of β-MnO$_2$ nanorods and sintering with lithium acetate and aluminum nitrate. λ-MnO$_2$ nanorods were also obtained by delithiating the LiMn$_2$O$_4$ nanorods with acid. For the first time, the effect of the electrolyte on LiMn$_2$O$_4$ is tracked at a single-particle level by both SEM characterization and electronic transport measurement. Al dopants reduce the dissolution of Mn$^{3+}$ ions, and make Li$_{0.1}$Mn$_{1.9}$O$_4$ nanorods more stable than LiMn$_2$O$_4$ against electrolyte etching. Li$_{0.1}$Mn$_{1.9}$O$_4$ nanorods shows high stability during fast cycling, with 96% capacity retention after 100 cycles at 1C rate at room temperature, and 80% at 60 °C. Moreover, the temperature-dependent $I$–$V$ measurement indicates that LiMn$_2$O$_4$, Li$_{0.1}$Mn$_{1.9}$O$_4$ and λ-MnO$_2$ nanorods all behave like semiconductors and the activation energy of λ-MnO$_2$ is just half of that of LiMn$_2$O$_4$ and Li$_{0.1}$Mn$_{1.9}$O$_4$, which is consistent with the rule that smaller Mn–Mn distance leads to easier hopping among Mn ions. The sharp drop in conductivity at 290 K of LiMn$_2$O$_4$ has not been observed in our measurement, which might suggest good battery performance at low temperatures. In addition to the specific materials discussed in this study, we suggest that our method of single-nanorod measurement could be used as a new way to investigate the interaction between electrolyte and electrode materials, revealing relationships that current methods have not ascertained.

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Supporting Information Available: Additional figures depicting experimental results. This material is available free of charge via the Internet at http://pubs.acs.org.

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