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Li$_{2+x}$MnSi$_{1-x}$Al$_x$O$_4$/C nanoparticles for high capacity
lithium-ion battery cathode applications

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Abstract Li$_{2+x}$MnSi$_{1-x}$Al$_x$O$_4$/C (0 < x < 0.09) nanoparticles were synthesized under supercritical fluid conditions at 550 °C for 3 h using water and ethanol as mixed solvent. Compared with Li$_2$MnSiO$_4$, Li$_{2+x}$MnSi$_{1-x}$Al$_x$O$_4$ displays higher theoretical capacity and better electrochemical properties. Li$_{2.09}$MnSi$_{0.91}$Al$_{0.09}$O$_4$/C nanoparticles with an average diameter of ca. 11 nm deliver a high discharge capacity of 308 mAh·g$^{-1}$, which indicates approximate two Li$^+$ ion exchange can be realized during the cycling process.

Introduction

Recently, lithium-ion batteries have attracted great interest due to their widely applications in mobile phones, laptop computers and other portable electronic devices. Especially, the lithium-ion batteries have been also considered to one of promising power sources for electric vehicles and hybrid-electric vehicles.1-3 It is well known that the increased capacity of lithium-ion batteries can improve the performances of the above electronic and electric equipments. However, the low theoretical specific capacity of conventional cathode materials, such as LiCoO$_2$ (274 mAh·g$^{-1}$), LiMn$_2$O$_4$ (148 mAh·g$^{-1}$) and LiFePO$_4$ (170 mAh·g$^{-1}$), limits the increase of battery capacity. Compared with conventional cathode materials, lithium metal silicate (Li$_2$MSiO$_4$, M = Fe, Mn, Co and Ni) have high theoretical specific capacity (> 330 mAh·g$^{-1}$, which is possible while extracting two Li$^+$ ion per formula unit).

Mn$^{3+}$/Mn$^{4+}$ redox couples, due to its lower cell voltage,4 is convenient to study rather than Fe$^{3+}$/Fe$^{4+}$, Co$^{3+}$/Co$^{4+}$ or Ni$^{3+}$/Ni$^{4+}$. Therefore, Li$_2$MnSiO$_4$ can be a more suitable choice as one of high capacity cathodes. Up to now, some methods have been developed to optimize the electrochemical properties of Li$_2$MnSiO$_4$, for example, controlling the particle size and morphology,5-7 doping certain cations8,9 and surface...

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coating with conductive materials\textsuperscript{10-15} and the deintercalation of more than one Li\textsuperscript{+} ion already is achieved in Li\textsubscript{2}MnSiO\textsubscript{4} cathodes, which makes Li\textsubscript{2}MnSiO\textsubscript{4} show high capacity than other cathodes.\textsuperscript{16} Then, can the higher theoretical specific capacity be realized in cathodes?

The existence of aluminum silicate inspires us the doping of Al\textsuperscript{3+} may be a correct choice. When part of [SiO\textsubscript{4}]\textsuperscript{4-} has been replaced into [AlO\textsubscript{4}]\textsuperscript{5-} in Li\textsubscript{2}MnSiO\textsubscript{4}, to compensate for the extra negative charge, corresponding lithium-ion is introduced and forms Li\textsubscript{2+x}MnSi\textsubscript{1-x}Al\textsubscript{x}O\textsubscript{4}, which leads a substantial increase of theoretical specific capacity \((\frac{(2 + x)F}{3.6(161 + 6x)} \text{mAh\cdot g}^{-1}, F \text{ is Faraday constant})\) in cathodes. The substitution process can be described by the following defect equation:

\[
\frac{1}{2} \text{Al}_2\text{O}_3 + \text{Si}^{x\text{Si}} + \frac{1}{2} \text{Li}_2\text{O} \rightarrow \text{Al}^{/\text{Si}} + \text{Li}^{./i} + \text{SiO}_2
\]

Islam group once calculated the trivalent ion substitution on the Si site, and found the introducing additional lithium into Li\textsubscript{2}MnSiO\textsubscript{4} was feasible\textsuperscript{17}. Nevertheless, to our best knowledge, there have been few reports on the synthesis of Li\textsubscript{2+x}MnSi\textsubscript{1-x}Al\textsubscript{x}O\textsubscript{4} cathodes\textsuperscript{8}, probably since the substitution of [AlO\textsubscript{4}]\textsuperscript{5-} tetrahedron instead of [SiO\textsubscript{4}]\textsuperscript{4-} tetrahedron decreases the structure stability of Li\textsubscript{2+x}MnSi\textsubscript{1-x}Al\textsubscript{x}O\textsubscript{4}. Here we develop a supercritical mixed solvothermal approach to fabricate Li\textsubscript{2+x}MnSi\textsubscript{1-x}Al\textsubscript{x}O\textsubscript{4}/C (0 < x <0.09) nanoparticles. In this approach, the carbon coating (arising from the carbonization of ethanol) plays a pivotal role. Li\textsubscript{2+x}MnSi\textsubscript{1-x}Al\textsubscript{x}O\textsubscript{4} are covered by the carbon coating, which not only increases the products’ structure stability, but also improves the products’ electrochemical properties.

**Experimental**

Li\textsubscript{2+x}MnSi\textsubscript{1-x}Al\textsubscript{x}O\textsubscript{4}/C (x = 0, 0.05, 0.07, 0.09) nanoparticles were synthesized by mixed solvothermal process. LiOH·H\textsubscript{2}O, MnCl\textsubscript{2}·4H\textsubscript{2}O, SiO\textsubscript{2} (Cabosil) and AlCl\textsubscript{3}·6H\textsubscript{2}O were dissolved in 10 ml mixed solution of ethanol (5 ml) and water (15 ml) in a molar ratio of (2+x):1:(1-x):x. After being stirred for few minutes, the mixture was transferred and sealed into a 40 ml stainless steel autoclave and kept at 550 °C for 3 h, and then being cooled to room temperature naturally. The product was
centrifuged and washed with distilled water and absolute ethanol several times, and finally dried in vacuum at 60 °C for 5 h.

The phase structure of samples was examined by Powder X-ray diffraction (XRD; Philips X’pert MPD with Cu Kα radiation, λ = 1.5418 Å). The morphology and size of samples were analyzed by high-resolution transmission electron microscopy (HRTEM; JEOL-2011, using an accelerating voltage of 200 kV). The chemical compositions of samples were determined by energy dispersive X-ray spectroscopy (EDX), inductive coupled plasma emission spectroscopy (ICP-ES; Optima 7000DV) and atomic absorption spectroscopy (AAS; AAanlyst 800). The carbon content of samples was evaluated from thermal gravimetric analysis (TGA; DT-50) data.

Electrochemical tests were conducted with coin cells (type 2016). The cathode was fabricated by pasting 80 wt% active material, 10 wt% acetylene black, and 10 wt% polyvinylidene fluoride (PVDF) dissolved in N-methylpyrrolidone (NMP) onto Al foil by doctor blade technique, and then the foil was dried under vacuum at 100 °C for 12 h. Metallic lithium was used as the anode. The electrolyte was 1 M LiPF₆ in a 1:1 mixture of ethylene carbonate (EC)/dimethyl carbonate (DMC); the separator was Celgard 2400. All the cells were assembled in a glove box filled with pure argon. The cells were cycled on an Arbin BT2043 battery test system between 1.5 and 4.5 V versus Li⁺/Li.

Results and discussion

The XRD patterns of the as-prepared samples (Li₂⁺ₓMnSi₁₋ₓAlₓO₄/C) with different ratios of Al doping are shown in Fig. 1, and the calculated lattice parameters and cell volumes from the XRD patterns are listed in table 1. The diffraction peaks of the product obtained without Al doping (x = 0) are well assigned to the orthorhombic phase Li₂MnSiO₄ (space group Pmn2₁, fig. 1a). The calculated lattice parameters and cell volumes are in agreement with the previous reports. Fig. 1b and c show the XRD patterns of Li₂.03MnSi₀.95Al₀.05O₄ and Li₂.07MnSi₀.93Al₀.07O₄, respectively. In the two cases, the additional Li⁺ ions have been introduced into Li₂MnSiO₄ along with a small amount of Al³⁺ ions doping(x ≤ 0.07). The two products both maintain the orthorhombic structures. The increase of cell volumes can be attributed to the Li⁺
interstitial compensation and the larger ionic radius of Al³⁺ (0.5 Å) than Si⁴⁺ (0.41 Å). At x = 0.09, the crystal structure of Li₂₀.₉₉MnSi₀.₉₁Al₀.₀₉O₄ shifts to the monoclinic phase (fig. 1d),¹⁹ which indicates the higher doping concentration be able to change the crystal structure of Li₂₊ₓMnSi₁₋ₓAlₓO₄.

As listed in Table 2, the data derived from AAS, ICP-ES and EDX analysis confirm the chemical compositions of the Li₂₊ₓMnSi₁₋ₓAlₓO₄ samples. The results are close to the expected value.

Fig. 2a and b give the TEM images of Li₂₀.₅₅MnSi₀.₉₄Al₀.₀₅O₄/C and Li₂₀.₉₉MnSi₀.₉₁Al₀.₀₉/C nanoparticles, respectively. The two products both display irregular spherical polyhedron morphologies and relatively narrow size distributions. From Fig. 2c and d, one can see about 80% of the Li₂₀.₅₅MnSi₀.₉₄Al₀.₀₅O₄/C particles are in range of 7–12nm, with an average diameter of ca. 9 nm, and about 80% of the Li₂₀.₉₉MnSi₀.₉₁Al₀.₀₉/C particles are in range of 7–16nm, with an average diameter of ca. 11 nm.

The microstructures of Li₂₀.₅₅MnSi₀.₉₄Al₀.₀₅O₄/C composites were further investigated by HRTEM (Fig. 3a). By analyzing the image contrast, we deduced the dark core part is a Li₂₀.₅₅MnSi₀.₉₄Al₀.₀₅O₄ nanoparticle, and the well-resolved lattice planes illustrate that the nanoparticles are single crystalline and structurally uniform. The measured interplanar spacing is ca. 0.42 nm, which corresponds to the separation of {110} planes of orthorhombic phase Li₂₀.₅₅MnSi₀.₉₄Al₀.₀₅O₄. Outside of the core region, the light part is a layer of carbon coating with 1-2 nm thickness. The amounts of the carbon coating are measured by TGA in air atmosphere. As shown in Fig. 3b, 12.4% weight loss corresponds to the presence of carbon. The first weight loss below 120 °C is probably due to evaporation of absorbed water on the the surface of cathode materials.

In our reaction system to fabricate Li₂₊ₓMnSi₁₋ₓAlₓO₄ cathode materials, the carbon, which roots in the carbonization of ethanol under supercritical conditions: C₂H₅OH → 2C + H₂O + 2H₂,²⁰ plays an important role at least in three: (1) improving the structural stability of Li₂₊ₓMnSi₁₋ₓAlₓO₄: the comparative experiments without ethanol showed that no Li₂₊ₓMnSi₁₋ₓAlₓO₄ were obtained; (2) controlling the size and
morphology of products: the neonatal Li$_{2+x}$MnSi$_{1-x}$Al$_x$O$_4$ seeds were covered by the carbon coating, which blocked the growth of these seeds and generated Li$_{2+x}$MnSi$_{1-x}$Al$_x$O$_4$/C nanoparticles$^{21}$ (3) enhancing the Li-ion diffusion and electronic conductivity.$^{10}$

Supercritical fluid, which exhibits both a density close to that of liquid and high diffusivity and low viscosity similar to that of gas, is helpful for the formation nanoparticles.$^6$ In our sealed system, the supercritical state can be easily realized and provides a constant environment for the formation Li$_{2+x}$MnSi$_{1-x}$Al$_x$O$_4$/C nanoparticles.

The first charge–discharge curves for the samples at 0.05 C between 1.5 and 4.5 V are shown in Fig. 4a-d. In the initial cycle, the Li$_{2+x}$MnSi$_{1-x}$Al$_x$O$_4$/C nanoparticles deliver a discharge capacity of more than 170 mAh·g$^{-1}$, which demonstrates there is beyond one Li$^+$ ion exchange in per formula unit (fig. 4e). Moreover, the three aluminum silicate samples exhibit higher discharge capacity than that of silicate sample, and the increase of Li$^+$ ions exchange exceeds the amount of additional lithium, which reveals the Li$_{2+x}$MnSi$_{1-x}$Al$_x$O$_4$ (x$\neq$0) not only present more lithium but also are in favor of the intercalation and deintercalation of Li$^+$ ions. That may be caused by the enlarged Li$_{2+x}$MnSi$_{1-x}$Al$_x$O$_4$ (x$\neq$0) cell volumes, which expand the transfer channels and contribute to the ion diffusion in the electrode reaction.$^{22}$

Furthermore, all the samples have similar voltage profiles with the two discharge potential plateaus at around 2.8 V and 3.2 V, and two charge potential plateaus at around 3.25 V and 4.2 V, which correspond to the Li$^+$ ions intercalation and deintercalation process, respectively.$^6,11$ The two discharge/charge potential plateaus suggest the existence of two stage Li$^+$ ions intercalation/deintercalation process, which respectively originate from the Mn$^{2+}$/Mn$^{3+}$ and Mn$^{3+}$/Mn$^{4+}$ redox reactions.$^6$

Figure 5 represents the cycle performance of the samples at 0.05 C between 1.5 and 4.5 V. The discharge capacity dramatically decreases at the second cycle. This phenomenon mainly results from the decomposition of the electrolyte and subsequent formation of an organic layer deposited on the surface of particles that occur in the low-potential region for transition metal oxides. The capacity fading rate becomes slow during the subsequent cycles. That can be attributed to the existence of the
carbon coating, which prevents direct contact between the active cathode materials and the electrolyte,\textsuperscript{21} and thus slows down the erosion of Li\textsubscript{2+x}MnSi\textsubscript{1-x}Al\textsubscript{x}O\textsubscript{4} upon cycling. Among these samples, Li\textsubscript{2.05}MnSi\textsubscript{0.95}Al\textsubscript{0.05}O\textsubscript{4}/C appears higher capacity retention than the other samples and 204 mAh·g\textsuperscript{-1} capacity can be retained after 50 cycles (fig. 5b). In our opinion, the introduction of aluminum decreases the crystal field splitting energy and Jahn-Teller effect of [MnO\textsubscript{4}] tetrahedron, which preserves the crystal structure of cathode materials during cycling. However, the substitution of [AlO\textsubscript{4}]\textsuperscript{5-} in place of [SiO\textsubscript{4}]\textsuperscript{4-} decreases the structure stability of Li\textsubscript{2}MnSiO\textsubscript{4}, so the excess aluminum doping is not conducive to the improvement of the cycle performance (fig. 5c and d).

The rate performance of the samples has been shown in Fig. 6. It is obvious that the Li\textsubscript{2.05}MnSi\textsubscript{0.95}Al\textsubscript{0.05}O\textsubscript{4}/C exhibits an excellent discharge capability in any measured rate. We think that the improvement of the rate performance should be put in relation to monodisperse nanoparticles, conductive carbon coating and slight Jahn-Teller effect.

Conclusions

In summary, Li\textsubscript{2+x}MnSi\textsubscript{1-x}Al\textsubscript{x}O\textsubscript{4}/C nanoparticles have been synthesized under supercritical fluid conditions. Benefiting from the enlarged cell volumes and ion diffusion channels, Li\textsubscript{2+x}MnSi\textsubscript{1-x}Al\textsubscript{x}O\textsubscript{4} cathodes have not only the increased theoretical specific capacity but also the improved electrochemical properties. At x=0.09, the obtained Li\textsubscript{2.09}MnSi\textsubscript{0.91}Al\textsubscript{0.09}O\textsubscript{4}/C nanoparticles display higher discharge capacity; at x=0.05, the obtained Li\textsubscript{2.05}MnSi\textsubscript{0.95}Al\textsubscript{0.05}O\textsubscript{4}/C nanoparticles display lower capacity fading and better rate performance. The carbon coating promotes the formation of Li\textsubscript{2+x}MnSi\textsubscript{1-x}Al\textsubscript{x}O\textsubscript{4} nanoparticles and accelerates electron and ion transportation. To develop Li\textsubscript{2+x}MnSi\textsubscript{1-x}Al\textsubscript{x}O\textsubscript{4} into one of the practical cathode materials with high specific capacity, the further studies are required to improve their stability, reversibility and conductivity.

Acknowledgements

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Department of Henan Province (13A150186).

References


3 L. Lu, X. Han, J. Li, J. Hua and M. Ouyang, *J. Power Sources*, 2013, **226**, 272.


Table 1 Calculated lattice parameters ($a$, $b$, $c$ and $\beta$), cell volumes ($V$) and structural units per cell ($Z$) for orthorhombic ($x = 0, 0.05, 0.07$) and monoclinic ($x = 0.09$) Li$_{2+x}$MnSi$_{1-x}$Al$_x$O$_4$

<table>
<thead>
<tr>
<th>Samples</th>
<th>$a$ (Å)</th>
<th>$b$ (Å)</th>
<th>$c$ (Å)</th>
<th>$\beta$ (degree)</th>
<th>$V$ (Å$^3$)</th>
<th>$Z$</th>
</tr>
</thead>
<tbody>
<tr>
<td>Li$_2$MnSiO$_4$</td>
<td>6.311</td>
<td>5.380</td>
<td>4.970</td>
<td>90.00</td>
<td>168.747</td>
<td>2</td>
</tr>
<tr>
<td>Li$<em>{2.05}$MnSi$</em>{0.95}$Al$_{0.05}$O$_4$</td>
<td>6.334</td>
<td>5.358</td>
<td>5.100</td>
<td>90.00</td>
<td>173.082</td>
<td>2</td>
</tr>
<tr>
<td>Li$<em>{2.07}$MnSi$</em>{0.93}$Al$_{0.07}$O$_4$</td>
<td>6.350</td>
<td>5.401</td>
<td>5.102</td>
<td>90.00</td>
<td>174.980</td>
<td>2</td>
</tr>
<tr>
<td>Li$<em>{2.09}$MnSi$</em>{0.91}$Al$_{0.09}$O$_4$</td>
<td>6.382</td>
<td>10.966</td>
<td>5.547</td>
<td>91.06</td>
<td>388.140</td>
<td>4</td>
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</table>

Table 2 Chemical composition of the Li$_{2+x}$MnSi$_{1-x}$Al$_x$O$_4$ samples ($x = 0, 0.05, 0.07, 0.09$)

<table>
<thead>
<tr>
<th>Samples</th>
<th>Li</th>
<th>Mn</th>
<th>Si</th>
<th>Al</th>
</tr>
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<tbody>
<tr>
<td>Li$_2$MnSiO$_4$</td>
<td>1.995</td>
<td>0.999</td>
<td>1.000</td>
<td>0.000</td>
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<td>Li$<em>{2.05}$MnSi$</em>{0.95}$Al$_{0.05}$O$_4$</td>
<td>2.046</td>
<td>0.994</td>
<td>0.955</td>
<td>0.047</td>
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<tr>
<td>Li$<em>{2.07}$MnSi$</em>{0.93}$Al$_{0.07}$O$_4$</td>
<td>2.066</td>
<td>0.992</td>
<td>0.941</td>
<td>0.068</td>
</tr>
<tr>
<td>Li$<em>{2.09}$MnSi$</em>{0.91}$Al$_{0.09}$O$_4$</td>
<td>2.085</td>
<td>0.987</td>
<td>0.922</td>
<td>0.089</td>
</tr>
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</table>

Fig. 1 XRD patterns of Li$_{2+x}$MnSi$_{1-x}$Al$_x$O$_4$/C samples with (a) $x = 0$, (b) $x = 0.05$, (c) $x = 0.07$ and (d) $x = 0.09$
Fig. 2 TEM images of (a) Li\textsubscript{2.05}MnSi\textsubscript{0.95}Al\textsubscript{0.05}O\textsubscript{4}/C and (b) Li\textsubscript{2.09}MnSi\textsubscript{0.91}Al\textsubscript{0.09}/C; particle size distributions of (c) Li\textsubscript{2.05}MnSi\textsubscript{0.95}Al\textsubscript{0.05}O\textsubscript{4}/C and (d) Li\textsubscript{2.09}MnSi\textsubscript{0.91}Al\textsubscript{0.09}/C.

Fig. 3 (a) HRTEM image of Li\textsubscript{2.05}MnSi\textsubscript{0.95}Al\textsubscript{0.05}O\textsubscript{4}/C; (b) TGA curve of Li\textsubscript{2.05}MnSi\textsubscript{0.95}Al\textsubscript{0.05}O\textsubscript{4}/C.
Fig. 4 The first charge–discharge curves of Li_{2-x}MnSi_{1-x}Al_xO_4/C samples with (a) x = 0, (b) x = 0.05, (c) x = 0.07 and (d) x = 0.09 at 0.05 C; (e) graphic illustration of the number of Li^+ ion exchange in per Li_{2-x}MnSi_{1-x}Al_xO_4 unit and corresponding discharge capacity while the first discharging (green column is the experimental value, and red column is the theoretical value.).
Fig. 5 Cyclic performance of Li$_{2+x}$MnSi$_{1-x}$Al$_x$O$_4$/C samples with (a) $x = 0$, (b) $x = 0.05$, (c) $x = 0.07$ and (d) $x = 0.09$ at 0.05 C.

Fig. 6 Rate performance of Li$_{2+x}$MnSi$_{1-x}$Al$_x$O$_4$/C samples with (a) $x = 0$, (b) $x = 0.05$, (c) $x = 0.07$ and (d) $x = 0.09$. 