Title
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Improving the Performance of Lithium Manganese Phosphate

Through Divalent Cation Substitution

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Abstract

Highly crystalline samples of LiMnPO₄ and its analogs with partial substitution of Mn by divalent Mg, Cu, Zn, and Ni were prepared by hydrothermal synthesis and characterized by x-ray diffraction and infrared spectroscopy. Chemical oxidation produced two-phase mixtures of the initial phases LiMnₙ₋₁ₓMₓPO₄ and the delithiated forms, LiₓMnₙ₋₁ₓMₓPO₄, all with the olivine structure. The extent of oxidation depended upon the quantity of oxidizing agent used and on the identity of the substituent ions. Mg, Ni and Cu were found to increase the level of delithiation relative to that in pure LiMnPO₄. Mg was also shown to reduce the tendency of the oxidized phase to absorb water.

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Introduction

Olivine-type LiMPO₄ (M = Fe, Mn, Co, Ni) compounds have an orthorhombic comprised of close-packed phosphate anions with M²⁺ ions in corner-sharing MO₆ sites and Li ions in edge-sharing LiO₆ sites.¹ The strong P-O covalent bond gives these materials good stability and makes them attractive as positive electrodes for lithium batteries in vehicle applications. LiMnPO₄ (lithiophilite) cycles at a nearly constant potential of 4.1 V vs. Li/Li⁺, which is well-suited to use in standard battery configurations. In contrast with LiFePO₄, however, the manganese compound suffers from poor Li extraction and insertion kinetics. Many factors have been considered to contribute to the slow phase transition, including the electronic and ionic conductivities, the Jahn-Teller effect in Mn³⁺, interface strain due to the large volume change between LiMnPO₄ and MnPO₄, and the metastable nature of the delithiated phase.²⁴ Although the mechanism of the two-phase reaction has not been identified, it is likely that it is similar to that of the LiFePO₄/FePO₄ system.⁵ Previous efforts⁶⁻⁹ to improve the performance of stoichiometric LiMnPO₄ have mostly been limited to particle size minimization, which increases the rate and utilization, but inevitably decreases the volumetric energy density of the electrode.

Studies have shown that M-site doping with divalent cations such as Mg²⁺, Zn²⁺, Cu²⁺, Ni²⁺ and Co²⁺ can improve the performance of LiFePO₄.¹⁰⁻¹⁴ Yamada has intensively investigated the Li(Mn,Fe)PO₄ solid solution system¹¹⁻¹⁴, and found that although Fe-substituted LiMnPO₄ has kinetics and utilization properties superior to those of pure LiMnPO₄, for Mn content above 80 %, Fe does not provide sufficient stabilization, and some decomposition takes place on extraction of lithium. Solid-state
synthesis of LiMnPO$_4$ and LiMn$_{(1-y)}$M$_y$PO$_4$ (M = Mg, Ca, Zn, Ni, Co, Cu, Al, B, Cr and Nb) samples at high temperatures has been reported in patents by Valence Technology Inc.,$^{19-20}$ Phase-pure, well-formed LiMnPO$_4$ crystals have recently been synthesized by low-temperature hydrothermal methods.$^{21-23}$ Chen et al.$^{22-23}$ prepared LiMn$_{(1-y)}$Mg$_y$PO$_4$ crystals with a wide range of Mg substitution. Here we report hydrothermal syntheses of M$^{2+}$-substituted LiMnPO$_4$ crystals with M = Ni, Cu and Zn. The reactivities of these samples and one Mg-substituted sample were examined through chemical oxidation, and a mechanism is proposed to describe the effects of substitution. Electrochemical properties of the synthesized phosphates will be reported in a future publication.

**Experimental**

LiMnPO$_4$ crystals were synthesized using the hydrothermal method described previously for LiFePO$_4$.$^5$ Equimolar amounts of MnSO$_4$.H$_2$O (Mallinckrodt, Inc) and H$_3$PO$_4$ (85%, J. T. Baker) were mixed in 30 ml deionized and deoxygenated water to give an Mn concentration of 1.0 M. A 1.5 M LiOH (Spectrum) solution was added slowly with stirring to give Mn:P:Li equal to 1:1:3 and a final pH of 10. Substantial precipitation occurred during this step. After stirring under helium for another 5 min, the reaction mixture was transferred to a 125 ml Teflon-lined reactor, which was tightly sealed after purging with helium, then held at 220 °C for 5 h. On cooling to room temperature, the off-white precipitate was filtered, thoroughly washed with deionized water, and dried in a vacuum oven at 60 °C for 24 h. The substituted LiMnPO$_4$ samples were prepared using the same procedure, except that a 9:1 molar mixture of MnSO$_4$.H$_2$O with Mg(NO$_3$)$_2$.6H$_2$O (>99%, EM Science), Ni SO$_4$.6H$_2$O (Johnson Matthey Electronics),
CuSO$_4$.5H$_2$O (99%, J. T. Baker) or ZnSO$_4$.7H$_2$O (99%, Aldrich), was dissolved before adding the H$_3$PO$_4$ and LiOH solutions. Delithiation was achieved by stirring the samples in aliquots of a 0.1 M solution of nitronium tetrafluoroborate (NO$_2$BF$_4$, 95+, Aldrich) in acetonitrile for 24 h at room temperature in an argon-filled glovebox with O$_2$ < 1 ppm and H$_2$O < 2 ppm.

X-ray diffraction (XRD) patterns were acquired using a Panalytical Xpert Pro diffractometer equipped with monochromatized Cu K$_\alpha$ radiation. The scan rate was 0.0025°/s from 10° to 70° 2θ in 0.01° steps. Lattice parameters and phase ratios in the oxidized samples were determined by pattern refinement using Riqas software (Materials Data, Inc.). Scanning electron microscopy (SEM) images were obtained using a Hitachi S-4300 SE/N microscope at 20 and 10 kV accelerating voltages, with energy dispersive X-ray analysis (EDX) and elemental mapping using the Noran System SIX. Fourier transform infrared spectroscopy (FTIR) measurements were performed on KBr pellets using a Nicolet 6700 spectrometer in transmission mode with a spectral resolution of 4 cm$^{-1}$.

Results and Discussion

Synthesis and Characterization

XRD patterns of substituted and unsubstituted LiMnPO$_4$ samples are shown in Fig. 1. The unsubstituted and Mg-substituted samples were phase pure, while small amounts of unidentified impurities were detected in Ni-, Cu- and Zn-substituted samples. This is probably due to lower solubility of the Ni, Cu or Zn precursor in the reaction mixture, resulting in formation of the substituted olivine before complete uptake was
possible, rather than a true limit of solubility in the parent compound. Lattice parameters obtained from Rietveld refinement of the whole patterns are listed in Table 1. The unsubstituted LiMnPO$_4$ had $a = 10.4474$ Å, $b = 6.1016$ Å, $c = 4.7506$ Å and $V = 302.83$ Å$^3$, in good agreement with the reported values.$^{24}$ M$^{2+}$ substitution decreased all three lattice parameters. The cell volume decreased 0.8% for Mg$^{2+}$, 0.6% for Ni$^{2+}$, 0.3% for Cu$^{2+}$ and 0.4% for Zn$^{2+}$ substitution, consistent with their smaller ionic radii. The lattice parameters of our LiMg$_{0.1}$Mn$_{0.9}$PO$_4$ sample were slightly smaller than those reported by Chen et al., whose crystals were much larger and needle-like.$^{23}$ Site occupancy refinement showed no excess electron density on the lithium site in each case. The Scherrer crystallite sizes were about 30 nm for all samples except the Mg-substituted one, which was 48 nm.

SEM images of the hydrothermal synthesized samples are shown in Fig. 2. The unsubstituted LiMnPO$_4$, Ni-, Cu- and Zn-substituted samples were elongated diamond-shaped aggregates composed of small crystals about 200 nm in size. The morphology of the Mg-substituted sample was quite different. It consisted of smooth, hexagonal-shaped crystal plates measuring about 0.8 μm x 0.4 μm x 0.1 μm. EDX and elemental mapping (Fig. 3) showed a uniform distribution of Mn, Mg, P and O. The atomic ratio of Mn to Mg was 8.8:1, consistent with the intended ratio of 9:1, and was constant among crystals and at different locations on individual crystals.

**Chemical delithiation**

To investigate the effects of substitution on reactivity, the extent of lithium extraction by the NO$_2$BF$_4$ solution was determined by XRD. In each case, the oxidized
samples contained of a second olivine-type phase along with unreacted starting material. The potential of the NO$_2^+/\text{NO}_2$ couple is ca. 5.1 V vs. Li/Li$^+$, more than sufficient to remove Li from LiMnPO$_4$ according to:

\[
\text{LiMnPO}_4 + x\text{NO}_2\text{BF}_4 \rightarrow (1-x)\text{LiMnPO}_4 + x\text{MnPO}_4 + x\text{LiBF}_4 + x\text{NO}_2 \quad [1]
\]

In the case of a 10 % substituted olivine in which the substituent, M, is not oxidized, we may expect the reaction to proceed as follows:

\[
\text{LiM}_{0.1}\text{Mn}_{0.9}\text{PO}_4 + 0.9x\text{NO}_2\text{BF}_4 \rightarrow (1-x)\text{LiM}_{0.1}\text{Mn}_{0.9}\text{PO}_4 + x\text{Li}_{0.1}\text{M}_{0.1}\text{Mn}_{0.9}\text{PO}_4 + 0.9x\text{LiBF}_4 + 0.9x\text{NO}_2 \quad [2]
\]

XRD patterns of samples treated with an amount of NO$_2$BF$_4$ sufficient to extract 50 % of the active lithium are shown in Fig. 4. The lattice parameters of the delithiated phases are given in Table 1. Ten percent of the unsubstituted LiMnPO$_4$ had been converted to the oxidized phase after 24 h. Under the same conditions, the Zn$^{2+}$-substituted, Cu$^{2+}$-substituted and Ni$^{2+}$-substituted samples were 2 %, 14% and 18% oxidized, respectively. The Mg$^{2+}$-substituted sample was the most reactive, with 23% of the delithiated phase formed. Thus, Mg$^{2+}$, Ni$^{2+}$ and Cu$^{2+}$ substitution were beneficial, while Zn$^{2+}$ had a negative effect.

The Mg$^{2+}$-substituted crystals were several times larger than those of the other phosphates, which were aggregated crystals about 200 nm size. The observed improvement is clearly not due to a size effect, but rather to the presence of Mg$^{2+}$ in the
structure. At this point it is unclear why the Zn$^{2+}$-substituted sample was so difficult to delithiate. One possibility is the presence of an insoluble impurity on the surface that hinders the reaction. Synthesis experiments carried out using different metal precursors, solution concentrations and/or pH may provide the answer.

FTIR absorption spectra of the unsubstituted and substituted samples are compared in Fig. 5a. The bands at frequencies $> 900$ cm$^{-1}$ are attributed to the symmetric and antisymmetric stretching vibration of PO$_4^{3-}$ anion, while those at lower frequencies include bending vibrations of the anion and lattice modes. The low frequency bands were largely unaffected by substitution. The higher frequency bands, however, were shifted toward higher energy, suggesting an increase in P-O bond strength in the substituted samples. The FTIR spectra of the partially oxidized samples are shown in Fig. 5b. In each case, a new band appeared at about 1075 cm$^{-1}$, increasing in intensity with the amount of the oxidized phase present.

When unsubstituted samples were treated with increasing quantities of the NO$_2$BF$_4$ solution, the extent of delithiation increased, reaching 30 % and 70 % for NO$_2$BF$_4$:LiMnPO$_4$ ratios of 1:1 and 2:1 respectively after 24 h reaction (Fig. 6a). The result is consistent with the report by Yamada et al$^{16}$ where 70% of delithiation was achieved after reaction for 48 hr at a ratio of 2:1. In contrast, oxidation of Mg-substituted samples reached 60 % and 100 % under the same conditions (Fig. 6b). The domain sizes of the delithiated phases increased monotonically with the extent of delithiation in each case, to a maximum of 10 nm for unsubstituted LiMnPO$_4$, and to 18 nm for the fully oxidized Mg-substituted sample.
It has been reported\textsuperscript{1-2} that partially and fully delithiated LiMnPO\textsubscript{4} samples absorb water in air. We indentified the hydration products by XRD as MnPO\textsubscript{4}·1.5H\textsubscript{2}O and Mn\textsubscript{0.5}Mg\textsubscript{0.1}PO\textsubscript{4}·1.5H\textsubscript{2}O. This instability, attributed to the elastic energy accumulated inside the Jahn-Teller deformed lattice, complicates the isolation and characterization of the pure MnPO\textsubscript{4}.\textsuperscript{3} Li\textsubscript{0.1}Mg\textsubscript{0.1}Mn\textsubscript{0.9}PO\textsubscript{4} was found to be stable in air when coexisting with LiMg\textsubscript{0.1}Mn\textsubscript{0.9}PO\textsubscript{4}, slowly converting to Li\textsubscript{0.1}Mg\textsubscript{0.1}Mn\textsubscript{0.9}PO\textsubscript{4}·1.5H\textsubscript{2}O only in fully oxidized samples. Our crystal samples of both Li\textsubscript{0.1}Mg\textsubscript{0.1}Mn\textsubscript{0.9}PO\textsubscript{4} and MnPO\textsubscript{4} were stable when kept in an inert atmosphere.

\textit{Possible mechanism of the substituent effect}

The ionic radius of Mn\textsuperscript{2+} in octahedral coordination is 0.98 Å,\textsuperscript{26} which decreases to 0.785 Å upon oxidation to Mn\textsuperscript{3+}. The substituent ion radii are 0.83 Å for Ni\textsuperscript{2+}, 0.86 Å for Mg\textsuperscript{2+}, 0.87 Å for Cu\textsuperscript{2+} and 0.88 Å for Zn\textsuperscript{2+}, and they remain the same in the lithiated and delithiated phases. Assuming that the phase transformation progresses through the “reactive transition zone” mechanism as described in LiFePO\textsubscript{4},\textsuperscript{5} Mn-site substitution may influence LiMnPO\textsubscript{4} oxidation in several ways. The presence of the larger ions, which are more compatible with the olivine structure, may stabilize the lattice against the strain created by the small Jahn-Teller Mn\textsuperscript{3+} ions. The decreased reactivity of Li\textsubscript{0.1}Mg\textsubscript{0.1}Mn\textsubscript{0.9}PO\textsubscript{4} in air, as well as the presence of larger crystalline domains in the Mg-substituted material point toward such stabilization. This may, in turn, create a more favorable boundary between the two phases and facilitate the conversion of one phase to the other without loss of coherence.
Substitution also decreases the volume change between the two end members (Fig. 7), in part because $M^{2+}$ is smaller than $Mn^{2+}$ but larger than $Mn^{3+}$, and also because the Li associated with the electrochemically inactive $M^{2+}$ remains in the substituted delithiated phase. For the unsubstituted sample, the volume change between LiMnPO$_4$ and MnPO$_4$ is 9.5%, while the changes are 8.2% for Zn substitution, 8.0% for Cu, 7.9% for Ni and 7.8% for Mg. The mismatch in the $ac$ plane, in which the phase boundary is probably located, is also reduced, with the largest decrease observed for Mg.

Finally, if the products of decomposition of the delithiated phases contains only $Mn^{3+}$, a substantial de-mixing of ions is required, presenting a kinetic barrier to rearrangement.

**Conclusions**

Divalent cation-substituted LiMnPO$_4$ crystals were synthesized by a hydrothermal method and characterized by XRD and FTIR. The substituents resided only on Mn sites. Chemical delithiation produced two-phase mixtures of lithiated and delithiated olivine-type phases. The presence of Mg$^{2+}$, Ni$^{2+}$ and Cu$^{2+}$ resulted in increased conversion to the delithiated phases. Further, Mg-substitution improved the stability of the delithiated phase with respect to hydration in air. These effects are attributed to stabilization of the oxidized phases due to the ion sizes and the incomplete removal of lithium from the lattice.
Acknowledgements

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Table 1. Cell parameters of substituted and unsubstituted LiMnPO$_4$

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<th>$c$ (Å)</th>
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References


Figure captions

1. X-ray diffraction patterns of hydrothermal synthesized samples: a) LiMnPO₄, b) Mg-substituted LiMnPO₄, c) Ni-substituted LiMnPO₄ d) Cu-substituted LiMnPO₄, and e) Zn-substituted LiMnPO₄. * indicates unidentified impurity peaks • indicates Al peaks from sample holder.

2. SEM images of hydrothermal synthesized samples: a) LiMnPO₄, b) Mg-substituted LiMnPO₄, c) Ni-substituted LiMnPO₄, d) Cu-substituted LiMnPO₄, and e) Zn-substituted LiMnPO₄.

3. SEM image and the corresponding element maps of Mg-substituted LiMnPO₄.

4. XRD patterns of the oxidized samples prepared using a 1:2 ratio of NO₂BF₄ to the indicated phosphates: a) LiMnPO₄, b) Mg-substituted LiMnPO₄, c) Ni-substituted LiMnPO₄, d) Cu-substituted LiMnPO₄, and e) Zn-substituted LiMnPO₄.

5. FTIR absorption spectra of a) hydrothermally synthesized samples and b) samples oxidized using a 1:2 ratio of NO₂BF₄ to the phosphates.

6. XRD patterns of samples oxidized using indicated NO₂BF₄ to phosphate ratios: a) LiMnPO₄ and b) Mg-substituted LiMnPO₄; c) oxidation percentage comparison.
Chen and Richardson, Figure 1
Chen and Richardson, Figure 2
Chen and Richardson, Figure 3
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