Stabilizing the Structure of LiCoPO4 Nanocrystals via Addition of Fe3+: Formation of Fe3+ Surface Layer, Creation of Diffusion-Enhancing Vacancies, and Enabling High-Voltage Battery Operation

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Stabilizing the Structure of LiCoPO_4 Nanocrystals via Addition of Fe^{3+}: Formation of Fe^{3+} Surface Layer, Creation of Diffusion-Enhancing Vacancies, and Enabling High-Voltage Battery Operation

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ABSTRACT: Factors affecting the cyclability of the Fe-substituted LiCoPO_4 (LiCo_{0.8}Fe_{0.2}PO_4, LCFP) material were elucidated, including both the structural and electrode/electrolyte stability. Electrochemical characterization of the synthesized LCFP nanoparticles lends clear evidence for improved electrochemical stability of LCP, as well as enhanced rate capability, with Fe^{3+} substitution. Surface analysis using X-ray photoelectron spectroscopy (XPS) and electron energy loss spectroscopy (EELS) suggest that Fe enrichment on the surface of LCFP occurs through the oxidation of Fe^{2+} into Fe^{3+} in the synthesis process. The Fe^{3+}-rich phase on the LCP surface enhances the stability of the delithiated phase, preventing oxidative reactions with electrolytes during high-voltage operation. This surface protection persists as long as the electrochemical reduction of Fe^{3+} is avoided by ensuring that the full range of operating voltages lie above the Fe^{3+}/Fe^{2+} redox potential. Our findings may offer new approaches to stabilize the structure of LCP and other high-voltage positive electrodes for use in 5 V-class Li-ion batteries.

INTRODUCTION

Growing global demands for electric and hybrid electric vehicles necessitate the development of 5 V-class lithium ion batteries (LIBs) in view of their potential for higher specific energy and voltage compared with the 4 V-class LIBs. From this perspective, LiCoPO_4 (LCP) is an interesting material that offers advantages complementary to those of conventional cathode materials such as LiCoO_2 and LiFePO_4 (LFP): high reaction potential of 4.8 V vs. Li/Li^+, theoretical capacity of 167 mAh g^{-1}, and thermal stability thanks to the P=O covalent bonding.1-3 Although the low electron conductivity (<10^{-9} S cm^{-1}) and thermal stability thanks to the P=O covalent bonding1-3,8 and the primary focus of this study—is its poor cyclability: LIBs with LCP cathodes exhibit a dramatic decrease in capacity within just a few 10s of cycles. Mechanisms proposed to explain the plummeting capacity include instability of the ethylene carbonate (EC)-based electrolyte,11 cleavage of P=O bonds by nucleophilic attack of F^{-} in LiPF_6-containing electrolytes,12 and instability of the delithiated phase (Li_xCoPO_4, x << 1).13,14 Formation of a protective surface film on LCP using alternatives to EC-based solvents, such as fluorinated ethylene carbonate15 and other electrolyte additives,16 has succeeded in prolonging useful life to 100 cycles. Allen et al. recently showed that Fe substitution into LCP in combination with a phosphite electrolyte additive...
enhances electrochemical performance, extending cyclability to 500 cycles with 80% retention of initial capacity. 16, Fe substitution—specifically, aliovalent Fe3+ substitution in Li+ (M1) and transition metal M2+ (M2) sites obtained by such simple air-annealing process—has been well-known for other phosphates such as LFP and LiMn1−xFexPO4 (LMFP) to cause changes in the structural characteristics (defective) and in the electrochemical signature (slope).17–19 In case of LCP, the Fe3+ substitution yields structural stabilization at high potential such as 4.5 V vs. Li/Li+ and creation of vacancies in the Li+ (M1) sites.16,20 It is reported that these vacancies contribute to increased Li mobility, thus enhancing rate capability.16,21 Hanafusa et al. demonstrated that Fe3+-substituted LCP can charge and discharge with negligible capacity loss (<5%) over 150 cycles even in conventional EC-based electrolyte.20 However, achieving such an excellent cyclability in Fe3+-substituted LCP requires restricting the operating voltage range between 3.5 and 5.0 V vs. Li, thus sacrificing capacity contributed by the Fe3+/Fe2+ redox below 3.5 V vs. Li. Moreover, the detailed mechanism responsible for improved cyclability, and the role played by the presence of Fe3+ within the LCP crystal structure, remains unclear.

This study aims at elucidating the influence of Fe3+ substitution on LCP cyclability by focusing on electrode/electrolyte interface. Using LCP nanoparticles (100 nm in average diameter) highly dispersed within multiwalled carbon nanotube (MWCNT) matrix prepared via ultracentrifugation,25 we have synthesized different LCP-based materials with different Fe3+ amounts. Electrochemical characterizations of the synthesized materials give clear evidence of improved electrochemical stability of LCP as well as enhanced rate capability, with Fe3+ substitution. Surface analysis using X-ray photoelectron spectroscopy (XPS) and electron energy loss spectroscopy (EELS) suggest that Fe enrichment of the surface of LCP nanoparticles occurs through the oxidation of Fe2+ into Fe3+ along with the creation of vacancies at Co2+ (M2) sites during the synthesis process. The Fe3+-rich phase on the LCP surface enhances the stability of the delithiated phase, preventing oxidative reactions with electrolytes during high-voltage operation. This surface protection persists over 1000 cycles at 0.2 C and 5000 cycles at 1.0 C as long as the electrochemical reduction of Fe3+ is avoided by ensuring that the full range of operating voltages lies above the Fe3+/Fe2+ redox potential. Additionally, we first demonstrated for LCP or LCfP materials that the vacancies in Co/Fe M2 sites are responsible for the Li+ diffusivity enhancement within LCPF crystals and thus for the improvements of C-rate capability. Our findings may offer new approaches for stabilizing the structure of LCP and other high-voltage positive electrodes for use in 5 V-class Li-ion batteries.

### EXPERIMENTAL SECTION

**Materials.** Co(CH3COO)2·H2O (>99%, Wako Pure Chemicals), Fe(CH3COO)2 (>99%, Sigma-Aldrich Corp.), CH3COOLi (>98.0%, Wako Pure Chemicals), and H3PO4 (>85.0%, Wako Pure Chemicals) were used to prepare LiCo0.8Fe0.2PO4 (LCPF). Citric acid (C6H8O7·H2O) (>99.5%, Sigma-Aldrich Corp.) was used as a chelating agent. Multiwalled carbon nanotubes (MWCNTs) with a specific surface area of 240 m2 g−113 were used for the preparation of carbon composite. Distilled water (17 MΩ) was used as a medium for the entire preparation scheme.

**Preparation of LiCo0.8Fe0.2PO4/MWCNT Composite under UC Treatment.** First, two kinds of solution (solution A and B) were prepared. Solution A was composed of 0.4000 g of MWCNT and 0.6714 g of H3PO4 (1.0 equiv) mixed in 50 mL of ultrapure water under ultrasonication. Solution B was composed of 1.1725 g of Co(CH3COO)2·4H2O (0.8 equiv), 0.2027 g of Fe(CH3COO)2 (0.2 equiv), 0.3922 g of CH3COOLi (1.0 equiv), and 1.1247 g of citric acid (1.0 equiv) dissolved in 20 mL of ultrapure water. Solution A was subjected to the ultracentrifugation (uc) treatment17 at 80°C for 5 min before and after the addition of solution B. The uc-treated sol was further dried at 130°C under vacuum for 12 h. The obtained powder made of LCPF precursor and MWCNT was preannealed at 350°C in air for 1 h to remove absorbed water and citric-acid-derived amorphous carbon, then left to cool to room temperature (RT). The preannealed powder was then fast annealed at 650°C under N2 flow (RT to 650°C in 3 min, dwell for 5 min, 20 min to cooling to RT) to form the LCPF/MWCNT composite (pristine LCPF/MWCNT). The pristine LCPF/MWCNT composite was further annealed at 350°C for 3 h under air in order to oxidize Fe2+ into Fe3+ without incinerating MWCNTs (air-annealed LCPF/MWCNT).

**Physicochemical Characterization of LiCo0.8Fe0.2PO4/MWCNT Composite.** Structure analysis on pristine and air-annealed LCPF samples was performed by X-ray diffraction (XRD, Smart-lab (Rigaku), Cu Kα radiation (λ = 1.54 Å), operating at 45 kV, 200 mA). XRD patterns were recorded in the 10°−90° 2θ range at a scan speed of 0.0033° s−1. Time-of-flight neutron powder diffraction (NPD) measurements were performed at the iMATERIA (BL20) facility of the Japan Photon Accelerator Research Complex (J-PARC, Ibaraki, Japan) to obtain NPD patterns corresponding to 0.5−4.8 Å in d-spacing.26 Fe Mössbauer spectra were collected at room temperature in transmission geometry on a constant acceleration spectrometer using a 57Co source in a Rh matrix. Isomer shift (IS) calibrations were performed using Fe as a standard at room temperature.

**Electrochemical Characterization of LiCo0.8Fe0.2PO4/MWCNT Composite.** Half-cells were assembled using a negative Li metal electrode and a positive LiCo0.8Fe0.2PO4/MWCNT electrode. Distilled water (17 MΩ) was used as an electrolyte for the composite. Distilled water (17 MΩ) was used as an electrolyte for the composite. Half-cells were assembled using a negative Li metal electrode and a positive LiCo0.8Fe0.2PO4/MWCNT electrode in 2032 coin-type cells. The electrolyte was a mixture of ethylene carbonate, propyl carbonate, and dimethyl carbonate (EC: PC: DMC = 1:1:3) containing 1.0 M of lithium hexafluorophosphate (LiPF6). LiCo0.8Fe0.2PO4/MWCNT electrode was prepared by mixing 90 wt % of the composite and 10 wt % of polyvinylidene difluoride (PVdF) in N-methyl pyrrolidone (NMP). The mixture was coated using an NFR-2000 (Nikon Corp.) to evaluate the ratio of Co and Fe species using references selected to exhibit different oxidation states, such as Co3+ (LiCoPO4 bulk sample), Co2+ (Cobalt acetylacetonate), Fe3+ (LiFePO4 bulk sample), and Fe2+ (FePO4 bulk sample oxidized by the chemical method using NO3−, Fe3+ (LiFePO4 bulk sample), Co2+ (Cobalt acetylacetonate), Fe3+ (LiFePO4 bulk sample), and Fe2+ (FePO4 bulk sample oxidized by the chemical method using NO3−, Fe2+ (FePO4 bulk sample oxidized by the chemical method using NO3−, FePO4 bulk sample oxidized by the chemical method using NO3−, FePO4 bulk sample oxidized by the chemical method using NO3−, FePO4 bulk sample oxidized by the chemical method using NO3−).
Table 1. Rietveld Refinement Results of Neutron Powder Diffraction Patterns for Air-Annealed LCFP/MWCNT Composite

<table>
<thead>
<tr>
<th>space group</th>
<th>atom</th>
<th>Wyckoff</th>
<th>x/a</th>
<th>y/b</th>
<th>z/c</th>
<th>Biso</th>
<th>occupancy</th>
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</thead>
<tbody>
<tr>
<td>Pnma</td>
<td>Li(^+)</td>
<td>4a</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>2.6(2)</td>
<td>0.984(7)</td>
</tr>
<tr>
<td></td>
<td>Fe(^{3+})</td>
<td>4a</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>2.6(2)</td>
<td>0.024(1)</td>
</tr>
<tr>
<td></td>
<td>Co(^{2+})</td>
<td>4c</td>
<td>0.2763(2)</td>
<td>0.25</td>
<td>0.9892(4)</td>
<td>1.19(5)</td>
<td>0.762(2)</td>
</tr>
<tr>
<td></td>
<td>Fe(^{3+})</td>
<td>4c</td>
<td>0.2763(2)</td>
<td>0.25</td>
<td>0.9892(4)</td>
<td>1.19(5)</td>
<td>0.129(2)</td>
</tr>
<tr>
<td></td>
<td>P</td>
<td>4c</td>
<td>0.0925(2)</td>
<td>0.25</td>
<td>0.4200(2)</td>
<td>0.89(2)</td>
<td>1</td>
</tr>
<tr>
<td></td>
<td>O(^-)</td>
<td>4c</td>
<td>0.10209(9)</td>
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<td>0.7443(2)</td>
<td>0.84(2)</td>
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<tr>
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<td>0.1982(2)</td>
<td>1.13(2)</td>
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<td>O(^-)</td>
<td>8d</td>
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<td>0.0466(2)</td>
<td>0.2817(2)</td>
<td>1.09(2)</td>
<td>1</td>
</tr>
</tbody>
</table>

\(R_{wp} = 5.76, R_p = 4.49, R_e = 2.34, \chi^2 = 6.07\)

Figure 1. Structural characterization of air-annealed LiCo\(_{0.8}\)Fe\(_{0.2}\)PO\(_4\) (LCFP)/MWCNT composite. (a) XRD patterns and (b) neutron powder diffraction patterns for air-annealed LCFP/MWCNT composite. (c) Mössbauer spectrum fitted with two Fe environments (Fe\(^{2+}\) and Fe\(^{3+}\)). (d) Lower-magnification (20 000x) higher-resolution TEM image for highly dispersed LCFP nanocrystalline within the MWCNT matrix. (e) Magnified (400 000x) HRTEM image of the LCFP nanocrystalline. Clear lattice fringes of \(d = 3.86, 5.10, \) and 5.92 Å appear, corresponding to the (210), (200), and (010) planes of LCFP, respectively.
Results and Discussion

Figure 1a shows the X-ray diffraction pattern of the synthesized air-annealed LiCo0.8Fe0.2PO4 (LCFP)/multiwalled carbon nanotube (MWCNT) composite. Main peaks of the air-annealed LCFP composite can be indexed with an olivine structure (S.G. Pnma LiCoPO4: JCPDS card no. 85-0002). Apart from peaks attributed to MWCNT and Co3O4 with very small intensity, no other peaks due to possible impurities such as Fe2O3, LiFePO4 and FePO4 were detected, indicating successful substitution of Fe into the LiCoPO4 (LCP) crystalline structure. Neutron powder diffraction (NPD) analysis of the air-annealed LCFP (Figure 1b) confirmed the existence of vacancies in the synthesized crystal structure. Lattice parameters, atomic positions, and occupancies, including Li and expected vacancy sites, for air-annealed LCFP were determined through Rietveld refinements on the NPD data. The observed NPD peaks can be indexed by an olivine structure with lattice constants of \(a = 10.1592(7) \text{ Å}, b = 5.9114(4) \text{ Å}, \) and \(c = 4.7091(3) \text{ Å} \) and a unit-cell volume of \(V = 282.80(3) \text{ Å}^3\), as shown in Table 1. These values of the \(a\) and \(b\) lattice parameters and the cell volume are smaller than those of LiCoPO4 (LCP, see Table S1), in good agreement with previous reports.\(^{10}\)

The obtained stoichiometry of the air-annealed LCFP crystal structure is \((\text{Li}_{0.98}\text{Fe}_{0.02})\text{M}1(\text{Co}_{0.76}\text{Fe}_{0.13}\text{O}_{0.085})\text{M}2\text{PO}_4\), where \(\text{M}1\), \(\text{M}2\), and \(\square\) respectively, represent Li, Co, and vacancy sites. Fe\(^{3+}\) is present primarily at transition metal sites (M2), while the elemental mapping of the pristine LiCo0.5Fe0.5PO4 shows uniform distribution of Fe, Co, O, and P within the particles. While an average Co/Fe ratio of 83/17 is calculated from NDPI refinement, the XPS experiments show that the distribution of Fe and Co varies from the surface to the bulk of the particles. When a higher Co/Fe ratio of 83/17 is calculated from NDP refinement, the XPS experiments show that the surface (no Ar ion etching) ratio is equal to 75.4/42.6 and progressively increases when moving toward the center of the particle (increasing Ar ion etching duration). In contrast, the Co/Fe ratio of pristine LCFP/MWCNT composite remains fixed at approximately 80/20 independent of location between surface and center of the particle (independent of Ar ion etching duration) (Figure S2). This suggests that the air-annealing process induces, beside the oxidation of the Fe and creation of Co vacancies to balance the charge, the migration of Fe\(^{3+}\) ions toward the surface of the particles in the amorphous domain. Such phenomenon is confirmed by the analysis of air-annealed samples of a composite with higher Fe concentration, namely, LiCo0.8Fe0.2PO4/MWCNT. Elemental mapping (Figure S4a) and line analysis (Figure S4c) of this sample confirms the existence of a Fe-rich surface layer on the nanoparticle surface, while the elemental mapping of the pristine LiCo0.8Fe0.2PO4 shows uniform distribution of Fe, Co, O, and P within the particle (Figure S4b). This inhomogeneous distribution of Fe allows, in addition, interpreting the apparently overestimated ratio of Co and Fe calculated from Co 2p\(^{3/2}\) and Fe 2p\(^{3/2}\) peaks.

Figure 2. Consecutive XPS measurements with data acquisition at 5 s intervals during 0−30 s Ar ion etching, yielding a depth profile of air-annealed LCFP/MWCNT composite: (a) Co 2p\(^{3/2}\) peak and Co satellite peak, (b) Fe 2p\(^{3/2}\) peak and Co LMM Auger electron peak, (c) atomic ratio of Co and Fe calculated from Co 2p\(^{3/2}\) and Fe 2p\(^{3/2}\) peaks.
The refinement of NDP take to account only the vacancy-rich part of the particle thus explaining the apparent over-estimation. The electrochemical behavior of LCFP/MWCNT composites was investigated in half-cells versus Li anode at a rate of...
0.2C (1C = 167 mA g\(^{-1}\)) (Figure 3). During the charge/discharge processes, a voltage shoulder around 3.5 V and two plateaus were observed at 4.76 and 4.83 V (Figure 3a). The shoulder corresponds to the redox reaction of Fe\(^{2+}/Fe^{3+}\) in LiCo\(_{0.8}\)Fe\(_{0.2}\)PO\(_4\). The two plateaus at high potentials correspond to a two-step redox reaction of Co\(^{2+}/Co^{3+}\) in LiCo\(_{0.8}\)Fe\(_{0.2}\)PO\(_4\).
Li$_{0.6}$Co$_{0.8}$Fe$_{0.2}$PO$_4$ which well agrees with previous reports.\textsuperscript{20,24,25} The air-annealed LCFP/MWCNT composites showed a high initial discharge capacity of 130 mAh g$^{-1}$ (Figure 3a) and increased stability of the cycle performance with 86% capacity retention after 50 cycles compared to LCP/MWCNT and pristine LCFP/MWCNT composites (Figure 3d). During cycling between 2.5 and 5.0 V, dQ/dV plots show an increase near the Fe$^{2+}$/Fe$^{3+}$ redox peak at 3.5 V and a decrease near the broad redox peak at 3.4 V (Figure 3c). Such change in redox peaks suggests that Fe$^{3+}$ in the air-annealed LCFP is gradually reduced into Fe$^{2+}$ at low potentials and that the energy level of resultant Fe$^{3+}$ produced by reoxidation of Fe$^{2+}$ is different from that for Fe$^{3+}$ originally existed in the air-annealed LCFP. To avoid this phenomenon, we perform charge and discharge tests of the composites with operating voltages confined to the range 4.3–5.0 V. This results in excellent cycle performance, with 99% capacity retention at 0.2C after 1000 cycles, and 85% at 1C after 5000 cycles (Figure 3e) largely exceeding the best reported one by Hanafusa et al.\textsuperscript{27}

Additionally, air-annealed LCFP/MWCNT composite enabled discharge at a 100C rate with capacity of 45 mAh g$^{-1}$, larger than observed for LCP/MWCNT and pristine LCFP/MWCNT composites (Figure 3f). Comparison of Li$^+$ diffusion coefficients ($D_{li^+}$) between the pristine (Fe$^{2+}$) and air-annealed LCFP (Fe$^{3+}$) composites shown in Figure S5 supports such improvement in C-rate capability. The obtained $D_{li^+}$ for the air-annealed LCFP is higher in both charge and discharge compared to the pristine. Such results indicate that the creation of vacancies in M2 sites resulting from the oxidation/migration of Fe ions enhances the Li$^+$ diffusivity within LCFP crystal, unlike previous reports which suggest the enhancement of Li$^+$ mobility due to the vacancies in Li (M1) sites.\textsuperscript{16,21} EIS analysis using symmetric cells\textsuperscript{26} (see Figure S6) shows that the pristine LCFP shows higher impedance behavior than air-annealed LCFP (with Fe$^{3+}$-rich surface). Both the charge transfer and mass transport processes were improved for Fe$^{3+}$-rich LCFP: the relevant absolute impedance decreases at low voltage is also responsible for the cation mixing.

The evolution of Co and Fe valence state along cycling has been determined by X-ray absorption measurements. The X-ray absorption near edge structure (XANES) spectra of Co and Fe K-edge for air-annealed LCFP/MWCNT composite before cycling and at cut off voltages of the two investigated voltage ranges (2.5–5.0 V and 4.3–5.0 V) are shown in Figure S8, while calculated average oxidation states for Co and Fe are reported in Table S3.

In the air-annealed LCFP/MWCNT composite material before cycling, valence states of Co and Fe are +2.1 and +2.9, respectively, in good agreement with previous Mössbauer spectroscopy results (Figure 1c). During cycling in the (2.5–5.0 V) voltage range, the Fe valence state changes between +2.3 and +3.0 at lower and upper cutoff voltages, respectively, and these values remain stable along cycles. The Co valence state at low cut off voltage is equal to +2.1 and remain stable along cycles while at 5.0 V at the end of the first charge it is equal to +2.8 and progressively decreases along cycles to reach +2.5 at the 30th charged state. During cycling in the (4.3–5.0 V) voltage range, Fe valence state, as expected, increases during the first charge from +2.9 in the sample before cycling up to +3. The valence state of +3 is maintained then whatever the voltage along cycles. For the Co, the valence state is +2.1 and +2.8 at 4.3 and 5.0 V, respectively. These values are stable along cycles. This stable cycle performance confirms the specific role of the Fe$^{3+}$ which together with vacancies at M2 sites of LCFP effectively prevents irreversible changes in crystal structure. This stabilizing effect added to the previously detailed prevention of decomposition of electrolyte during charge and discharge processes, result in stable cycle performance with improved rate performance for Fe-substituted samples.

Based on the results of XRD, neutron diffraction analysis, Mössbauer, XANES, and XPS studies, we propose the schematic picture illustrated by Figure 5 for the structural changes occurring during cycling. Upon cycling, LCP as a reference material suffers both structural degradation (Li–Co cation mixing) and SEI accumulation on the particle surface, while only the latter can be restrained by a high cutoff voltage operation within the range of 4.3 and 5.0 V (Figure 5a). Fe$^{3+}$ was introduced as a partial substitute for Co in our material. It

**Figure 3a** shows the pristine LCFP shows higher impedance behavior than air-annealed LCFP (with Fe$^{3+}$-rich surface). Both the charge transfer and mass transport processes were improved for Fe$^{3+}$-rich LCFP: the relevant absolute impedance becomes roughly 1/10 for the charge transfer process and 1/5 for the diffusion process when moving from pristine LCFP to air-annealed LCFP.
was observed that, upon air annealing, some of the introduced Fe$^{3+}$ cations migrated to the nanoparticle surface, producing vacancies at M2 sites. However, the vacancies disappeared during cycling between 2.5 and 5.0 V, due to the reduction of Fe$^{3+}$ into Fe$^{2+}$ at 3.5 V, resulting in a destabilization of the charged Co$_3$Fe$_5$PO$_4$ structure. In contrast, the Fe$^{3+}$-rich phase on the surface stabilizes the unstable vacancies at M2 sites. However, the vacancies disappeared due to the increased repulsion between Fe$^{3+}$ and Co$^{2+}$. The Fe$^{3+}$-rich phase formed on the LCP surface plays a critical role in yielding stable charge–discharge performance. This structure results in stable cycle performance.

**CONCLUSIONS**

In conclusion, a detailed understanding of the mechanism of LCP structure stabilization by Fe$^{3+}$ substitution suggests that the Fe$^{3+}$-rich phase formed on the LCP surface plays a significant role in yielding stable charge–discharge performance. The oxidation of Fe$^{2+}$ into Fe$^{3+}$ during air annealing of the pristine LCPF results in diffusion of Fe ions to the surface due to the increased repulsion between Fe$^{3+}$ and Co$^{2+}$. The Fe$^{3+}$-rich phase on the surface stabilizes the unstable delithiated phase of (Li$_{0.02}$Fe$_{0.02}$)Co$_{0.76}$Fe$_{0.11}$Li$_{0.11}$PO$_4$ (x ≪ 1), preventing both the crystal structure degradation and continuous SEI formation on the LCPF nanoparticle surface. The phase stabilization yields excellent cyclability, with 85% capacity retention over 5000 cycles at 1C and 96% capacity retention over 1000 cycles even at a slow C-rate of 0.2C. The importance of Fe$^{3+}$ retention in the crystal structure, which requires to prevent Fe$^{3+}$ reduction to Fe$^{2+}$ around 3.5 V vs. Li/Li$^+$, was demonstrated by a combination of XRD and XAFS analysis on samples with and without operation-voltage extending below 3.5 V vs. Li. Meanwhile, differently from previous reports, it was found that the vacancies on Co/Fe M2 sites improve the overall electrochemical performances such as the capacity and C-rate capability, thanks to the enhancement of Li$^+$ diffusion within LCPF crystals. By demonstrating the existence of the Fe-rich phase on LCPF nanoparticles and its critical impact on electrochemical performance, this study may open the way to the design of stable 5 V cathode materials via fine-tuning of surface phenomena.

**ASSOCIATED CONTENT**

1. Supporting Information

The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/acs.chemmater.8b01965.

Additional experimental data: Neutron powder diffraction, XRD, Mössbauer spectroscopy, XPS, XAFS and elemental mapping line analysis for air-annealed LCP/MWCNT, LCPF/MWCNT and pristine LCPF/MWCNT composites (PDF)

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**Notes**

The authors declare no competing financial interest.

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Supporting Information

Stabilizing the structure of LiCoPO₄ nanocrystals via addition of Fe³⁺: Formation of Fe³⁺ surface layer, creation of diffusion-enhancing vacancies, and enabling high-voltage battery operation

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Neutron powder diffraction analysis on the air-annealed LiCoPO₄ (LCP) / Multiwalled carbon nanotube (MWCNT) composite.

To support the structure analysis on the air-annealed LiCo₀.₈Fe₀.₂PO₄ (LCFP), neutron powder diffraction (NPD) of the air-annealed LiCoPO₄ (LCP) was recorded. The synthesis procedure of the air-annealed LCP was as same as described in the main text without an use of Fe species (Fe(CH₃COO)₂). Time-of-flight neutron powder diffraction (NPD) measurements were performed at the iMATERIA (BL20) facility of the Japan Photon Accelerator Research Complex (J-PARC, Ibaraki, Japan) to obtain NPD patterns corresponding to 0.5-4.8 Å in d-spacing. Lattice parameters and atomic positions for the air-annealed LCP were determined through Rietveld refinements on the NPD data. The observed NPD peaks shown in Figure S1 can be indexed by an olivine structure with lattice constants of a = 10.2014(1) Å, b = 5.9237(7) Å, and c = 4.7007(5) Å and a unit-cell volume of V =
284.06(5) Å³, as shown in Table S1. These values of the \( a \) and \( b \) lattice parameters and the cell volume are larger than those of LCFP (see Table 1 in the main text). The atomic-position analysis on LCP can be well fitted without vacancies in the synthesized crystal structure unlike the air-annealed LCFP.

![Figure S1 Neutron powder diffraction patterns for the air-annealed LCP / MWCNT composite.](image)

**Table S1  Rietveld refinement results of neutron powder diffraction patterns from Figure S1 for the air-annealed LCP / MWCNT composite.**

<table>
<thead>
<tr>
<th>Space group</th>
<th>( a / \text{Å} )</th>
<th>( b / \text{Å} )</th>
<th>( c / \text{Å} )</th>
<th>( V / \text{Å}^3 )</th>
</tr>
</thead>
<tbody>
<tr>
<td>( Pnma )</td>
<td>10.2014(2)</td>
<td>5.9237(7)</td>
<td>4.7007(5)</td>
<td>284.06(5)</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Atom</th>
<th>Wyckoff</th>
<th>( x/a )</th>
<th>( y/b )</th>
<th>( z/c )</th>
<th>( B_{iso} )</th>
<th>Occupancy</th>
</tr>
</thead>
<tbody>
<tr>
<td>M1 site (Li site)</td>
<td>Li⁺</td>
<td>4a</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>1.01(2)</td>
</tr>
<tr>
<td>M2 site (Co site)</td>
<td>Co²⁺</td>
<td>4c</td>
<td>0.2775(3)</td>
<td>0.25</td>
<td>0.9781(7)</td>
<td>0.50(7)</td>
</tr>
<tr>
<td>P</td>
<td>4c</td>
<td>0.0943(2)</td>
<td>0.25</td>
<td>0.4190(3)</td>
<td>0.50(3)</td>
<td>1</td>
</tr>
<tr>
<td>O</td>
<td>4c</td>
<td>0.0972(2)</td>
<td>0.25</td>
<td>0.7420(3)</td>
<td>0.50(3)</td>
<td>1</td>
</tr>
<tr>
<td>O</td>
<td>4c</td>
<td>0.4543(2)</td>
<td>0.25</td>
<td>0.2037(3)</td>
<td>0.50(3)</td>
<td>1</td>
</tr>
<tr>
<td>O</td>
<td>8d</td>
<td>0.1663(1)</td>
<td>0.0430(1)</td>
<td>0.2817(2)</td>
<td>0.50(2)</td>
<td>1</td>
</tr>
</tbody>
</table>

\( R_w = 5.13 \), \( R_p = 3.71 \), \( R_e = 1.52 \), \( \chi^2 = 7.14 \)

X-ray diffraction and room temperature Mössbauer spectroscopy on the LCFP/MWCNT with and without the air-annealing process.

Mössbauer spectroscopy was conducted on the pristine LCFP/MWCNT, in order to verify the idea that the air-annealing of the pristine LCFP/MWCNT is the responsible process to create Fe³⁺ in the structure. To begin with, X-ray diffraction (XRD) pattern of the pristine LCFP crystal was recorded. Structure analysis on pristine and air-annealed LCFP samples was performed by X-ray diffraction [XRD, Smart-lab (Rigaku), Cu Kα radiation (\( \lambda = 1.54 \) Å), operating at 45 kV, 200 mA]. XRD patterns were recorded in the 10°-90° 2θ range at a scan speed of 0.0033° s⁻¹. The obtained XRD pattern of the pristine LCFP was shown along with that of the air-annealed LCFP in Figure S2(a). As same as for the air-annealing LCFP, all peaks of both the pristine LCFP composite correspond to a single phase of an olivine structure with \( Pnma \) space group (LiCoPO₄ : JCPDS card no. 85-0002). Excluding peaks
attributed to MWCNT, no other peaks due to possible impurities such as Co$_3$O$_4$, Fe$_2$O$_3$, LiFePO$_4$ and FePO$_4$ were detected, indicating successful substitution of Fe into the LiCoPO$_4$ (LCP) crystalline structure. Lattice parameters for the pristine LCFP were determined through Rietveld refinements on the XRD data, as shown in Table S2. Mössbauer spectroscopy for the pristine LCFP is shown in Figure S2(b) and compared with that for the air-annealing LCFP. $^{57}$Fe Mössbauer spectra were collected in transmission geometry on a constant acceleration spectrometer using a $^{57}$Co source in a Rh matrix operated at room temperature. Isomer shift (IS) calibrations were performed using a foil α-Fe as a standard at room temperature. The obtained spectrum for the pristine LCFP was best fitted by two Lorentzian doublets: one with isomer shift (IS) = 0.51 mm s$^{-1}$ and quadrupole splitting (QS) = 0.73 mm s$^{-1}$, and the other with IS = 1.2 mm s$^{-1}$ and QS = 3.0 mm s$^{-1}$. The former, accounting for 12% of the full spectra, corresponds to the Fe$^{3+}$ present on M1 and disordered M2 sites $^{1,2}$, and the latter, accounting for 88% of the spectra, corresponds to Fe$^{2+}$ on M2 sites. The drastic increase in Fe$^{3+}$ ratio from 12% to 88% indicates that the air-annealing is the main responsible process for the Fe$^{3+}$ creation in the LCFP crystal structure.

Figure S2  Structural characterization of pristine and the air-annealed LCFP / MWCNT composites. (a) XRD patterns for LCFP /MWCNT composites. (b) Mössbauer spectrum fitted with two Fe environments (Fe$^{2+}$ and Fe$^{3+}$).

Table S2  Rietveld refinement results of X-ray powder diffraction patterns from Figure S2(a) for pristine and the air-annealed LCFP / MWCNT composite.

<table>
<thead>
<tr>
<th></th>
<th>Space group</th>
<th>$a$ / Å</th>
<th>$b$ / Å</th>
<th>$c$ / Å</th>
<th>$V$ / Å$^3$</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pristine LCFP</td>
<td>Pnma</td>
<td>10.2237(8)</td>
<td>5.9376(5)</td>
<td>4.6985(4)</td>
<td>285.22(4)</td>
</tr>
<tr>
<td>Air annealed LCFP</td>
<td>Pnma</td>
<td>10.1589(7)</td>
<td>5.9105(4)</td>
<td>4.7090(3)</td>
<td>282.75(3)</td>
</tr>
</tbody>
</table>
X-ray Photoelectron Spectra of Co 2P_{3/2} and Fe 2P_{3/2} on the pristine LCFP/MWCNT composite.

The Co/Fe ratio of the pristine LCFP / MWCNT composite was evaluated from the fitting of X-ray photoelectron spectra. X-ray photoelectron spectroscopy (XPS JEOL Ltd. JPS-9200) was carried out with a pass energy of 40 eV for high-resolution scans using a monochromated Mg X-ray source. Prior to the XPS characterization, the sample electrodes were thoroughly washed by dimethyl carbonate (DMC) and dried overnight. The Co/Fe ratio was determined quantitatively by computing the areas under the Co 2p_{3/2} and Fe 2p_{3/2} peaks (Figure S2). The Co/Fe ratio at the surface of the pristine composite (0 sec) was 80.7/19.3, which well agrees with the dosage of the synthesis process (Co = 0.8 eq and Fe = 0.2 eq, see the paragraph of “Experimental” in the main text). Contrary to the obtained result of the air-annealed LCFP shown in Figure 2, the Fe/Co ratio remains almost fixed, independent of Ar ion etching duration over 30 sec.

![Figure S3](image)

**Figure S3** Consecutive XPS measurements with data acquisition at 5 sec intervals during 0–30 sec Ar ion etching, yielding a depth profile of pristine LCFP / MWCNT composite: (a) Co 2p_{3/2} peak and Co satellite peak, (b) Fe 2p_{3/2} peak and Co LMM Auger electron peak, (c) Atomic ratio of Co and Fe calculated from Co 2p_{3/2} and Fe 2p_{3/2} peaks.

**Electron energy-loss spectroscopy measurements on the pristine and air-annealed LCFP crystals (LiCo_{0.5}Fe_{0.5}PO_{4} / MWCNT)**

To support the hypothesis of an Fe-rich phase on the surface of air-annealed LCFP crystals based on the analysis of XPS spectra (Figure 2 in the main text), we conducted the electron energy-loss spectroscopy on the sample. Here, to compensate for the low ratio of Fe/Co in the composite and the corresponding low intensity of Fe in synthesized LCFP/MWCNT structures, we prepared the pristine and air-annealed samples of a composite with higher Fe concentration, namely LiCo_{0.5}Fe_{0.5}PO_{4} / MWCNT. As shown in Figure S4(a), elemental mapping of the air-annealed LiCo_{0.5}Fe_{0.5}PO_{4} /
MWCNT composite shows an Fe-rich surface layer on the nanoparticle surface which is supported by its line analysis [Figure S4(c)], where as that of the pristine in Figure S4(b) shows uniform distribution of Fe, Co, O, and P within the particle.

(a) Air annealed LiCo$_{0.5}$Fe$_{0.5}$PO$_4$ / MWCNT

(b) Pristine LiCo$_{0.5}$Fe$_{0.5}$PO$_4$ / MWCNT
Figure S4 Elemental mapping line analysis of (a) air annealed LiCo$_{0.5}$Fe$_{0.5}$PO$_4$/MWCNT and (b) pristine LiCo$_{0.5}$Fe$_{0.5}$PO$_4$/MWCNT composites, (c) line analysis of the air annealed LiCo$_{0.5}$Fe$_{0.5}$PO$_4$/MWCNT composite [shown in white frame in Figure S4(a)] obtained from electron energy loss spectroscopy.

Enhancement of lithium ion diffusion coefficients ($D_{Li^+}$): galvanostatic intermittent titration technique (GITT) on the pristine and air-annealed LCFP/MWCNT composites.

In order to elucidate the factor for the C-rate capability enhancement as shown in Figure 3(f) of the manuscript, GITT measurements were conducted and calculated $D_{Li^+}$ was compared between the pristine (Fe$^{2+}$) and air-annealed LCFP (Fe$^{3+}$ and resulted vacancies in Co/Fe M2 sites). As shown in Figure S5(a) and S5(b), $D_{Li^+}$ for the air-annealed LCFP (Fe$^{3+}$ giving vacancy in Co site) is higher compared to the pristine (Fe$^{2+}$) especially in the high potential region (high SOC and low SOD), which
is consistent with the results of C-rate capability tests. These results indicate that the existence of vacancies in M2 site enhances the kinetics of LCFP reaction, and the stable SEI mainly contributes to the cycling stability.

Figure S5. Plots of Li diffusion coefficients vs. state of charge/discharge calculated from GITT curves during (a) charge and (b) discharge for the pristine and air-annealed LCFP. GITT was conducted with 0.1C (1C = 167 mAh g\(^{-1}\) per active material) for 20 min. and rest time for 1h in the potential range of between 4.3-5.0 V vs. Li/Li\(^+\). Before GITT measurements, a pre-cycling was conducted at 0.1C in the same potential range as GITT.

Charge-transfer resistance and lithium ion diffusivity: electrochemical impedance spectroscopy (EIS) on the pristine and air-annealed LCFP/MWCNT composites using symmetric cells.

As shown from GITT measurement (Fig. S5), the apparent \(D_{Li^+}\) was enhanced for the air-annealed LCFP (with Fe\(^{3+}\) rich surface) compared with pristine LCFP. However, GITT analysis generally represents overall diffusion kinetics of active materials, thus, it is hard to distinguish between effects of charge transfer and diffusional properties. We made additional EIS experiments on (a) pristine LCFP and (b) Fe\(^{3+}\)-rich-surface LCFP at two different SOCs, namely 0 and 70\% (middle range of second plateau as shown in Figure 2b). As shown in the Figure S6, the pristine LCFP shows higher impedance behavior than the air-annealed LCFP (with Fe\(^{3+}\) rich surface). Both the charge transfer and mass transport processes have been improved for Fe\(^{3+}\)-rich LCFP: the relevant absolute impedance becomes roughly 1/10 for the charge transfer process and 1/5 for the diffusion process when moving from pristine LCFP to air-annealed LCFP.
Figure S6 Impedance spectroscopy on (a) pristine LCFP and (b) air-annealed LCFP with Fe$^{3+}$-rich-surface at two different SOCs, namely, 0 and 70% (middle range of second plateau). The measurement was made on symmetric cells comprising of both LCFP or LCFP (Fe$^{3+}$-rich) electrodes. Perturbation amplitude was ±10 mV (peak to peak). Frequency ranges was 10 mHz to 65 kHz.

Stability of LCP and LCFP crystal structure at different voltage range operation; 2.5-5.0 V and 4.3-5.0 V (Unit cell volume change).

In order to compare the cycling stability of crystal structure depending on the operation voltage ranges (2.5-5.0 V or 4.3-5.0 V), unit cell volumes for the air-annealed LCFP/MWCNT composites after 1, 10, and 30 cycles were plotted along with those for the air-annealed LCP/MWCNT. Unit cell volumes for the air-annealed LCFP operated within the high voltage range (4.3-5.0 V) remain constant, compared to those for other three samples which show clear increase because of Li-Co mixing during cycling. Such results strongly support the hypothesis that Fe$^{3+}$ and vacancies at M2 sites serve to stabilize the LCFP crystal structure. The air-annealed LCFP operated within 2.5-5.0 V, where its Fe$^{3+}$ undergoes Fe$^{3+}$/Fe$^{2+}$ redox [see Figure 3 (c) in the main text], cannot resist its initial volume increase. Meanwhile, unit cell volume for the air-annealed LCP continuously increase regardless of its operation voltage range.
Figure S7 Unit cell volumes of 4 different samples after charge and discharge up to 30 cycles: the air-annealed LCP / MWCNT and air-annealed LCFP / MWCNT composites at 2 different operation voltage ranges (2.5-5.0 V and 4.3-5.0 V). Each unit cell volume was calculated from the cell parameters obtained from corresponding XRD-pattern analysis.

XAFS measurements on air-annealed LCFP/MWCNT composites

To confirm the effect of charge and discharge tests between 4.3 and 5.0 V on air-annealed LCFP/MWCNT composites, redox states of Co and Fe were investigated by X-ray absorption measurements during cycling. The X-ray absorption near edge structure (XANES) spectra of Co and Fe K-edge for air-annealed LCFP / MWCNT composite are shown in Figure S8. The valence state of Co and Fe in the composite is determined by deconvolving the spectra using the contributions of Co$^{2+}$, Co$^{3+}$, Fe$^{2+}$ and Fe$^{3+}$. At 3.45 V, initial valence states of Co and Fe were +2.1 and +2.9, respectively, in good agreement with previous Mössbauer spectroscopy results [Figure 1(c) in the main text]. Valence numbers of Co and Fe during the 1st, 10th and 30th cycles were obtained from the XANES spectra shown in Figure S7 and Table S3. The Fe valence state over charge and discharge cycles between 2.5 and 5.0 V varied from +2.3 to +3.0, accompanied by a corresponding decrease in change of Co valence state. During cycling between 4.3 and 5.0 V, the valence number of Co exhibited minimal variation, ranging from +2.1 to +2.8 over all cycles. This stable cycle performance is made possible by the constancy of the valence state of Fe$^{3+}$ after the 1st, 10th and 30th cycles between 4.3 and 5.0 V.
Figure S8  XAFS measurements on a pellet of (a)-(b) air-annealed LCFP / MWCNT composite and (c)-(f) half-cell consisting of Li/1M LiPF₆ EC:PC:DMC(vol. 1:1:3)/(air annealed LCFP / MWCNT composite). The XANES spectra at the Co K-edge of the composite are shown in (a), (c) and (e). The XANES spectra at the Fe K-edge of the composite are shown in (b), (d) and (f). (c) and (d) are conducted in potential range between 2.5 and 5.0 V. (e) and (f) are conducted in potential range between 4.3 and 5.0 V.

Table S3  Valence states of Co and Fe of air annealed LCFP / MWCNT composite obtained from XANES spectra shown in Figure S6

<table>
<thead>
<tr>
<th>Cycle No.</th>
<th>Co valence state</th>
<th>Fe valence state</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Discharge</td>
<td>Charge</td>
</tr>
<tr>
<td>Before charge/discharge</td>
<td>+2.10</td>
<td>+2.91</td>
</tr>
<tr>
<td>2.5-5.0 V</td>
<td>1&lt;sup&gt;st&lt;/sup&gt;</td>
<td>+2.12</td>
</tr>
<tr>
<td>10&lt;sup&gt;th&lt;/sup&gt;</td>
<td>+2.11</td>
<td>+2.78</td>
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<tr>
<td>30&lt;sup&gt;th&lt;/sup&gt;</td>
<td>+2.08</td>
<td>+2.55</td>
</tr>
<tr>
<td>4.3-5.0 V</td>
<td>1&lt;sup&gt;st&lt;/sup&gt;</td>
<td>+2.11</td>
</tr>
<tr>
<td>10&lt;sup&gt;th&lt;/sup&gt;</td>
<td>+2.11</td>
<td>+2.82</td>
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<td>30&lt;sup&gt;th&lt;/sup&gt;</td>
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