

A New Low Cost Synthesis Method for LiFePO₄

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Lithium iron phosphate, LiFePO₄ has gained wide acceptance as the cathode material of choice for safe, large format Li-ion battery applications [1]. Since LiFePO₄ possesses a low intrinsic conductivity (around 10⁻⁹ S/cm) carbon-coating methods or suitable preparative approaches must be employed to produce a composite product incorporating a conductive component [2].

In this study we describe a new, solid state synthesis method for the preparation of high purity and highly conductive LiFePO₄. Cost analysis indicates that this preparative approach will be significantly less expensive than any currently employed solid-state manufacturing method. The proprietary approach does not require the inclusion of carbon to enhance the electronic conductivity [2], but does allow the use of low cost Fe³⁺ precursors such as Fe₂O₃ or FePO₄. Advantageously, the preparative approach may be carried out at relatively low synthesis temperatures (typically 500-600°C).

Figure 1 depicts the refined X-ray powder diffraction data for a representative sample of the LiFePO₄ prepared using the new preparative method. The Rietveld refinement of the powder X-ray data was carried out using the GSAS (EXP-GUI) software package [3]. The calculated unit cell parameters are $a = 10.3250(4) \text{ \AA}$, $b = 6.0055(2) \text{ \AA}$, $c = 4.6915(2) \text{ \AA}$, cell volume = 290.90 (2) \AA^3 , in close agreement with literature sources [4].

Figure 2 shows the first cycle constant current cycling data (C/10) for a LiFePO₄ sample prepared by the new low-cost method. These low rate data reveal a reversible specific capacity of around 160 mAh/g, a figure which compares favorably with the theoretical performance (170 mAh/g) and also with state-of-the-art literature data [4].

In an extension of this preparative method a similar approach has been adopted to synthesize lithium iron manganese phosphate, LiFe_{1-x}Mn_xPO₄, as well as several other polyanion active materials.

References:

- [1]. A. K. Padhi, N.S. Nanjundaswamy, C. Masquelier and J.B. Goodenough, *J. Electrochem. Soc.* **144**, 2581 (1997).
- [2]. J. Barker, M.Y. Saidi and J.L. Swoyer, *Electrochem. Solid-State Lett.*, **6**, A53, 2003.
- [3]. A.C. Larsen and R.B. von Dreele, General Structure Analysis System (GSAS), Los Alamos Laboratory Report, LAUR 2000.
- [4]. A. S. Andersson and J. O. Thomas, *J. Power Sources*, **97-98**, 498, (2001).

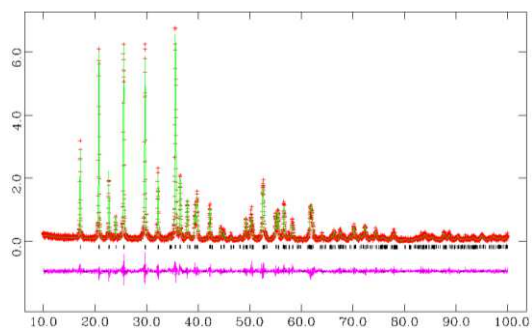


Figure 1: X-ray Diffraction. Rietveld refinement analysis (GSAS EXP-GUI) of a LiFePO₄ sample prepared by the new low-cost synthesis method.

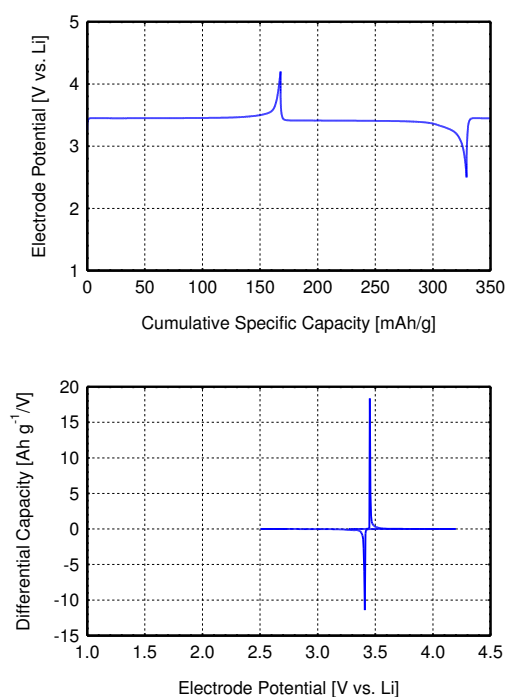


Figure 2: (Upper) First Cycle Electrode Potential versus Specific Capacity and (Lower) Differential Capacity Profile for a Li//LiFePO₄ cell cycled between 2.50 – 4.20 V. The LiFePO₄ was prepared by the new low-cost synthesis method.