RE-SYNTHESIS OF LiCoO₂ EXTRACTED FROM DISCARDED BATTERIES WITH LOW AND HIGH STATE OF HEALTH

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1. Introduction

To avoid ambient contamination and for sustainability reasons, discarded lithium-ion batteries (LIBs) should be recycled. Although several recycling processes are already known, no consider the effect of the battery state of health (SOH) on them. $LiCoO_2$ compounds extracted from cathodes of discarded LIBs with low (L) and high (H) SOH were re-synthesized and its structural and electrochemical properties are discussed.

2. Experimental

The batteries' SOH were measured from charge-discharge cycles performed in the 4.2-3.6 V. Crystalline phases and the lattice parameters of the LiCoO₂ were identified from Rietveld x-ray refinements. The as-extracted LiCoO₂ compounds were thermally decomposed at 700 °C in O₂ atmosphere, whose products were submitted to a solid-state reaction with Li₂CoO₃ at 750 °C in O₂ atmosphere. Galvanostatic charge-discharge cycles in a re-synthesized LiCoO₂/Li cell furnished its charge capacity and voltage profile.

3. Results and Discussions

The compounds extracted from the L and H batteries were identified respectively as $Li_{0.73}CoO_2$ and $Li_{0.96}CoO_2$, that after thermal decomposition resulted in the Li_1CoO_2 and Co_3O_4 compounds, as reaction products. Co_3O_4 concentrations equal to 33.5% and 11.8% in wt were measured for L and H decomposed cathodes, respectively [1]. The solid-state re-synthesis reaction transformed Co_3O_4 into the stoichiometric Li_1CoO_2 compound.

The LiCoO₂ lattice parameters as a function of processing temperatures are shown in Figure 1. The *c* parameter, higher for Li_{0.73}CoO₂ than for Li_{0.96}CoO₂ as-extracted compounds, and the inverse behavior for the *a* parameter is an effect of electrostatic repulsion between the O-Co-O layers, upon Li removal [2]. A Li₁CoO₂ single phase with *c*=14.0496(5) Å and *a*=*b*=2.81412(5) Å was identified after the LiCoO₂ re-synthesized. The c/a = 4.99 and the x-ray peak intensity ratio I(003)/I(104) > 2.6 indicates a well ordered Li₁CoO₂ layered structure, with few or none cationic exchange.

The voltage profiles as a function of the specific discharge capacity is shown in Figure 2a shows for the re-synthesized Li_1CoO_2 electrodes. Higher specific charge capacities were measured for the electrode re-synthesized from the L cathode, Figure 2b. Specific charge capacities of 130.0 and 125.0 mAh g⁻¹ were measured in the twentieth cycle for the re-synthesized L and H cathodes, respectively. We argue that the small size of the re-synthesized particles from the L cathode (as observed from SEM images) can explain the best performance of the corresponding Li₁CoO₂/Li cell, due the higher specific surface area of this electrode.

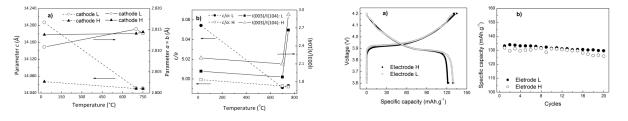
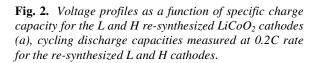


Fig. 1. Lattice parameters for the as-extracted (25 °C), thermal decomposed (700 °C) and re-synthesized (750 °C) LiCoO₂ compound from the L and H cathodes (a), the corresponding c/a and x-ray intensity $I_{(003)}/I_{(104)}$ ratios (b).



4. References

[1]- R. Floriano, A. O. dos Santos, A. Urbano, L. P. Cardoso, J. Scarminio, IJRRAS, 17, 158-166, (2013).
[2]- S. Laubach at al., Phys. Chem. Chem. Phys., 11, 3278–3289, (2009).

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