

Microstructure and Electrochemical Properties of $\text{LiNi}_{1/3}\text{Co}_{1/3}\text{Mn}_{1/3-x}\text{Al}_x\text{O}_2$ Cathode Material for Lithium Ion Batteries

Wei-Bo Hua, Zhuo Zheng, Xiao-Dong Guo, Ben-He Zhong

Abstract—The layered structure $\text{LiNi}_{1/3}\text{Co}_{1/3}\text{Mn}_{1/3-x}\text{Al}_x\text{O}_2$ ($x = 0 \sim 0.04$) series cathode materials were synthesized by a carbonate co-precipitation method, followed by a high temperature calcination process. The influence of Al substitution on the microstructure and electrochemical performances of the prepared materials was investigated by X-Ray diffraction (XRD), scanning electron microscopy (SEM), and galvanostatic charge/discharge test. The results show that the $\text{LiNi}_{1/3}\text{Co}_{1/3}\text{Mn}_{1/3-x}\text{Al}_x\text{O}_2$ has a well-ordered hexagonal $\alpha\text{-NaFeO}_2$ structure. Although the discharge capacity of Al-doped samples decreases as x increases, $\text{LiNi}_{1/3}\text{Co}_{1/3}\text{Mn}_{1/3-0.02}\text{Al}_{0.02}\text{O}_2$ exhibits superior capacity retention at high voltage (4.6 V). Therefore, $\text{LiNi}_{1/3}\text{Co}_{1/3}\text{Mn}_{1/3-0.02}\text{Al}_{0.02}\text{O}_2$ is a promising material for “green” vehicles.

Keywords—Lithium ion battery, carbonate co-precipitation, microstructure, electrochemical properties.

I. INTRODUCTION

LITHIUM-ION batteries (LIBs) have been considered as the most attractive power source because of their high energy density, high voltage, and long cycle life [1]. LiCoO_2 has been the most widely used positive electrode material since it was commercialized by Sony in 1991 [2]. Its further development, however, is seriously restricted by the toxicity, high cost, and instability, etc. [3], [4]. So, the $\text{LiNi}_{1/3}\text{Co}_{1/3}\text{Mn}_{1/3}\text{O}_2$, which was first reported by Ohzuku's research group in 2001, has attracted enormous attention due to its stable cycleability, safety, and high discharge capacity [5]-[11]. As far as we know, the $\text{LiNi}_{1/3}\text{Co}_{1/3}\text{Mn}_{1/3}\text{O}_2$ is one of the most promising candidates to replace LiCoO_2 cathode materials for LIBs [12]. In order to meet the requirements of the large-scale high-power system such as the electric vehicles (EVs), nevertheless, it is still a challenge to improve the structural stability and specific energy at high power rate and high voltage as the cycle properties and rate capability of this material become seriously deteriorated when charged to 4.6 V [12], [13]. Therefore, much research has been performed to optimize this series material by novel synthesis routes, using coating and employing metal substituted for Ni, Co and/or Mn [2], [14]-[24].

It is well-acknowledged that Al substitutions have a great impact on the cycle performance of layered lithium transition metal oxides [25], [26]. Dahn, J. R and other groups have

confirmed that the thermal stability of Al-substituted layered lithium transition metal oxides, such as $\text{LiNi}_{1/3}\text{Mn}_{1/3}\text{Co}_{(1/3-z)}\text{Al}_z\text{O}_2$ [27], [28], $\text{LiNi}_{1/3}\text{Mn}_{1/3}\text{Al}_{(1/3-x)}\text{Co}_x\text{O}_2$ [29], $\text{LiNi}_{0.5-z}\text{Mn}_{0.5-z}\text{Al}_{2z}\text{O}_2$ [30], $\text{Li}_{1-x}(\text{Ni}_{0.40}\text{Mn}_{0.40}\text{Co}_{0.2-z}\text{Al}_z)_{1-x}\text{O}_2$ [31], increases with increasing content of Al. Noted that Al in previous researches is usually employed to substitute part of Co. We are especially interested in substitution of Al for Mn, because the valence state of Al is lower than Mn, which may has a positive effect on the Li-ion diffusion. According to our knowledge, Al^{3+} substituted for Mn^{4+} in $\text{LiNi}_{1/3}\text{Co}_{1/3}\text{Mn}_{1/3}\text{O}_2$ has been reported [32], however, there has been no report about the advanced electrochemical properties at high voltage, especially high rate capabilities.

II. EXPERIMENTAL

A. Material Preparation

All chemical reagents were of analytical purity and used without further purification. Firstly, a Na_2CO_3 and NH_4HCO_3 aqueous solution (molar ratio = 1:1) and a mixed solution of $\text{Ni}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$, $\text{Co}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$, $\text{Mn}(\text{Ac})_2 \cdot 4\text{H}_2\text{O}$ and $\text{Al}(\text{NO}_3)_3 \cdot 9\text{H}_2\text{O}$ were simultaneously added drop-wisely to a batch reactor using a two-channel peristaltic pump. The molar ratio of Ni: Co: Mn: Al was 1: 1: (1-3x): 3x. The temperature, pH, and stirring speed are respectively controlled in 50-60°C, 8-9 and 800-1000 rpm. The carbonate precursors were filtered and washed thoroughly by centrifuging, and then was dried at 90°C overnight. The obtained powder and an appropriate of lithium carbonate were mixed completely by high energy mechanical milling. Lastly, the mixtures were firstly preheated at 500°C for 6 h to melt the lithium salt, then calcined at 900°C for 10 h in air. The products ($x = 0.00, 0.01, 0.02, 0.03, 0.04$) were denoted as Al 0, Al 1, Al 2, Al 3, Al 4, respectively.

B. Materials Characterization

The powder X-ray diffraction was made by using a Philip-produced PW 1730 diffractometer. XRD data were collected in the 2θ between 10° and 70° with a scanning speed of $0.06^\circ \text{ s}^{-1}$. Unit cell parameters were refined by the Rietveld method using the Jade 5.0 program. The scanning electron microscopy (SEM) was used to observe the morphology and particle size of the materials (S4800 machine).

C. Electrochemical Measurements

The electrochemical characterizations were tested in CR2032 coin type half-cell, which was comprised of a cathode, a lithium metal anode, a piece of porous polypropylene film and

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an electrolyte. The positive electrode materials contained an active material, carbon black, and PVDF binder in the weight ratio 80:13:7. The electrolyte employed to be analyzed was 1 mol L⁻¹ LiPF₆ solution in an EC-DMC (1:1 by volume) mixture. The fabricated coin cells were cycled at different current density between 3.0 – 4.6 V (vs Li/Li⁺) at room temperature. In this study, the capacity of 180 mA g⁻¹ was assumed to the 1 C ratio capacity.

III. RESULTS AND DISCUSSIONS

Fig. 1 shows the XRD patterns of the LiNi_{1/3}Co_{1/3}Mn_{1/3-x}Al_xO₂ ($x = 0 \sim 0.04$) samples in the scattering angle range from 10 to 70°. All of the peaks can be indexed to the hexagonal α -NaFeO₂ structure (R $\bar{3}m$ space group) and no impurity peaks are detected. It demonstrates that the LiNi_{1/3}Co_{1/3}Mn_{1/3-x}Al_xO₂ series are phase pure materials in the range of $x = 0.00 \sim 0.04$. In XRD patterns, the (006) / (102) and (108) / (110) doublet peaks are evidently split, which suggest a well-ordered layered structure [8], [19]. It should be noted that the (108) and (110) peaks become less sharp as x increases. The integrated intensity ratio of (003) and (104) peak is regarded as a measurement of the cation mixing in the layered structure. Because the radius of Li⁺ ($r_{Li^+} = 0.76 \text{ \AA}$) is approximate to the radius of Ni²⁺ ($r_{Ni^{2+}} = 0.69 \text{ \AA}$), the transition metal ions (Ni²⁺) at the 3a sites are could easily enter into the lithium metals at 3b sites. Ni²⁺ ions entering octahedral Li⁺ sites will produce NiO phase, which could weakens the intensity of the (003) peak but has no influence on (104) peak [33]. Thus, as the cation mixing is increased, the ratio of I_{003}/I_{104} is decreased. In General, undesirable cation mixing happens when I_{003}/I_{104} value is less than 1.2. Table I lists I_{003}/I_{104} value of LiNi_{1/3}Co_{1/3}Mn_{1/3-x}Al_xO₂ series, which are much larger than 1.2 for all samples. It can be observed that a small amount of Al substitution for Mn ($x \leq 0.02$) could restrain the cation mixing to some extent, especially when $x = 0.02$. Table I also gives the lattice parameters and structural parameters. The lattice parameters a , c and unit cell volume decreases monotonously with x increases because the radius of Al³⁺ ($r_{Al^{3+}} = 0.50 \text{ \AA}$) is smaller than that of Mn⁴⁺ ($r_{Mn^{4+}} = 0.53 \text{ \AA}$) [16]. The value of c/a , however, is the highest when $x = 0.02$, which indicates that LiNi_{1/3}Co_{1/3}Mn_{1/3-0.02}Al_{0.02}O₂ owns a better ordered hexagonal layered structure than other samples. Above all, we believe that Al has been incorporated successfully into the layered oxides crystal and Al 2 sample may deliver superior electrochemical properties.

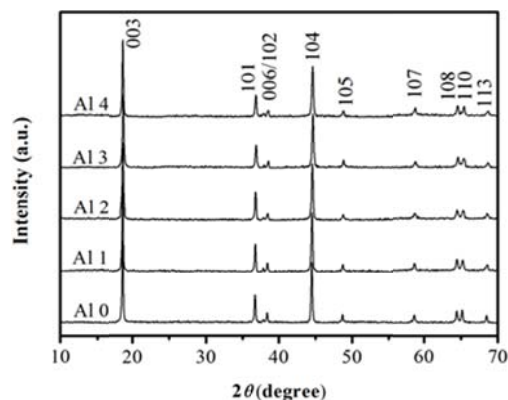


Fig. 1 XRD patterns of LiNi_{1/3}Co_{1/3}Mn_{1/3-x}Al_xO₂ samples (10 – 70°)

The surface morphology of the samples were observed using SEM, as shown in Fig. 2. The samples in Fig. 2 display extremely fine crystallites with spherical or polyhedral morphology, an average primary particle sizes are ~ 150 nm, and some of which are agglomerated into a bulk. The primary particle morphologies are similar for all prepared samples, but the particle agglomeration is seemingly aggravated as Al content increases, which are in accordance with the previous report [28], [34]. In general, the regular morphology of prepared particles is beneficial to the electrochemical performances.

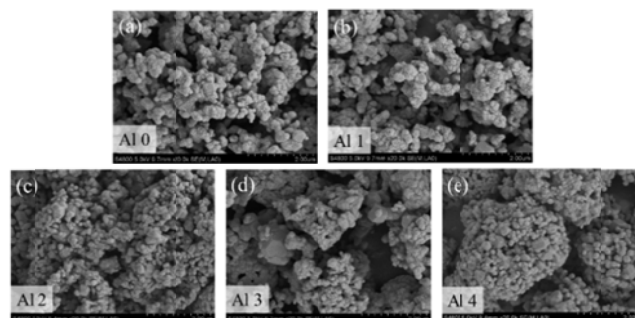


Fig. 2 SEM images of LiNi_{1/3}Co_{1/3}Mn_{1/3-x}Al_xO₂ samples

In order to study the impact of Al substitution on the electrochemical properties, the cells were first galvanostatically charged to 4.6 V and subsequently discharged to 3.0 V at a current density of 0.1 C (1 C = 180 mA g⁻¹) under room temperature. Fig. 3 shows the curves of specific discharge capacity vs cycle number for Li/LiNi_{1/3}Co_{1/3}Mn_{1/3-x}Al_xO₂ cells. Although Al 0 sample delivers the highest discharge capacity of 139.96 mAh g⁻¹ at 1 C, it suffers from a significant capacity drop during cycling and the capacity retention is 87.87% after 50 cycles. The poor cycling performances at high voltage mainly result from the structural instability, oxygen loss from the overlithiated oxide and side reaction caused by HF [2], [8], [25]. On the contrary, the Al doped electrodes exhibit superior cycleability though they own lower initial specific capacity. Among the Al doped samples, Al 2 shows the highest discharge capacity and has few capacity decrease during cycling. The initial discharge capacity is 136.17 mAh g⁻¹, and the capacity retention reaches 99.10% at the 50th cycle. Furthermore, the

TABLE I
LATTICE PARAMETERS AND STRUCTURAL PARAMETERS FOR
LiNi_{1/3}Co_{1/3}Mn_{1/3-x}Al_xO₂ SAMPLES

Samples	a (Å)	c (Å)	c/a	I_{003}/I_{104}	Unit cell volume (Å ³)
Al 0	2.8566	14.2206	4.9782	1.43	101.19
Al 1	2.8557	14.2178	4.9787	1.48	100.69
Al 2	2.8550	14.2150	4.9789	1.53	100.55
Al 3	2.8534	14.2033	4.9777	1.47	100.10
Al 4	2.8524	14.1955	4.9767	1.44	99.52

specific capacity of Al 2 exceeds that of Al 0 after 15 cycles. In addition, after 50 cycles, the capacity retention of Al 3 and Al 4 are 99.49% and 99.07%, respectively. The results confirm that Al constitution can efficiently enhance the cycling performance of materials.

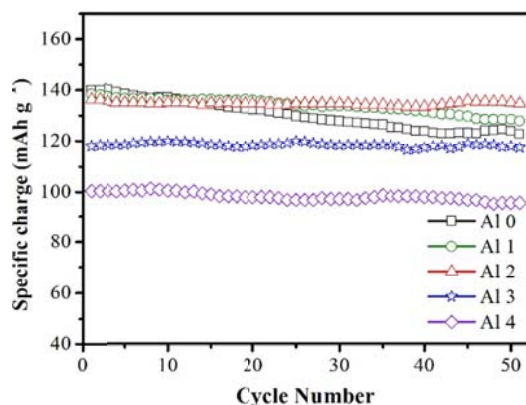


Fig. 3 Specific discharge capacity vs cycle number curves of Li/LiNi_{1/3}Co_{1/3}Mn_{1/3-x}Al_xO₂ cells in the potential range of 3.0 – 4.6 V at 1 C

IV. CONCLUSION

Al substituted layered LiNi_{1/3}Co_{1/3}Mn_{1/3-x}Al_xO₂ (x = 0 ~ 0.04) materials were successfully synthesized via a co-precipitation and annealing process. All samples have a hexagonal α -NaFeO₂ structure and no impurity phases are formed. As the Al content increases, the lattice parameters *a*, *c* and unit cell volume decrease correspondingly. The LiNi_{1/3}Co_{1/3}Mn_{1/3-x}Al_xO₂ materials show good cycling performance at high voltage. The Al-free sample has higher initial capacity, but shows inferior cycleability than Al-doped samples. The LiNi_{1/3}Co_{1/3}Mn_{1/3-0.02}Al_{0.02}O₂ (Al 2) material delivers an initial discharge capacity of 136.17 mAh g⁻¹ between 3.0 and 4.6 V at 1 C rate, and the capacity retention ratio after 50 cycles is 99.10 %. The superior electrochemical performances along with the better thermal stability of Al 2 make it a promising high power positive electrode material for lithium ion batteries.

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