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P2-Na_xCo_{1-y}Ti_yO₂: a high performance cathode material for sodium ion batteries

Noha Sabi^{a,b}, Angelina Sarapulova^c, , Sylvio Indris^c, Helmut Ehrenberg^c, Jones Alami^b and Ismael Saadoune^{a,b}

^{*a}LCME, FST Marrakesh, University Cadi Ayyad, Av. A. Khattabi, BP 549, 40000, Marrakech, Morocco*</sup>

^bMaterials Science and Nano-engineering Department, Mohammed VI Polytechnic University, Ben Guerir, Morocco

^c Institute for Applied Materials (IAM), Karlsruhe Institute of Technology (KIT), Hermann-von-Helmholtz Platz 1, 76344 Eggenstein-Leopoldshafen, Karlsruhe, Germany

Abstract:

Sodium ion batteries have attracted a wide attention in recent years thanks to the tempting properties that sodium, the main component of this technology, offers. It is worth noting that sodium has a homogeneous distribution in the earth crust and exhibits similar reaction mechanism to the conventionally used lithium [1].

Different compounds, in particular layered oxides, are successfully used as cathode materials for Lithium ion batteries. Oxides are, therefore, studied in the present work and their suitability for sodium ion batteries is investigated. LiCoO₂, typically used in Li-based batteries, was replaced by Na_xCoO₂. The intercalation/deintercalation of sodium ions in this material occurs with the existence of well-defined steps in the potential window 2.0 - 3.8 V, i.e. during the charge/discharge process. The existence of potential steps in V vs composition curve, is due to structural transitions, thus affecting the electrochemical process. [2]. Knowing that Na_{0.66}Co_{0.5}Ti_{0.5}O₂ exhibits a reduced number of plateaus in the potential range 2-4.2 V [3], the present work aims to investigate how the change of Co/Ti composition ratio in the cathode affects the performance of the Na-ion battery.

 $Na_yCo_{1-x}Ti_xO_2$ reveals a P2-type structure that delivers a first discharge capacity of 116 mAh/g with a good capacity retention. In-situ synchrotron XRD shows that the P2 type structure is practically preserved during the cycling, confirming judicious choice of titanium substitution.



Figure 1: Charge and discharge curves of $Na_xCo_{1-y}Ti_yO_2$ and the evolution of the (002) peak, data from the MSPD beamline at ALBA.

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[1] Su, H., Jaffer, S., Yu, H. Transition metal oxides for sodium-ion batteries, Energy Storage Materials, 116–131(2016)

[2] Berthelot, R., Carlierand, D., Delmas, C. Electrochemical investigation of the P2–Na_xCoO₂ phase diagram, nature materials **10**,74–80 (2011)

[3] Sabi, N., Doubaji, S., Hashimoto, K., Komaba, S., Amine, K., Solhy, A., Manoun, B., Bilal, E., Saadoune, I., Layered P2-Na_{2/3}Co_{1/2}Ti_{1/2}O₂ as a high-performance cathode material for sodium-ion batteries, Journal of Power Sources, (2017)

Contribution: Oral