Aberration corrected STEM and High Resolution EELS study Investigating Magnesium Intercalation in Vanadium Pentoxide Cathode

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Magnesium ion based batteries hold promise as a competitive alternative to conventional Lithium ion battery technology due to several key features. Theoretical volumetric capacity for magnesium metal anode is much higher compared to lithium metal. Furthermore, Mg is more readily available compared to Li, which can potentially lead to cost reduction and switching to Mg offers safety benefits over Li as well. Orthorhombic V₂O₅ is a well-known intercalation cathode host for Mg-ion batteries owing to its characteristic layered structure and weak vanadium oxygen bonding that facilitates ion intercalation between the layers.

The following contribution will focus on systematic characterization of electrochemically cycled thin film orthorhombic V₂O₅ cathode host to verify Mg intercalation employing aberration corrected scanning transmission electron microscopy (STEM) imaging. Our results indicate that upon electrochemical cycling the ε-phase is formed in agreement with previous density functional theory calculations.[¹][²] This is remarkably different from the δ-phase which is commonly observed in direct solid state synthesis of MgV₂O₅. Though the intercalation levels are low (which is possibly because the formation of fully magnesiated delta phase would involve structural rearrangement requiring high interfacial energy leading to low rates), this study aims to directly probe the Mg intercalation sites in orthorhombic V₂O₅, which is predicted to be one of the “beyond chevrel” phase high-voltage cathode host for Mg ion batteries. Multislice image simulations using the Kirkland code were also performed for the ε-MgₓV₂O₅ crystal in [001] orientation which closely matches our experimental data as well. Figure 1(a) and 1(b) present atomic resolution high angle annular dark field (HAADF) and Annular Bright field (ABF) images respectively for the cycled thin film orthorhombic V₂O₅ cathode. Simulated results of HAADF and ABF images are also shown in Fig 1(c) and Fig 1(d) respectively. The structural model for the electrochemically cycled ε-MgₓV₂O₅ phase is shown in Fig 1(e) while the structure for chemically synthesized δ- MgV₂O₅ is presented in Fig 1(f).

It was also recently reported that a new tunnel-structured polymorph of V₂O₅ (ζ-V₂O₅) has been stabilized, which could serve as a better cathode host for Li and Mg intercalation due to lower band gap.[³] Results will be presented discussing the detailed structure of the ζ-V₂O₅ nanowire using aberration-corrected STEM imaging and electron energy loss spectroscopy (EELS) and Mg intercalation sites will be investigated in these ζ-V₂O₅ nanowires after electrochemical cycling as well. In addition, the chemically-synthesized β-MgₓV₂O₅ nanowires will also be studied in order to compare the Mg intercalation sites with the electrochemically cycled ζ-V₂O₅. Figure 2(a) presents atomic-resolution HAADF image for ζ-V₂O₅ nanowires clearly showing the tunnel structure geometry [as shown by the structure in Fig 2(b)] and the EELS spectra [shown in Fig 2(c)] verifies the valence state of vanadium as V⁵⁺ (by comparing the energy difference between V L₃- and O K- edge onset) as expected.[⁴]
References
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Figure 1: (a) Filtered atomic-resolution HAADF image and (b) Filtered ABF image (c) Multislice image simulation of HAADF image and (d) ABF image for ε-Mg$_x$V$_2$O$_5$ crystal along [001] zone (e) ε-Mg$_x$V$_2$O$_5$ and (f) δ-MgV$_2$O$_5$ structure.

Figure 2: (a) Atomic resolution (filtered) HAADF image of ζ-V$_2$O$_5$ nanowire and (b) ζ-V$_2$O$_5$ structure showing only V atoms (c) EELS spectra from ζ-V$_2$O$_5$ nanowire