Particles and Binder Interaction in the Lithium-ion Cell Electrodes



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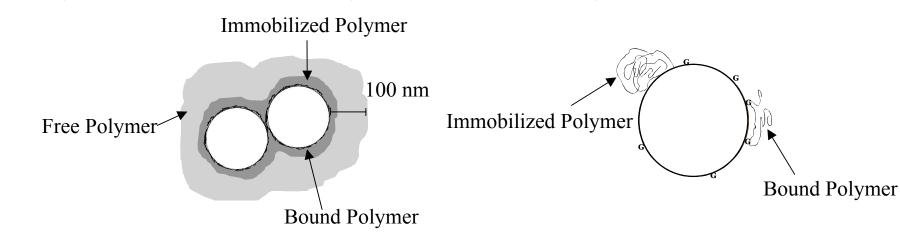
Introduction

Proper electrode design is critical to meet high power performance requirements for lithium-ion rechargeable battery applications. Binders and conductive additives, although not electrochemically active, are essential components in the electrodes. In a simple three component cathode system where polyvinylidene difluoride (PVDF) is used as a binder, along with

active material and acetylene black (AB) conductive additive, the dominant interaction between polymer and active material or polymer with AB changes proportionately with particle surface area. The PVDF binder interaction with the active material and AB on the micro or nano-scale plays an important role in determining the battery performance.

Particles A Cathode Electrode Binder iam.: 10 µm irface: 0.78 m²/g **Interaction** ontent: 90% (w%) Essential cale bar: 10 μm Content: 4% (w%) Scale bar: 100 nm Polymer Binder 4% (w%) NCA Electrode Scale bar: 4 µm Optional L 1-10 μm Diam.: 7.5 μm W~0.1-0.5 μm Surf.: 14.8 m²/g Surf.: $15.1 \text{ m}^2/\text{g}$ Scale bar: 1 µm

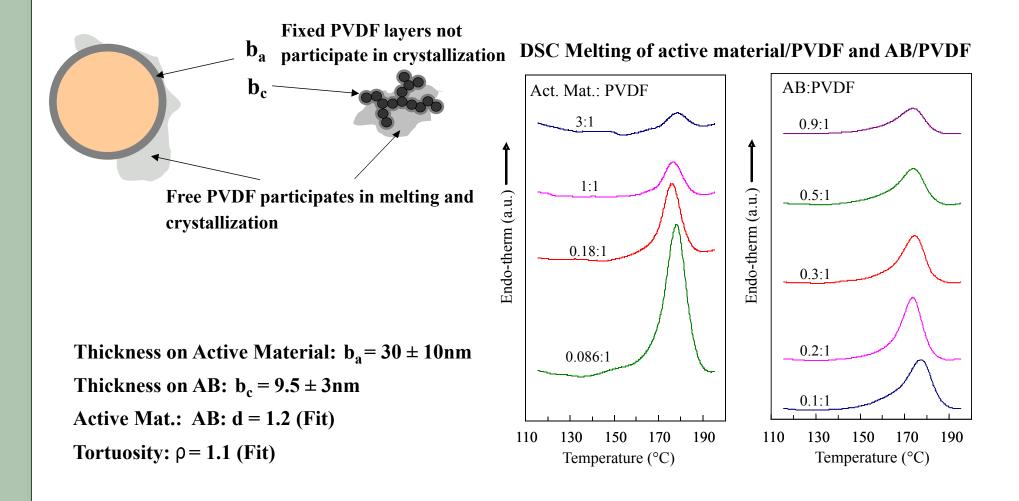
Three Physical States of Polymer in Contact with Particles Bound Polymer, Immobilized Polymer, and Free Polymer



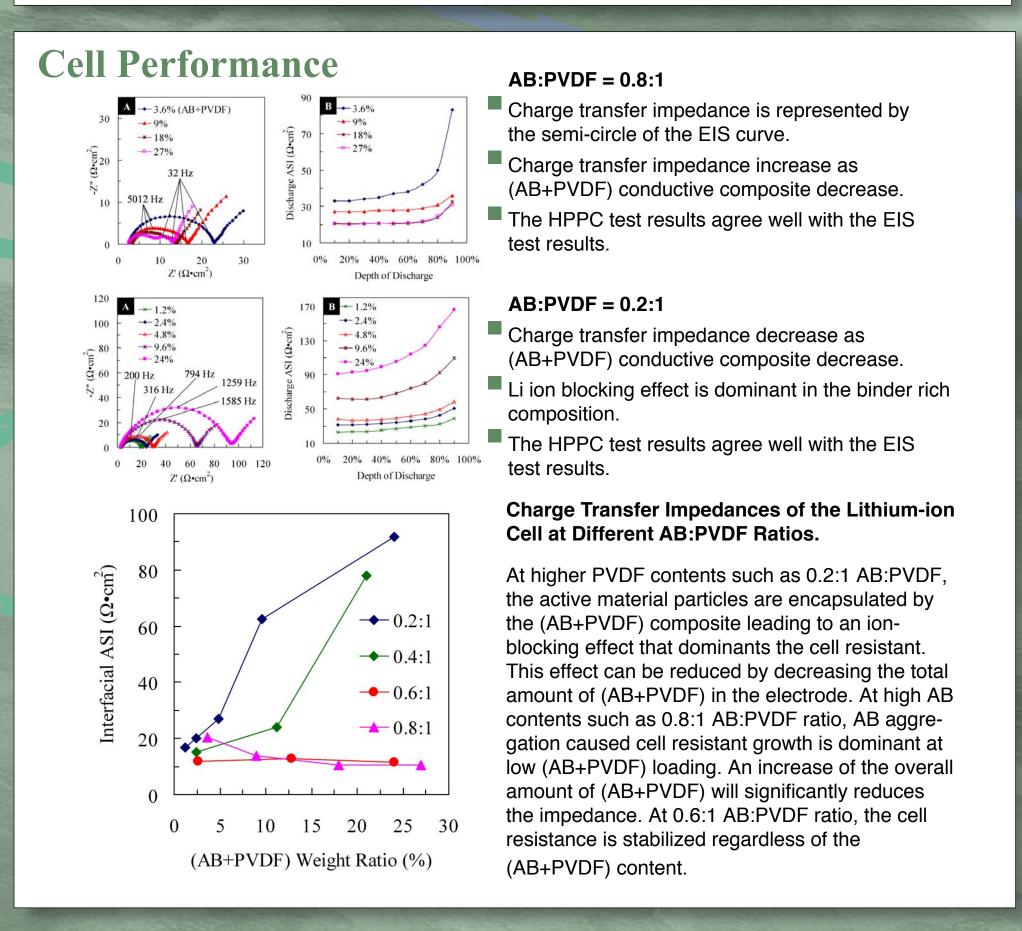
Bound and immobilized polymer are fixed layer on particles!

Nano-scale Interaction in the Electrode Composites AB/PVDF Conductive Matrix to Provide **Active Material Particles Electronic Conductivity Polymer binder AB** particles Conductivity Matrix 0.4:1 0.6:1 Acetylene Black:PVDF (by weight) $\mathbf{k} = \mathbf{k}_0(\mathbf{c}/\mathbf{b})$ $k_0(c/b)$ Active material and AB competing for binder **Changed conductivity** per unit volume ba: Fixed binder on **Active materials** Losing binder in the conductive composite $\mathbf{k} = \mathbf{k}_0(\mathbf{c}/(\mathbf{b}-\mathbf{b}_a))$

DSC Method to Determine Parameters: b_a & b_c



Electrode Electronic Conductivity Not Enough Binder **Electronic Conductivity** & Modeling Low impedance High impedance Li⁺ transport at the active material interface is affected by binder. $k_e = (c+b-d*b_a)^{\rho*}k_0(c/(b-d*b_a))$ $k_e = (c+b-b_a)^{\rho} * k_0(c/(b-b_a))$ Increase electronic conductivity d: Competing factor between active material and AB **Modeling and Experimental Results of the Electronic Conductivity of the Composite** → AB:PVDF 0.6:1 (M) △ (E) Electrode. DC-conductivity measured by 4-probe. Modeling was taken consideration of porosity and tortuosity factors of the AB/PVDF conductive matrix. The electronic conductivity is higher with AB:PVDF = 0.4:1; 0.6:1 compare to 0.8:1 withhigh loading of active materials. Active Material Fraction (wt.) Increase ionic conductivity



Conclusions

- Binder plays critical roles in the composite electrode for providing not only mechanical integrity, but also electronic conductivity.
- The competition for binder between the active material and acetylene black is a fundamental factor affects cell performance.
- High acetylene black content such as AB:PVDF = 0.8:1 tends to produce electrode with lower electronic conductivity at high active material loading, but tend to exert minimum Li ion blocking.
- Intermediate acetylene black content such as AB:PVDF = 0.6:1 tends to produce electrode with higher electronic conductivity at high active material loading, with constant low Li ion block effect.

Acknowledgments

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