## Abstract Submitted for the MAR08 Meeting of The American Physical Society

## Ion Conduction and Polymer Dynamics of Poly(2-vinylpyridine)

- Lithium Perchlorate Mixtures<sup>1</sup> PORNPEN ATORNGITJAWAT, Burapha University, JAMES RUNT, Penn State University — Ion conduction and polymer dynamics of single phase mixtures of poly(2-vinylpyridine) (P2VPy) with 0.1 to 10 mol% lithium perchlorate (LiClO<sub>4</sub>) were investigated using broadband dielectric spectroscopy. Interpretation of the relaxation behavior was assisted by findings from wide-angle and small-angle X-ray scattering experiments, and other techniques. Five dielectric relaxations were observed: a local  $\beta$  process in the glassy state, a segmental relaxation, a slow segmental process, an ion-mode relaxation, and electrode polarization. The local P2VPy relaxation was strongly suppressed with increasing LiClO<sub>4</sub> content arising from the formation of transient crosslinks, which lead to a subsequent decrease in the number of free pyridine groups, and/or a reduction in the local free volume in the presence of LiClO<sub>4</sub>. Ion conduction at low LiClO<sub>4</sub> concentrations (< 10 mol%) is governed by the diffusion of anions through the matrix, which is strongly coupled with the segmental relaxation. At relatively high LiClO<sub>4</sub> concentration (10 mol%), partial decoupling between ion motion and the segmental relaxation was observed, leading to increased conductivity.

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