Hot-pressed, nanocomposite, **PEO-based** dry, membranes. Ionic conductivity electrolyte characterization and battery tests.

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Poly(ethylene oxide) (PEO)-based lithium electrolyte membranes appear the most suitable candidates as separators for reliable, lithium rechargeable polymer batteries¹. Nevertheless, the PEO-LiX complexes perform well in terms of ionic conductivity only when the polymer is mainly in the amorphous state², i.e., when the mechanical properties of the polymer electrolytes are relatively poor.

In this work we report the results of the ionic characterizationof conductivity PEO-based. nanocomposite, polymer electrolytes prepared by hotpressing through a solvent-free procedure. Several P(EO)_nLiCF₃SO₃:filler electrolyte compositions were developed and characterized. In particular, the investigation was focused on the effect of the temperature, the PEO molecular weight, the EO/Li molar ratio, the nature and the content of filler. Nanoscale (< 7 nm) Al₂O₃ and SiO₂ were selected as fillers. Lithium polymer composite electrolytes, formed by a blend of poly(ethylene oxide) (PEO), LiCF₃SO₃ lithium salt and a selected, nanoparticle ceramic filler, were prepared by hot-pressing through a solvent-free procedure. These dry, ionically conducting membranes were characterized in terms of ionic conductivity in the 30-105°C temperature range. The results, reported in Figure 1, demonstrate the favorable basic properties of these membrane, which show an ionic conductivity higher than 10^{-4} Scm⁻¹ at 70°C, as well as high homogeneity and excellent mechanical properties.

The hot-pressed, composite PEO membrane were used as electrolyte separator in all-solid, lithium polymer cells by laminating a lithium foil anode, a selected hot-pressed, PEO electrolyte membrane and a LiFePO₄ composite cathode film prepared by hotpressing through a completely dry, solvent-free procedure. Among the materials under development as cathode in lithium polymer batteries, the lithium iron phosphate, LiFePO₄, originally proposed by Goodenough³, may regarded as very promising candidate due its capability to operate with a very flat voltage plateau. In addition, LiFePO₄ exhibits a large theoretical specific capacity (about 170 mAhg-1), thermal stability and benefits of low cost and environmental compatibility. The performance of the polymer battery was evaluated at different current densities and temperatures. The results are shown in Figure 2.

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References

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Fig. 1. Arrhenius plots of hot-pressed, nanocomposite, $P(EO)_n LiCF_3 SO_3$: x% wt filler electrolyte membranes. Panels from A to D are referred, respectively, to a filler content as well as 5% wt Al₂O₃ (panel A), 5% wt SiO_2 (panel B), 10% wt Al_2O_3 (panel C), and 10% wt SiO_2 (panel D). Each panel reports the data for different PEO molecular weights and EO/Li molar ratios.

Fig. 2. Delivered capacity vs. temperature dependence for a Li / PEO:LiCF₃SO₃:SiO₂ / LiFePO₄ polymer battery at different current densities (see legend).



