

Boron Nitride Modulates Polymer Electrolyte Conductivity

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The case for safer sodium-ion electrolytes

- Sodium is 100x less expensive than lithium
- Current electrolytes are flammable and risk thermal runaway
- Safer electrolytes are important for grid-scale energy storage



Thermal runaway is a major battery safety issue

Composite polymer electrolytes are promising

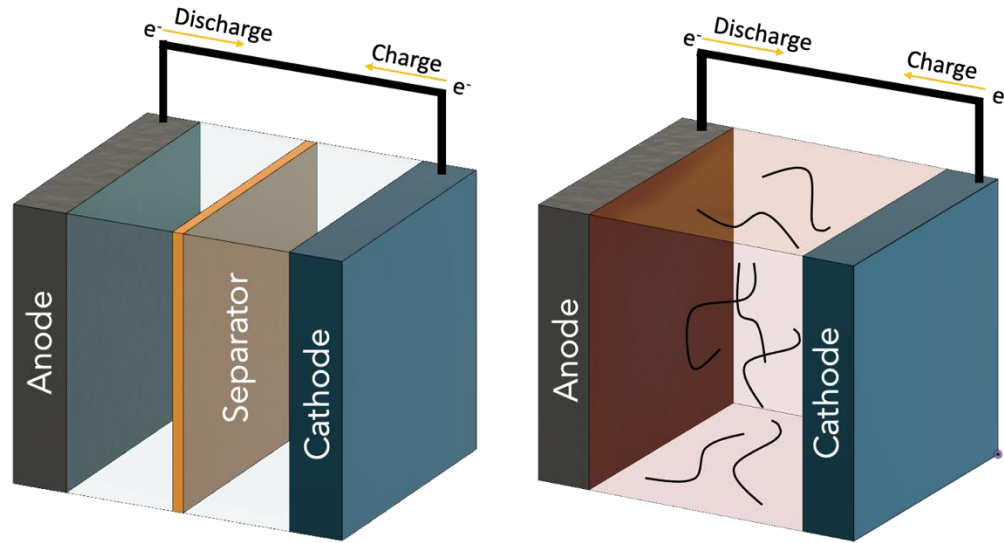


Fig. 1. Liquid (left) vs. polymer (right) electrolyte

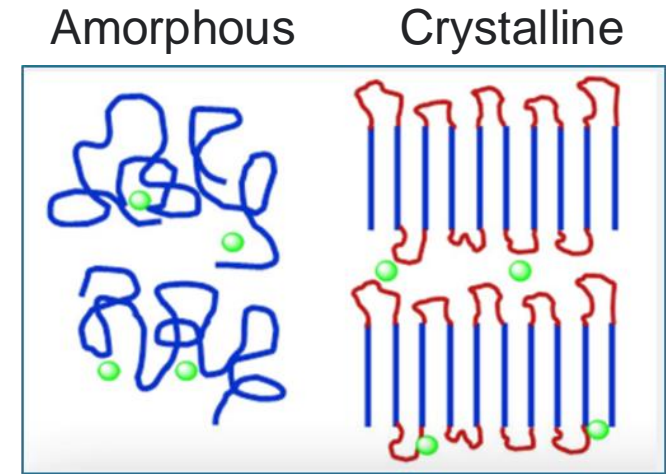
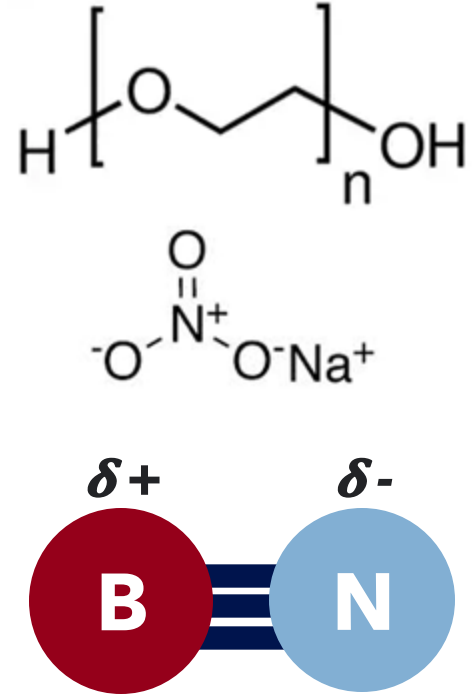


Fig. 2. (adapted from Cheng et al., 2014). Ions can only move through amorphous domains

Boron nitride is an interesting CPE filler

- Hexagonal boron nitride (BN) may interact with Na^+ and NO_3^- ions
- BN increases the transference number of PEO–LiTFSI electrolytes
 - Li^+ –BN binding energy: -157.6 kJ/mol
 - TFSI–BN binding energy: -166.51 kJ/mol



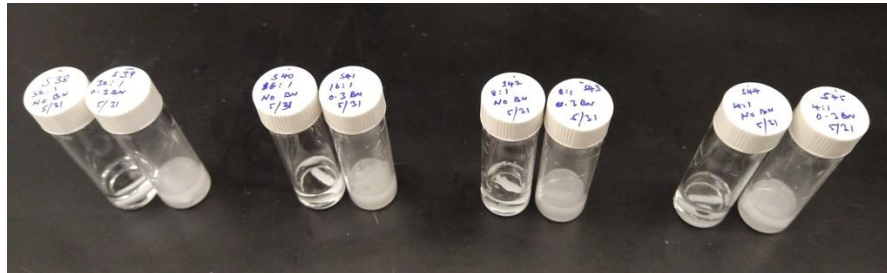


Methods

Phase 1: Prepare and cast solutions



- PEO, NaNO_3 , and h-BN mixed in water
- 0.3 wt.% and 3 wt% h-BN, 4:1 and 24:1 ether $\text{O}:\text{Na}^+$ ratios, and controls (9 samples)
- Solutions cast on 1 in² glass slides (381 μm -thick)



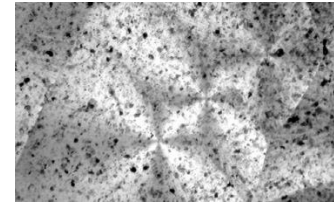
Phase 2: Dry in vacuum oven



- Hot plate at 120°C for 1h
- Vacuum oven at 120°C for 16h
- Vacuum chamber at 25°C for 1h



PEO Film, 25 mm²



PEO crystal with 3% BN,
1.5 mm²

Phase 3: Characterize Samples



Structural Characterization

- Differential Scanning Calorimetry (DSC)
- X-Ray Diffraction (XRD)
- Fourier Transform Infrared Spectroscopy (FTIR)

Electrochemical Characterization

- Electrochemical Impedance Spectroscopy (EIS)



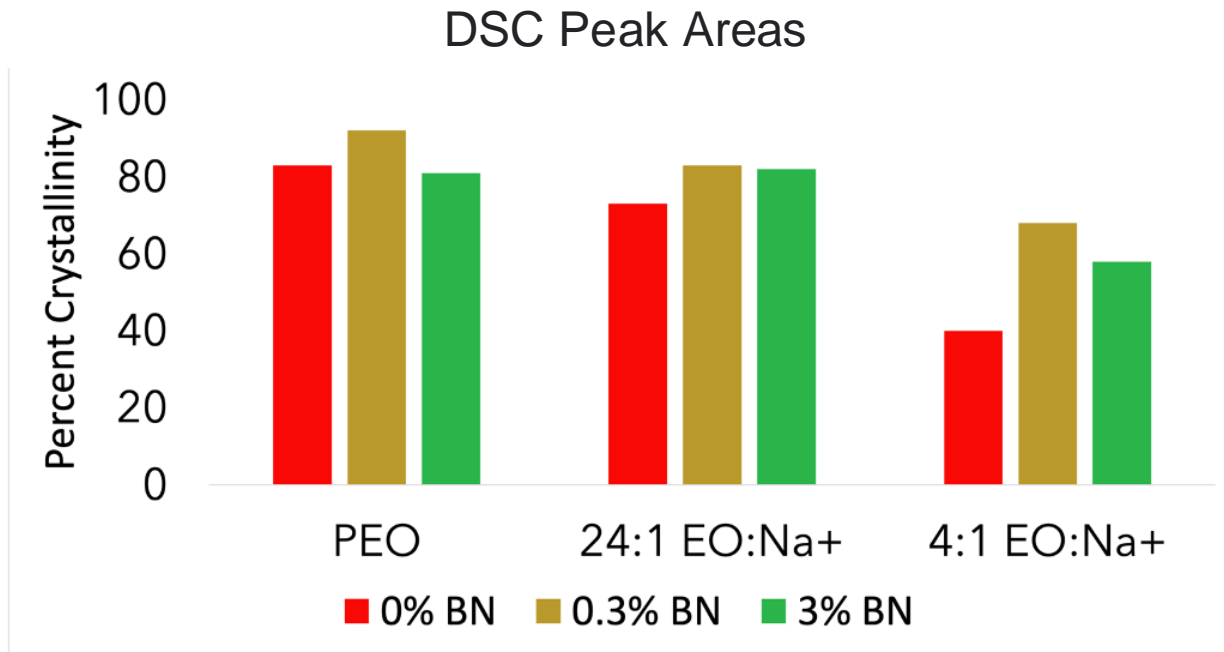
Cryostat (EIS)



Results

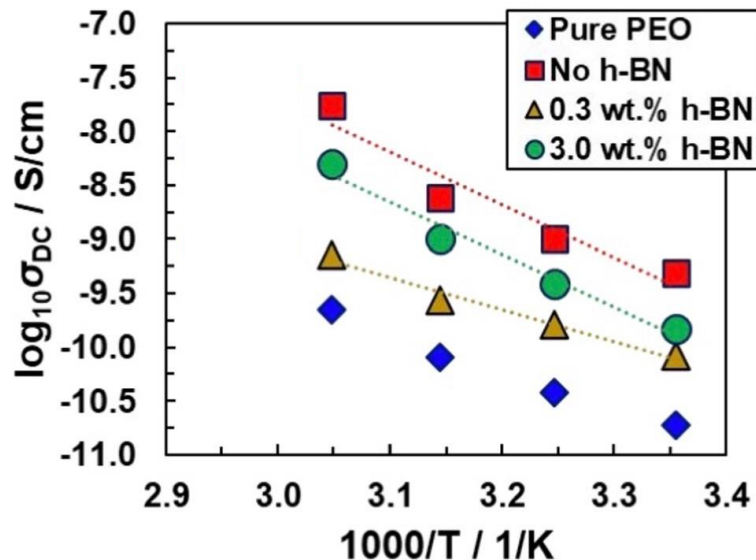
Adding BN affects PEO crystallinity

- **Differential Scanning Calorimetry** shows heat of melting
- Increasing BN to 0.3% increases crystallinity
- Crystallinity decreases at 3%
- Overall increased crystallinity could hinder ion mobility

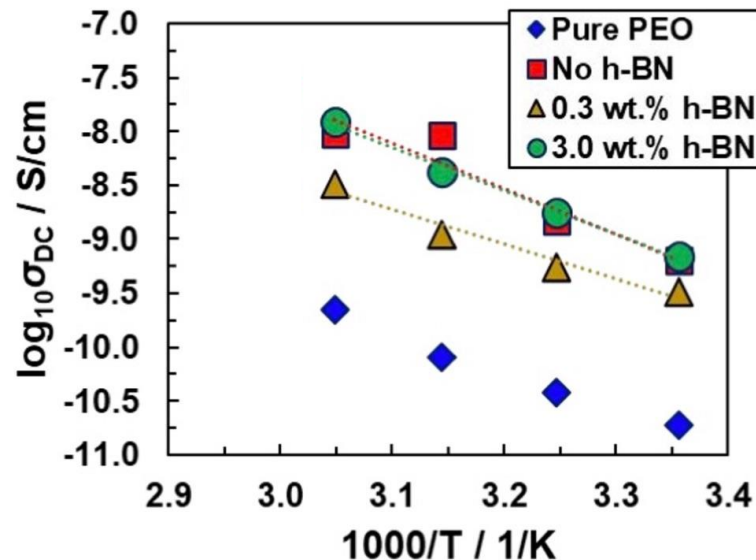


Crystallinity affects ionic conductivity

a) $(\text{PEO})_{24}:\text{Na}^+$



b) $(\text{PEO})_4:\text{Na}^+$

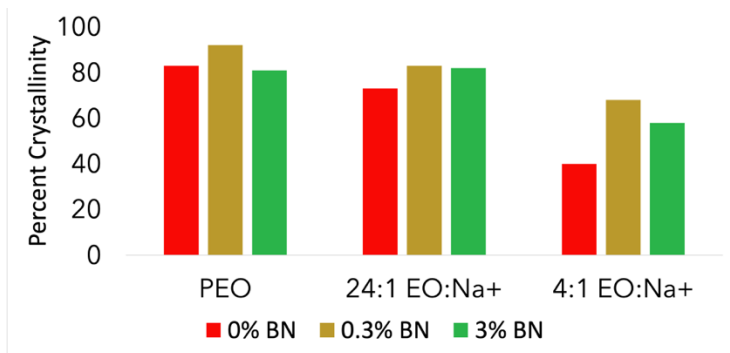


Ionic conductivity is highest for BN-free samples, showing the effect of increased crystallinity

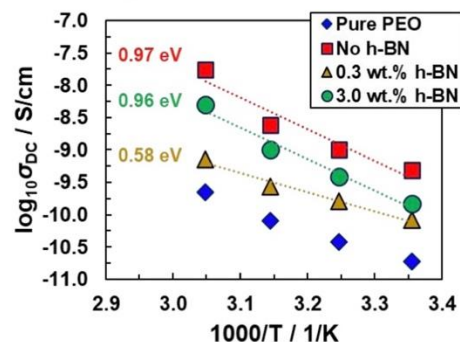
Conclusions

- Adding BN can increase PEO crystallinity via enhanced nucleation
- The effect of increased crystallinity dominates, decreasing ionic conductivity
- Sodium CPE properties may be tailored by changing the geometry of filler materials

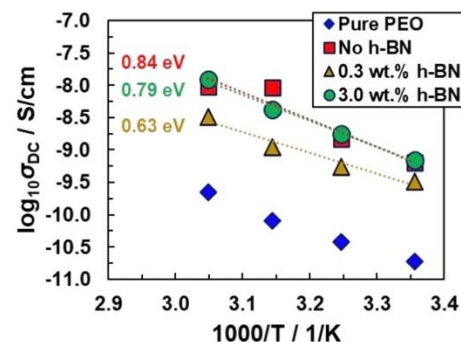
DSC Peak Areas



a) $(\text{PEO})_{24}:\text{Na}^+$



b) $(\text{PEO})_4:\text{Na}^+$



Acknowledgements

I'd like to thank the following people and organizations for making this project possible: Vagelos Integrated Program in Energy Research (VIPER), NSF grant FMRG-2134715, Ben Ferko, Steve Szewczyk, Mohamed Hassan, Professor Eric Detsi, Katie Sun, and my mentor, Shreyas Pathreker.





Supplementary Information

Phase 4: Compare to DFT Calculations

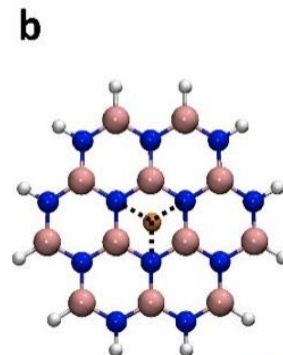


- Significant BN-salt binding energies
- Trends differ from lithium-ion literature

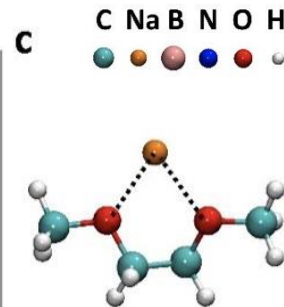
Complexation Energy (kcal/mol)



BN-NO₃⁻
-13.4



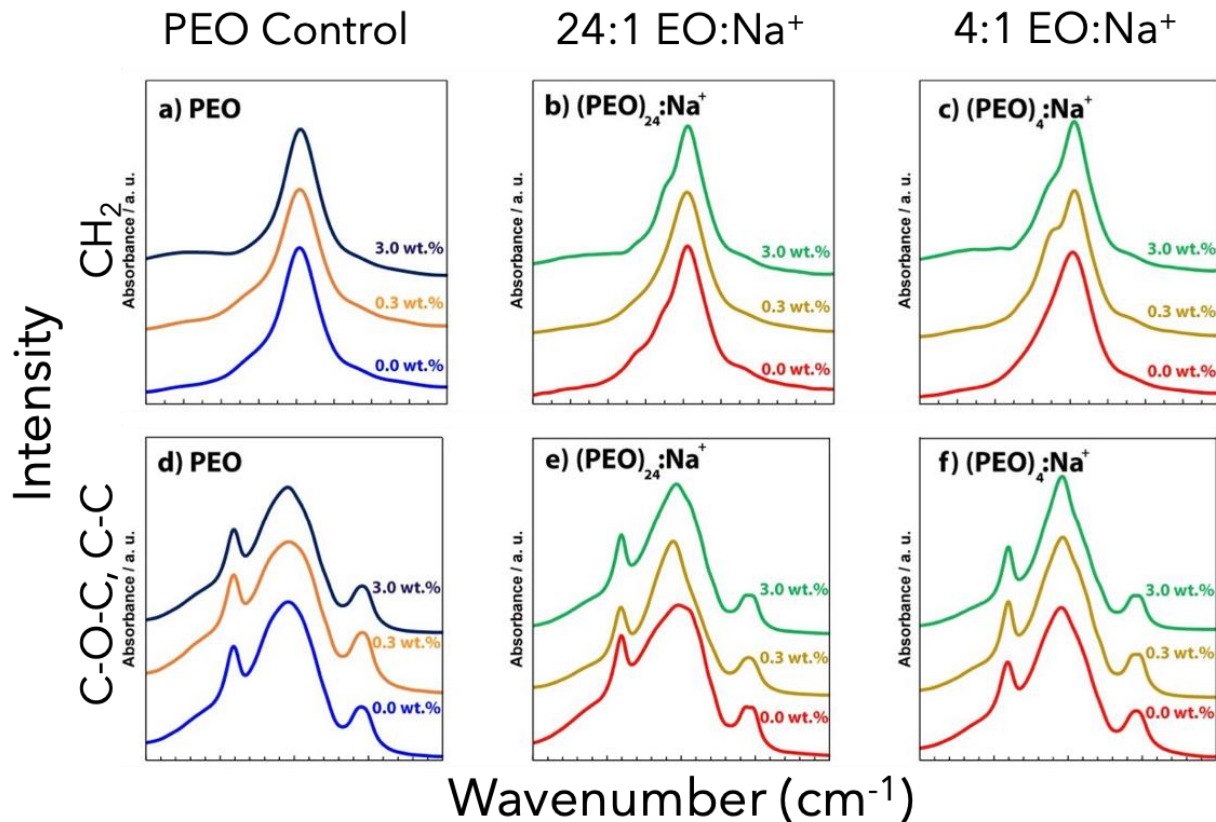
BN-Na⁺
-29.3



PEO-Na⁺
-44.1

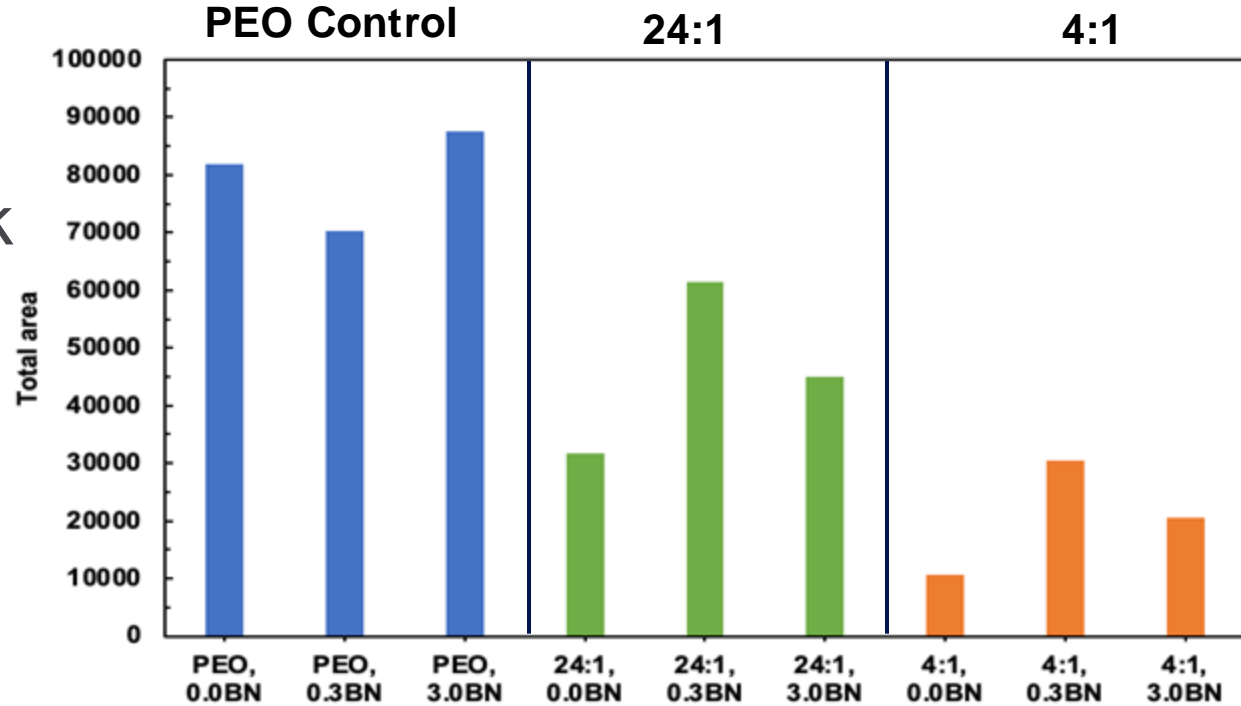
BN decreases PEO- NaNO_3 complexation

- PEO complexes with Na^+ , decreasing crystallinity
- FTIR peaks (840 and 1100 cm^{-1}) broaden with salt
- Peaks narrow when BN is added



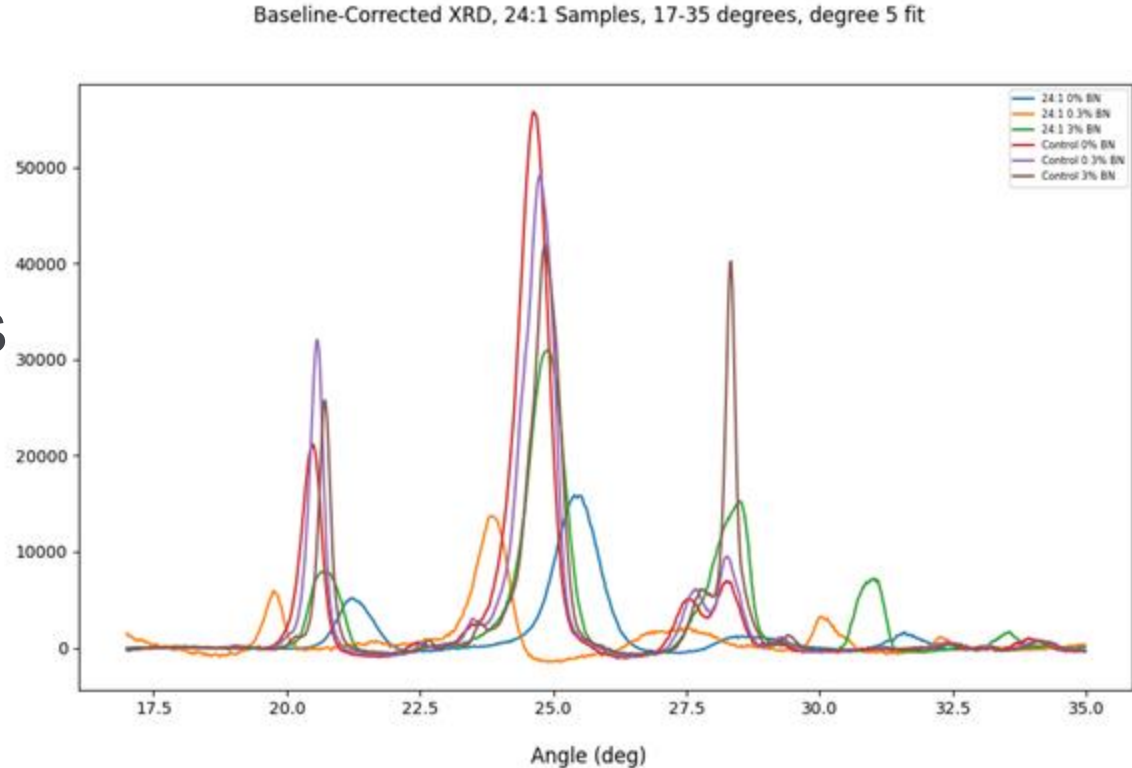
XRD peak area varies with h-BN loading

For salt-doped samples, total peak area (crystallinity) increases with 0.3% h-BN then decreases with 3% h-BN



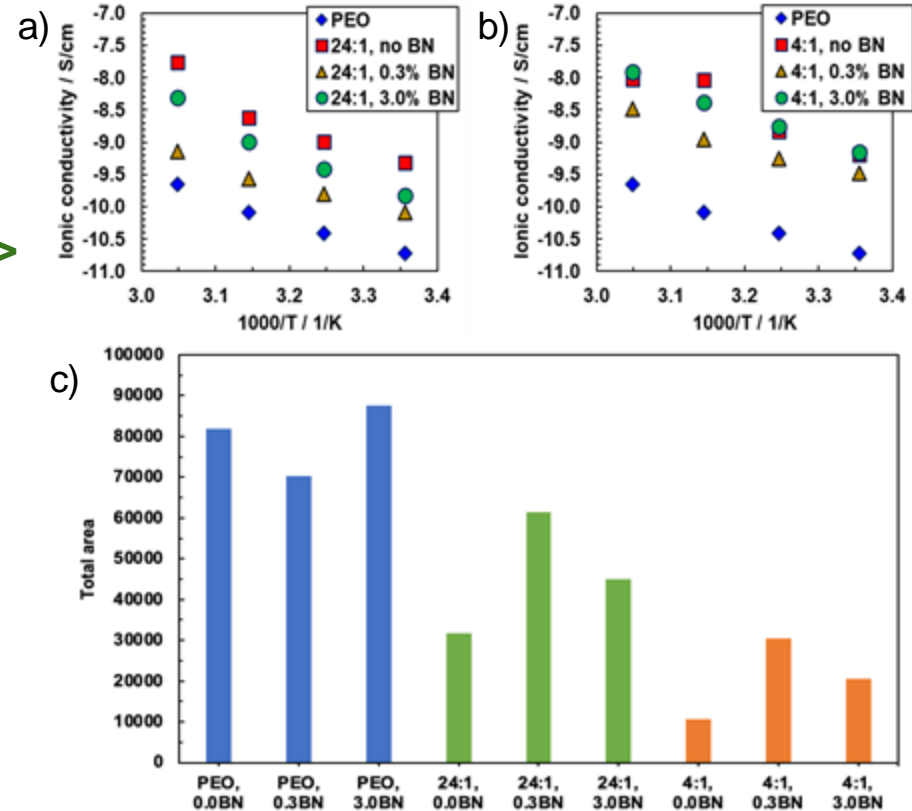
Additional XRD from Trial 2

No significant NaNO_3 peaks appear in the 24:1 XRD traces

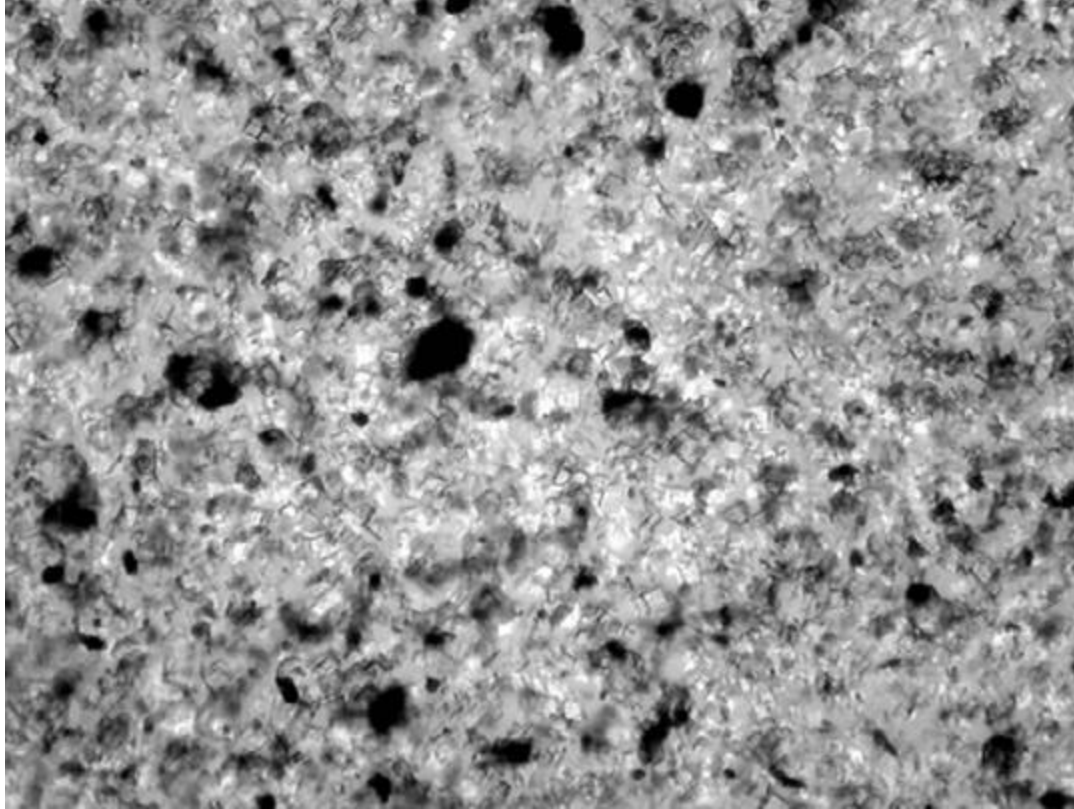


Ionic Conductivity and XRD Peak Area

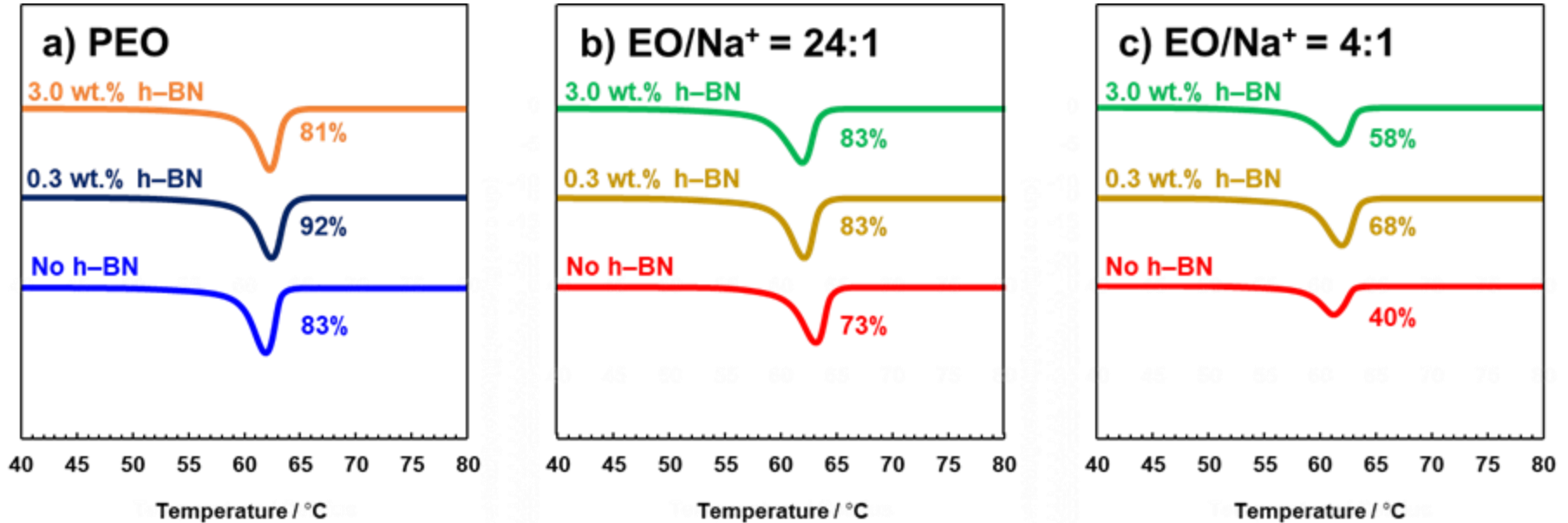
- Ionic conductivity follows the trend $IC_{No\ h-BN} > IC_{3.0\%} > IC_{0.3\%}$
- Interestingly, crystallinity follows the inverse trend $X_{No\ h-BN} < X_{3.0\%} < X_{0.3\%}$



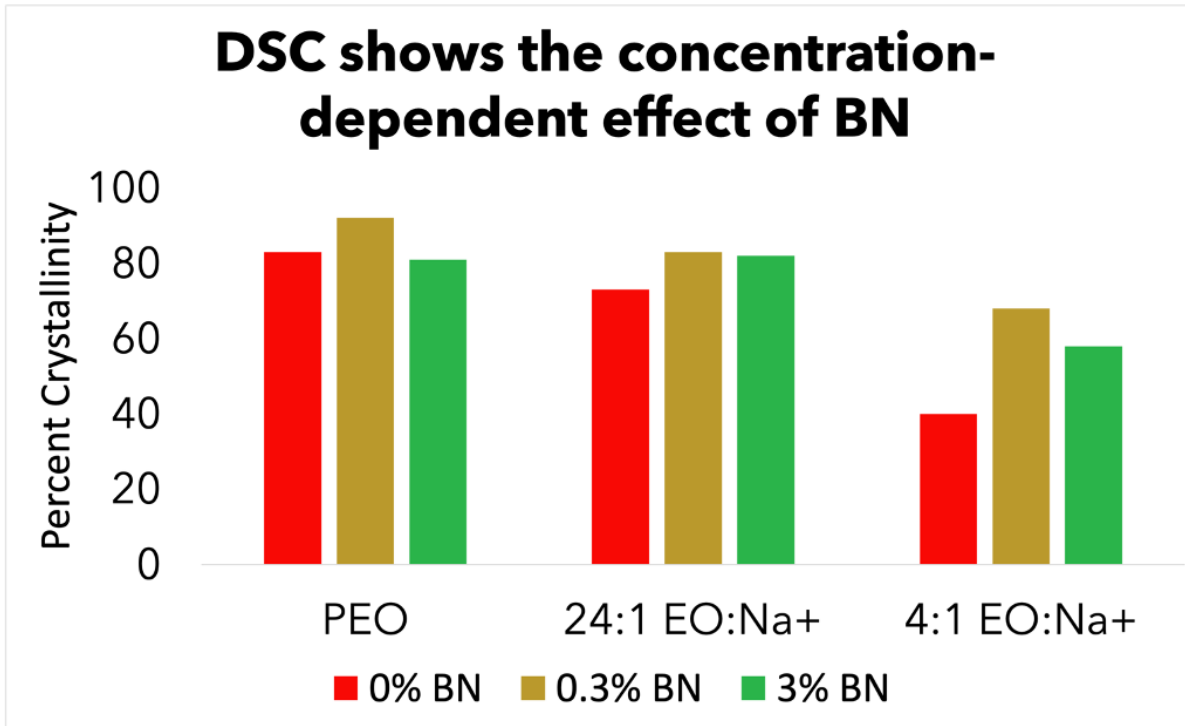
Salt Crystal image (4:1 NaNO₃, 3% BN)



DSC on polymer electrolytes



DSC shows the concentration-dependent effect of BN



PEO- NaNO_3 system is soluble in H_2O

