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Using the Principles of Electrochemistry to Understand and Overcome the Complicated Degradation Mechanisms of Silicon Anodes in Lithium-Ion Batteries.

Mike Carroll

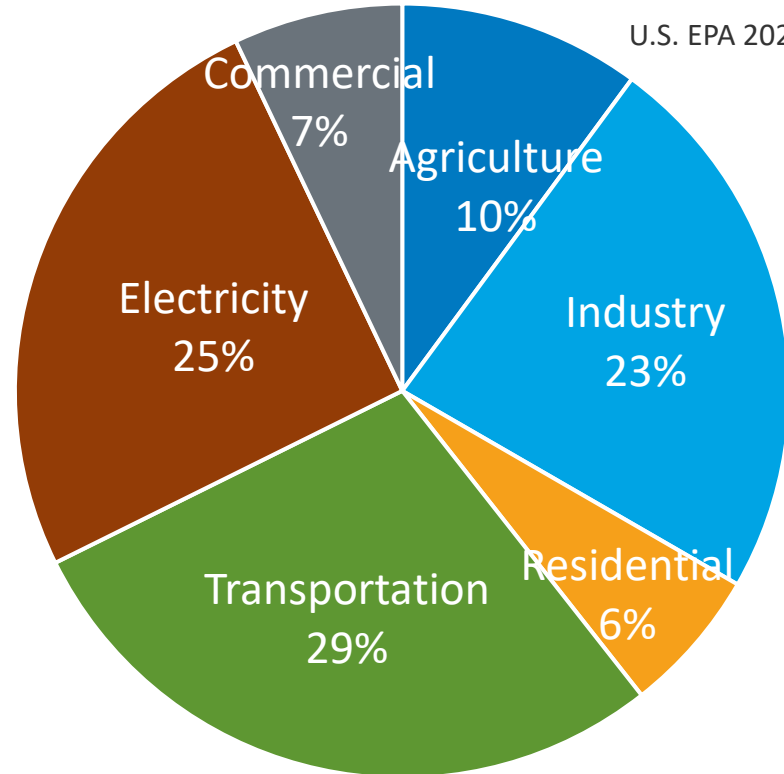
6/4/2024

2024 GRC Inorganic Chemistry

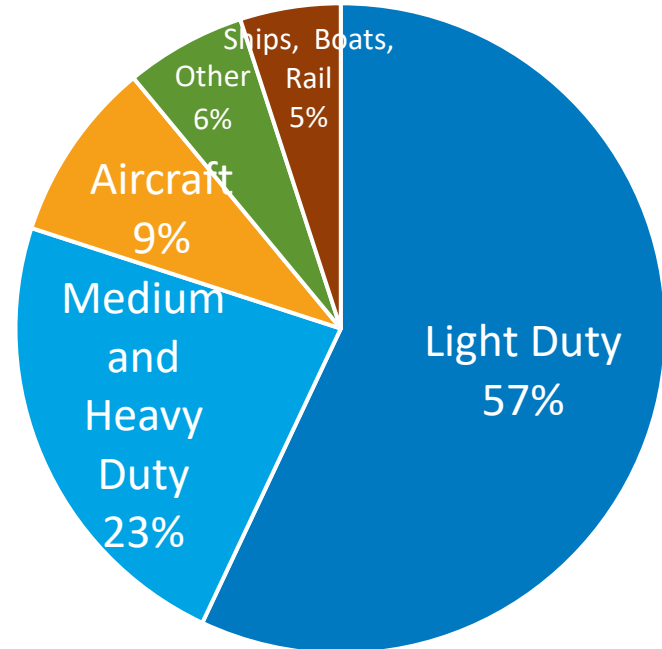
Decarbonization

US GHG emissions by sector

U.S. EPA 2022

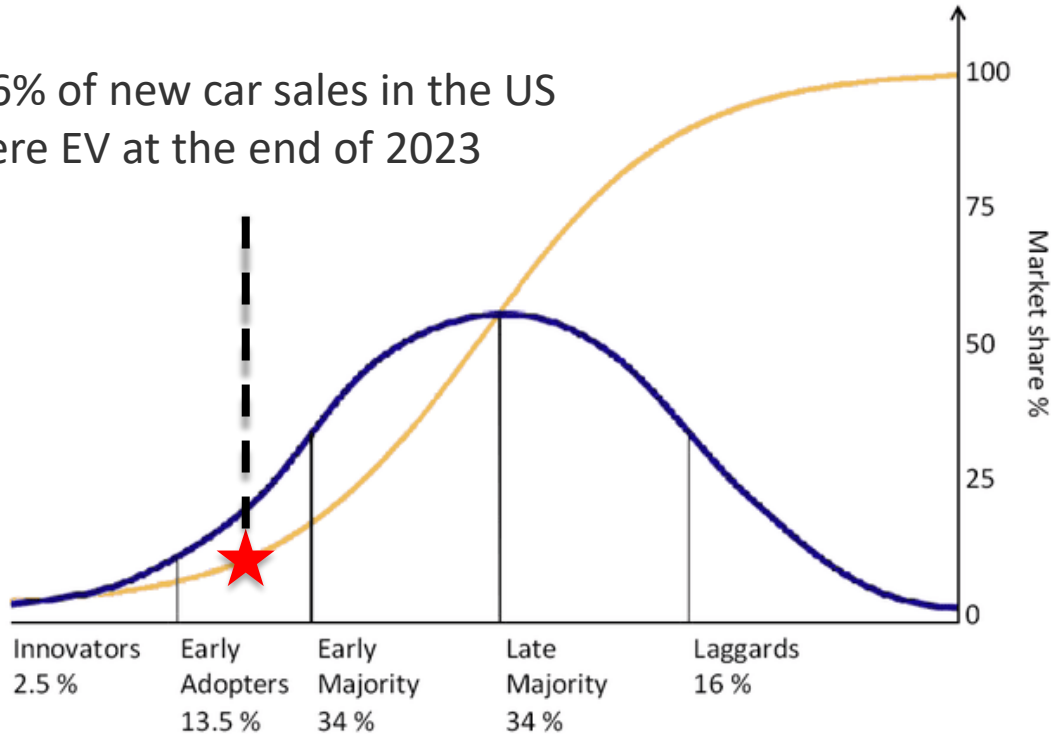


US transportation emissions



EV Adoption

7.6% of new car sales in the US were EV at the end of 2023



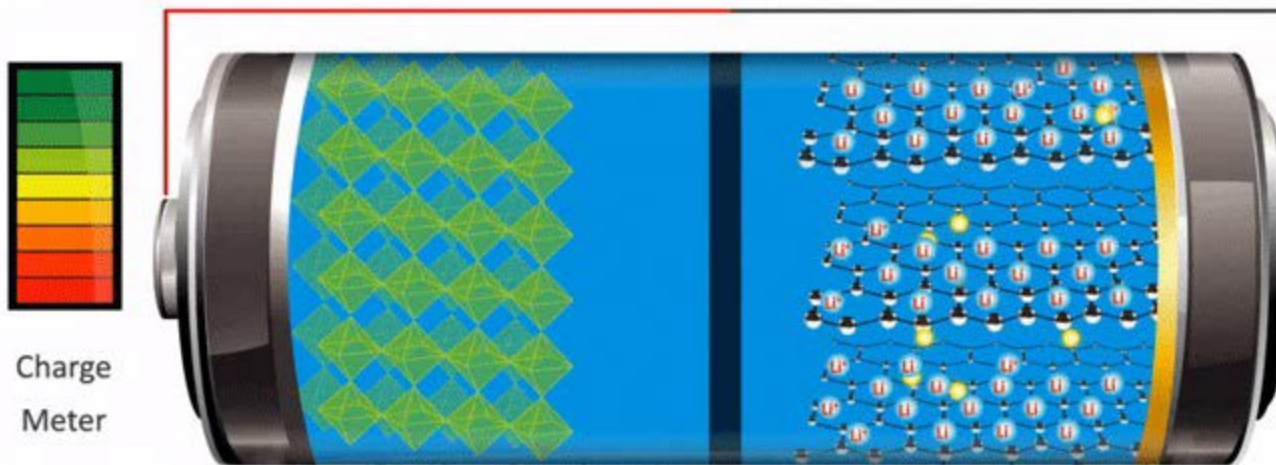
Major headwinds for EVs:

- (1) Range anxiety
- (2) Too expensive
- (3) Developing charging infrastructure

Battery technology is the bottleneck

How Lithium-ion Batteries Work

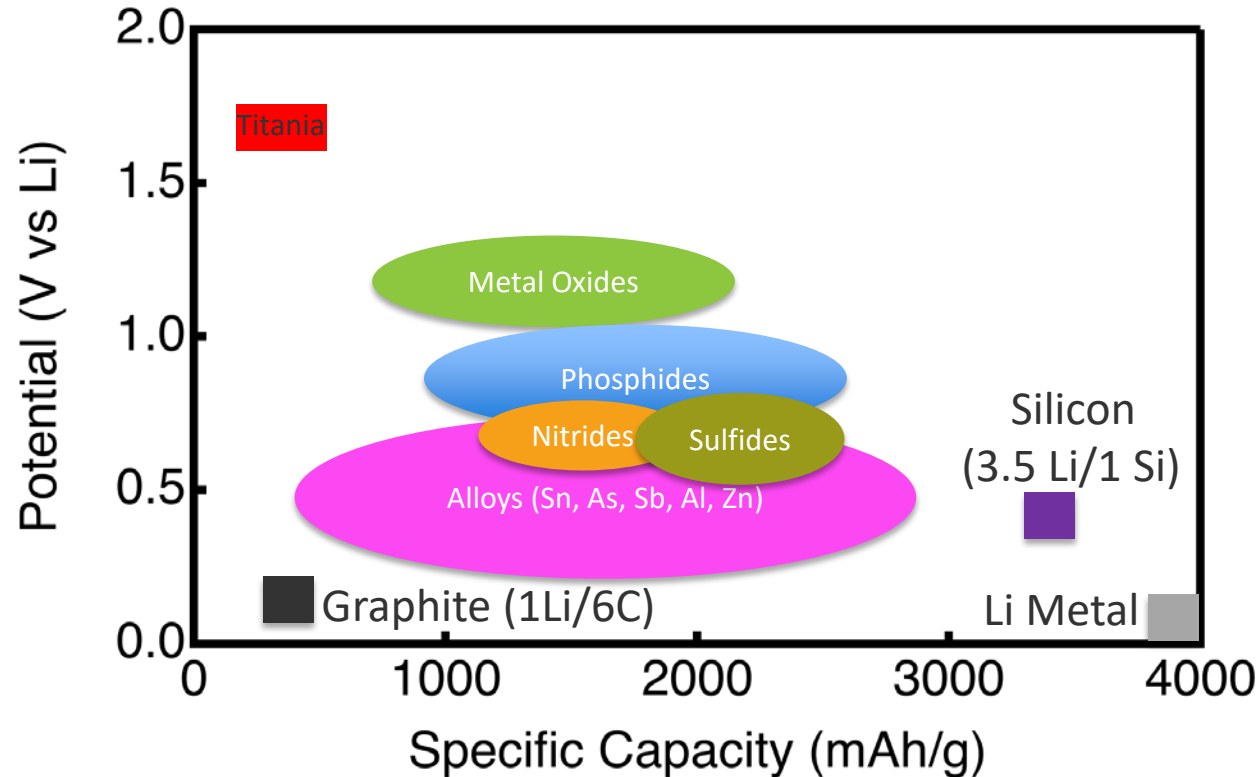
Discharge



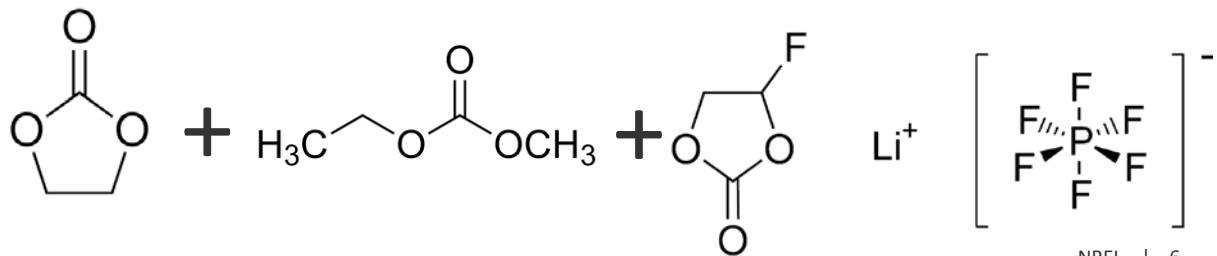
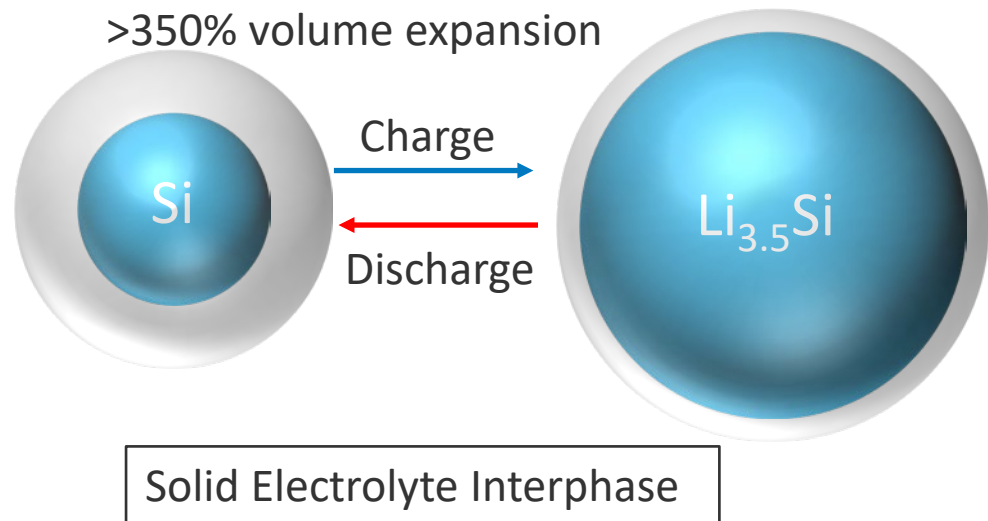
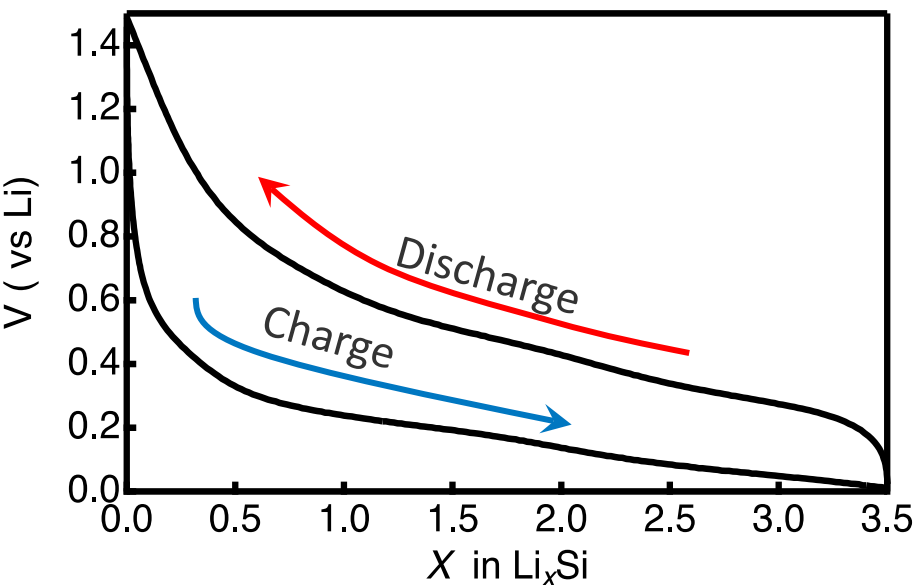
U.S. DEPARTMENT OF
ENERGY

Office of ENERGY EFFICIENCY
& RENEWABLE ENERGY

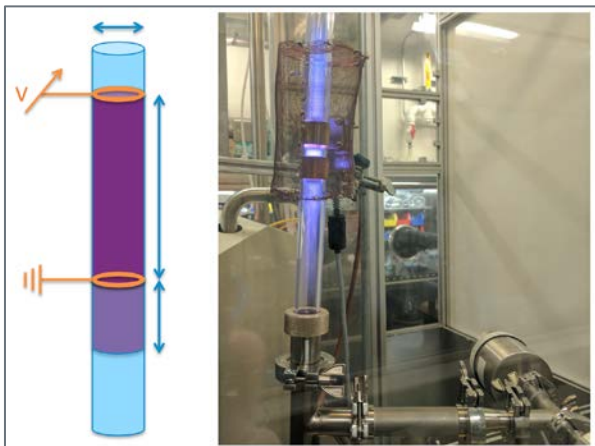
Anode Active Materials



Cycling Silicon Under Battery Operation



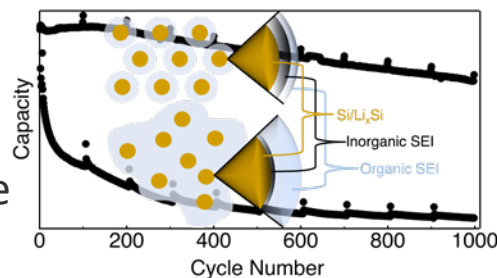
Plasma Enhanced Chemical Vapor Deposition Si NPs



-PECVD synthesis enables precise control over silicon size, surface chemistry, and composition

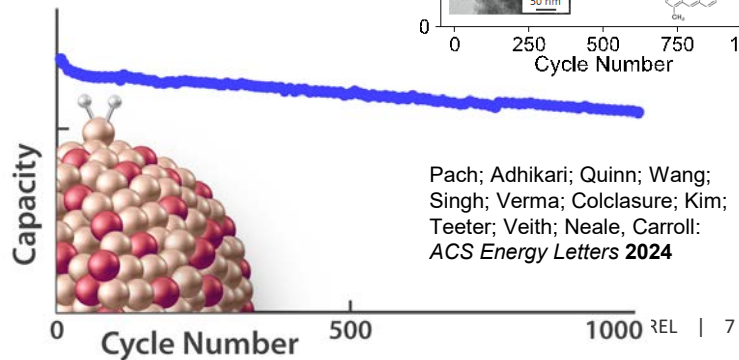
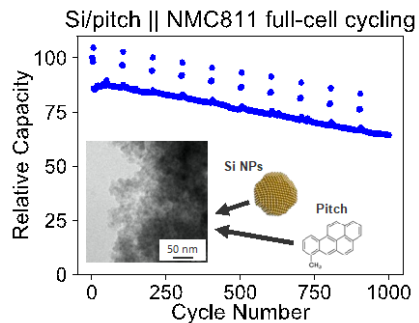
-Surfaces are terminated with SiH_x functional groups

-Radical chemistry to functionalize the silicon surface



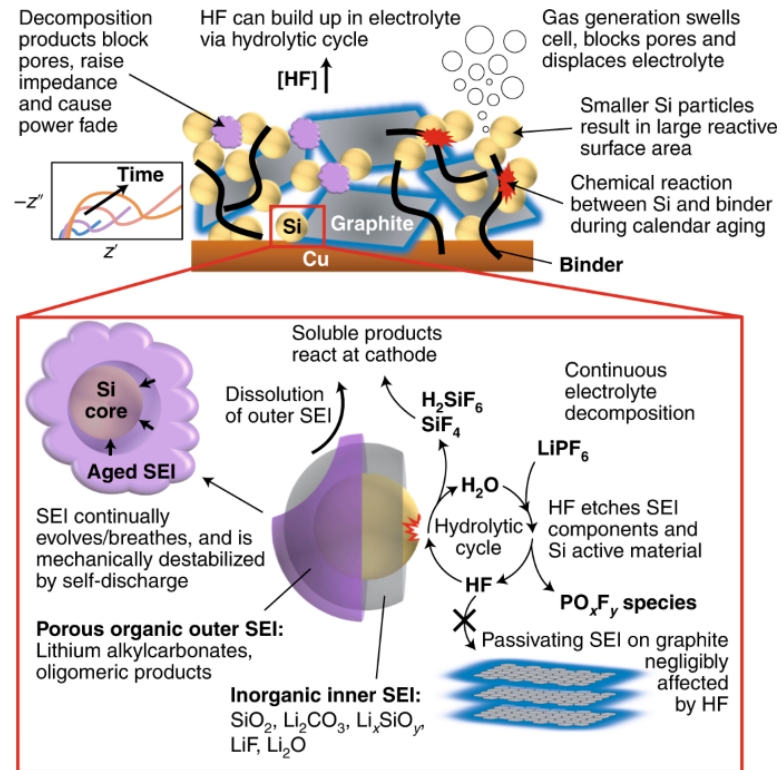
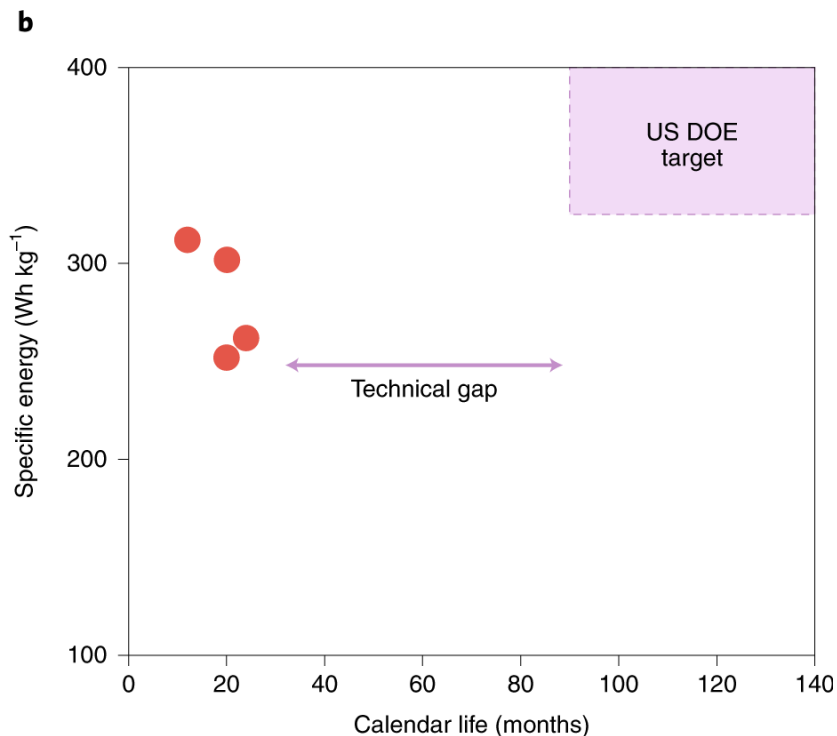
Schulze; Urias; Dutta; Huey; Coyle; Teeter; Doeren; Tremolet de Villers; Han; Neale; Carroll: *J. Mater. Chem. A* **2023**.

Schulze.; Fink; Palmer; Carroll; Dutta; Zwiefel; Engtrakul; Han; Neale, Tremolet de Villers: *Batteries and Supercaps*, **2023**.

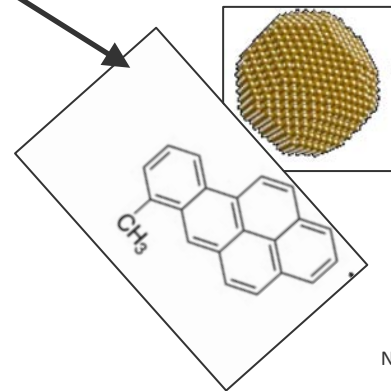
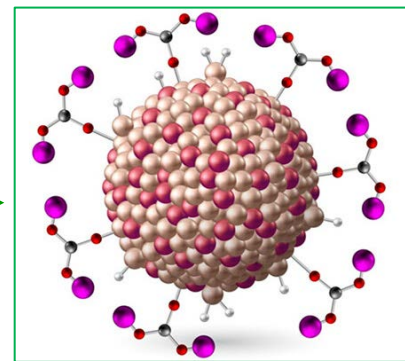
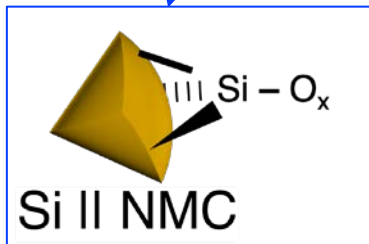
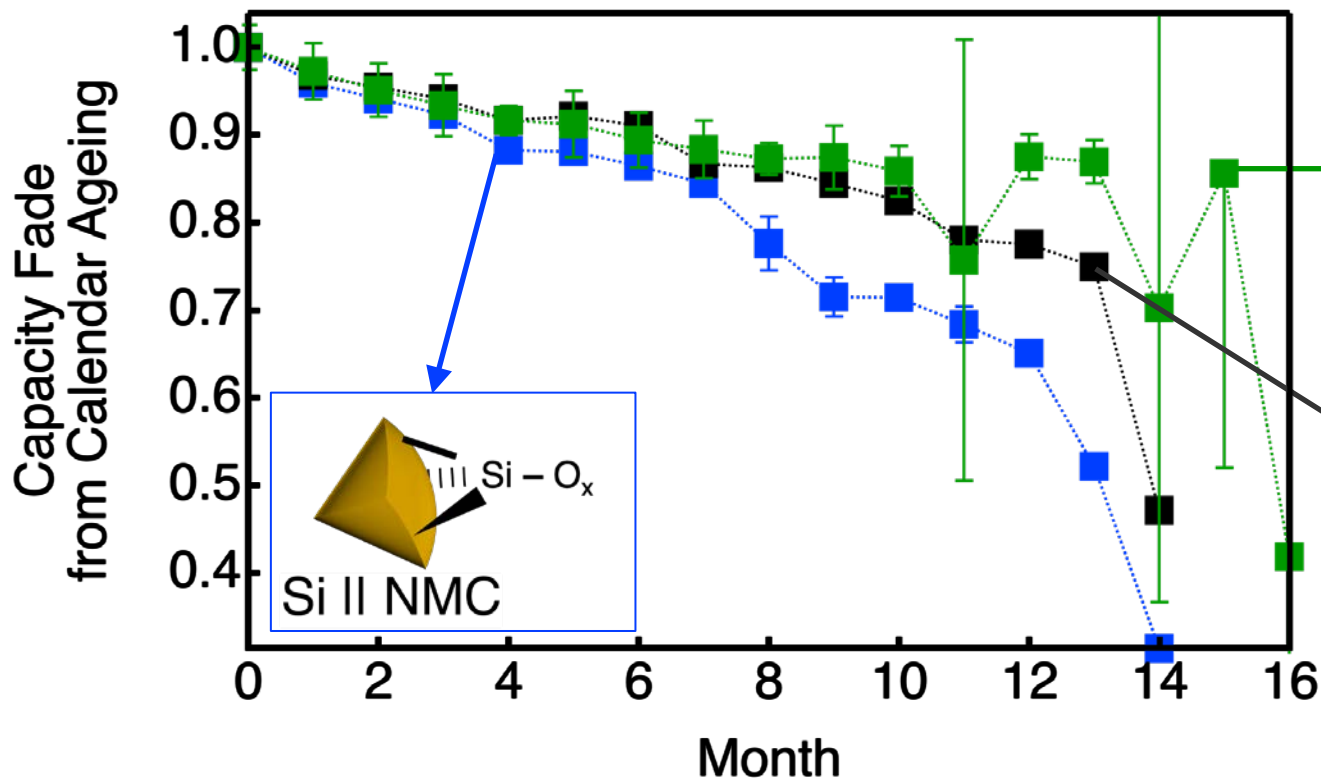


Pach; Adhikari; Quinn; Wang; Singh; Verma; Colclasure; Kim; Teeter; Veith; Neale, Carroll: *ACS Energy Letters* **2024**

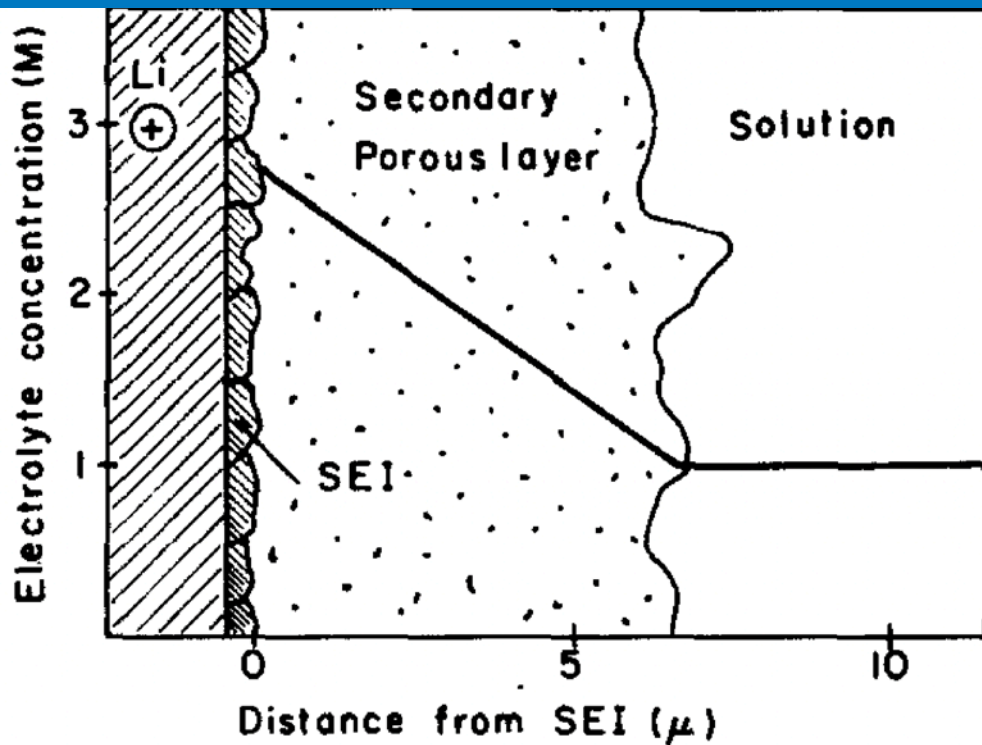
Calendar Life in Silicon Containing Anodes



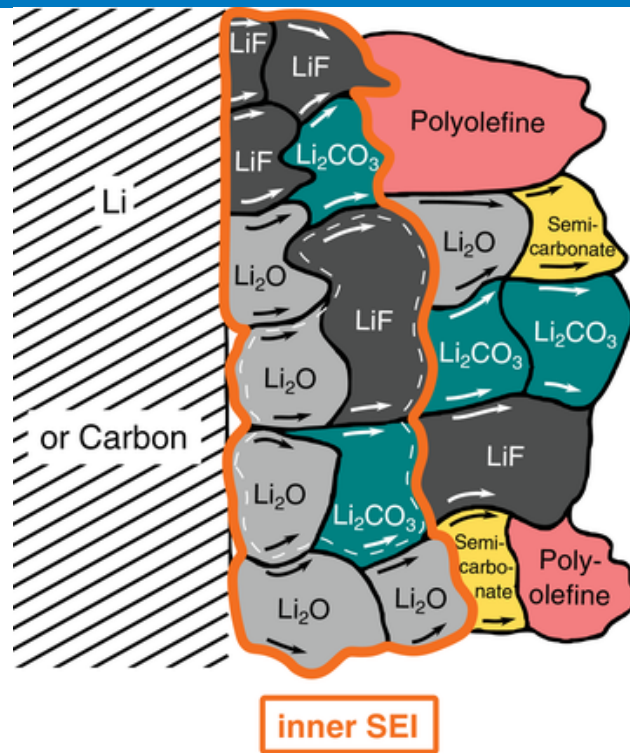
Calendar Life Issues in Silicon Containing Anodes



The SEI Model

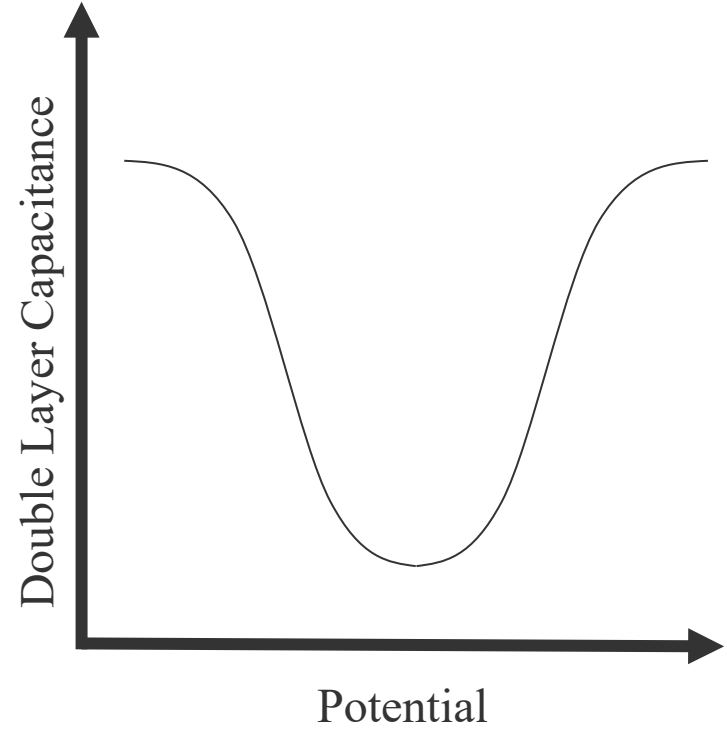
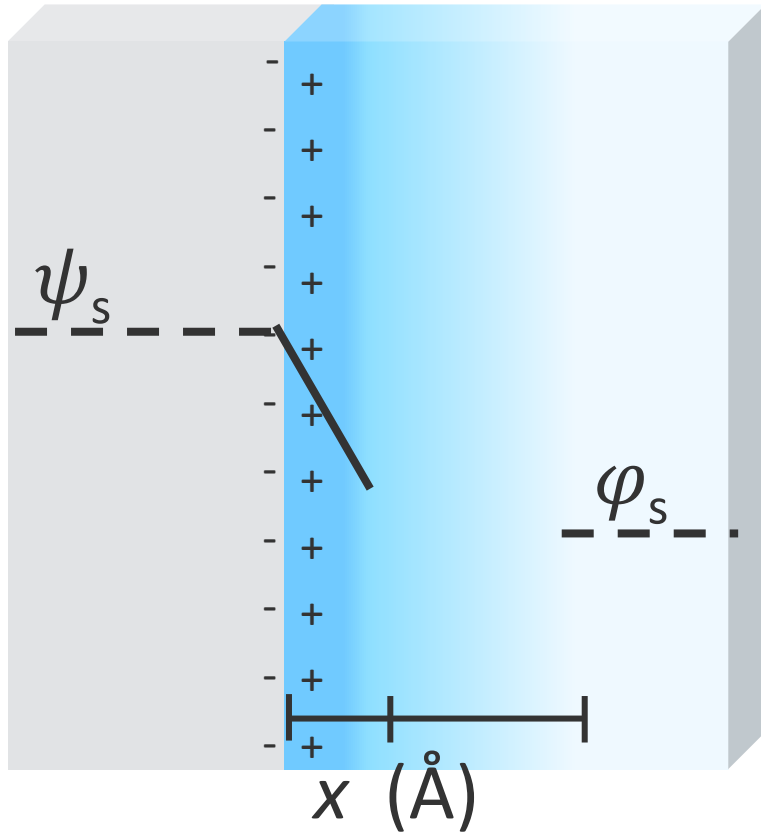


Emanuel Peled, *JPS* 1982

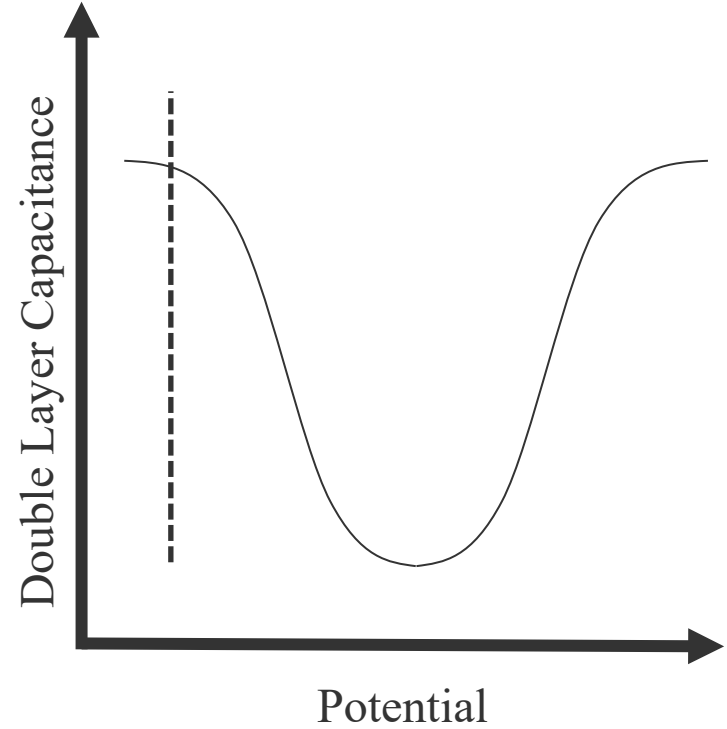
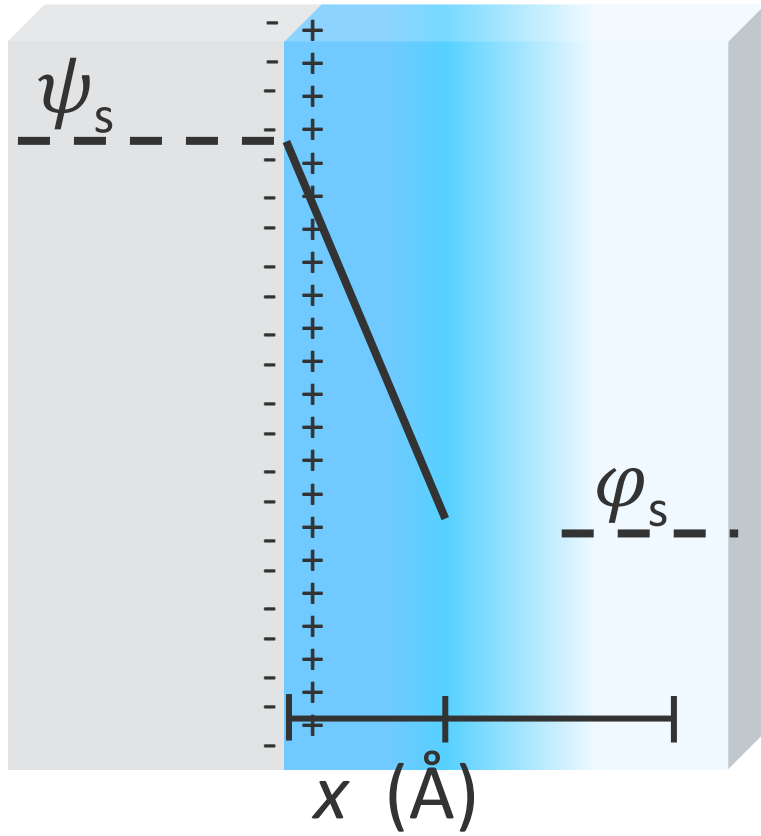


Krauss, et al., *Adv. Mater. Int.* 2022

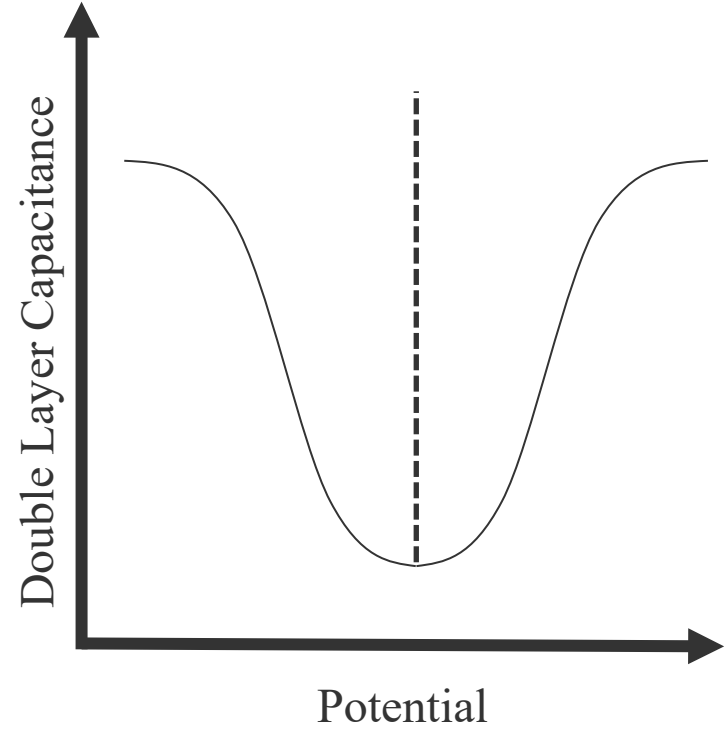
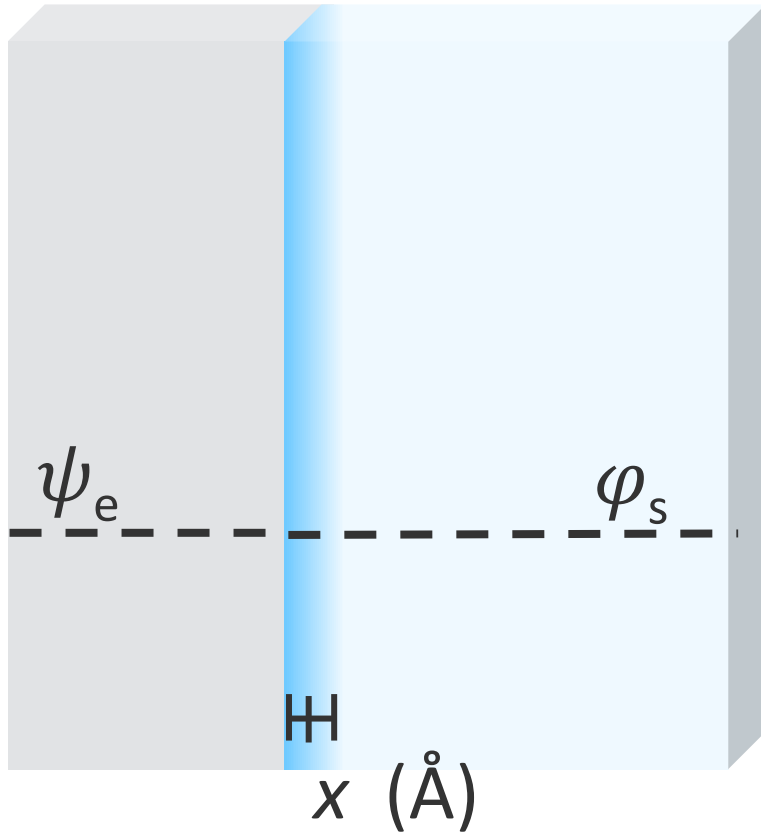
The Classical Model



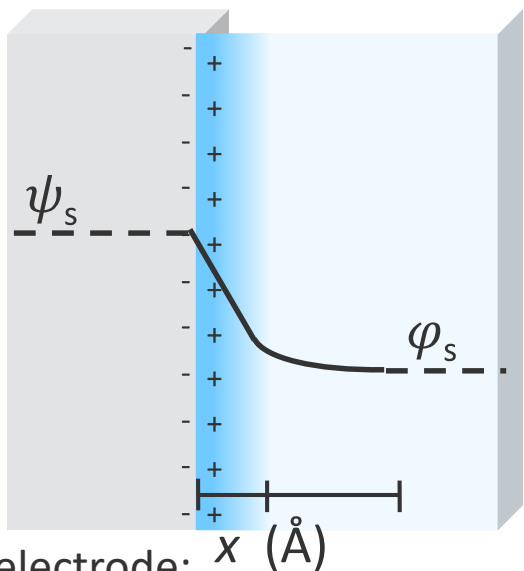
The Classical Model



The Classical Model



(Re-)Connecting Li_xSi to Model Interfaces



Ideal electrode:

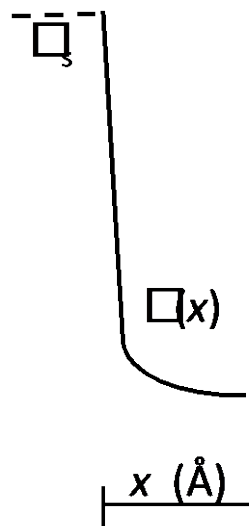
(1) Static and well-defined electrode interface

(2) No hysteresis between cycling

→ No change to electrode surface

→ No change to electrolyte

Li_xSi SEI Electrolyte



Silicon electrode:

(1) Dynamic Si electrode interface with lithiation

(2) Poorly-defined interface (Li_xSi ?, SEI?, $\text{Li}_x\text{Si}_y\text{O}_z$)

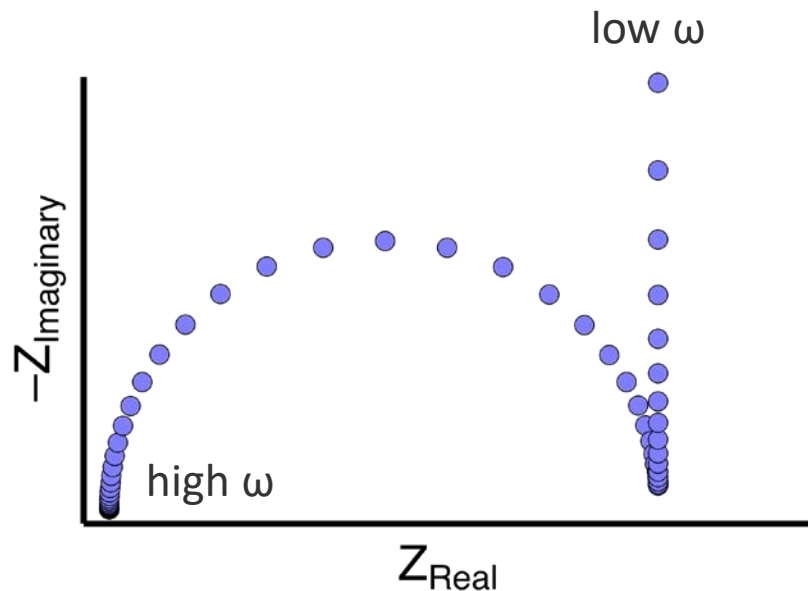
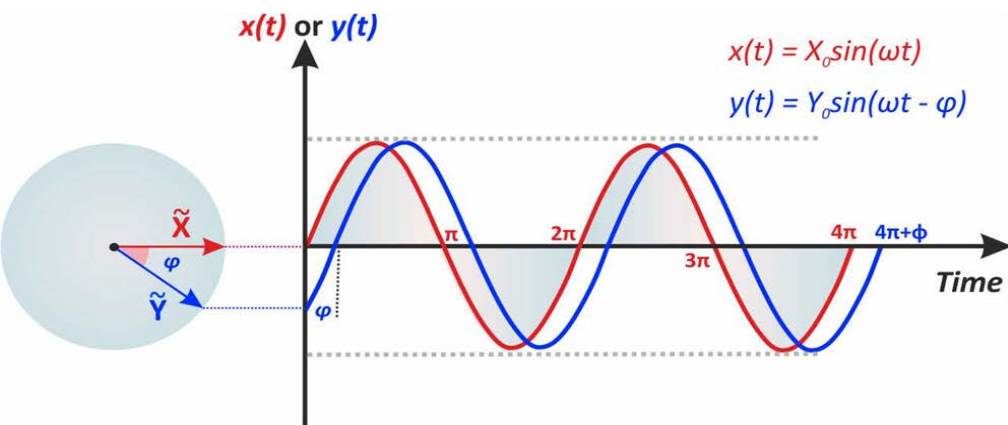
(3) Unstable

(intrinsically reactive)

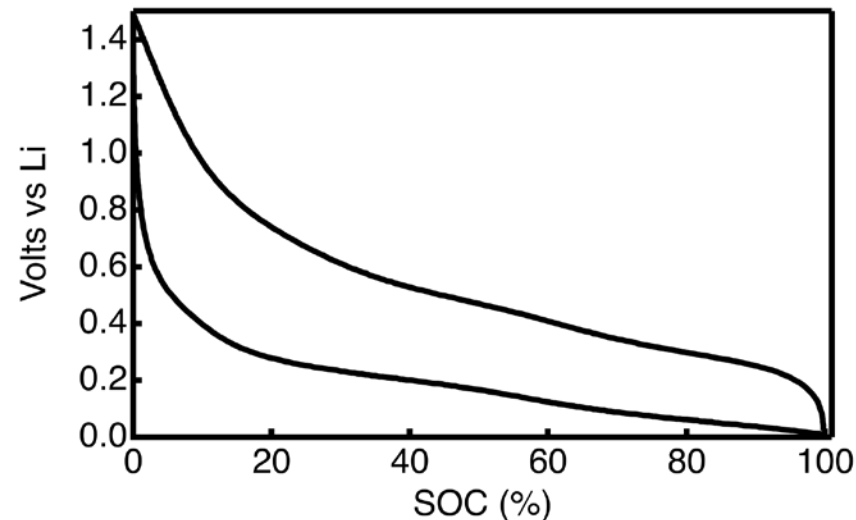
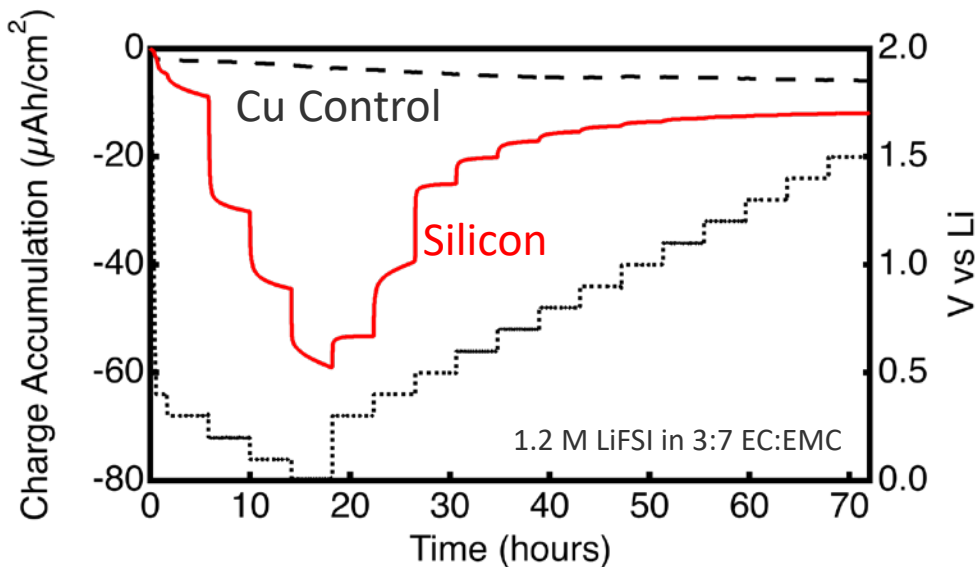
(4) Hysteresis between cycles

(5) Extremely heterogeneous

Electrochemical Impedance Spectroscopy

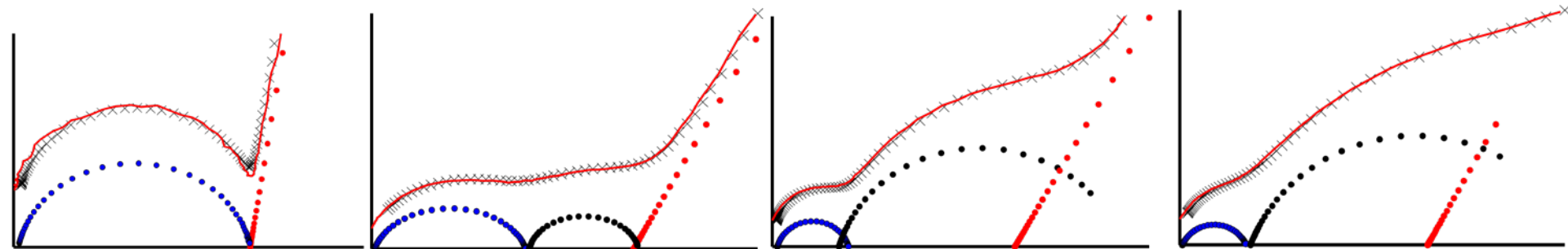


SOC – Dependent EIS



Silicon lithiation states held at a constant potential
Potentiostatic EIS performed at end of V-hold every 4 hours.
Delithiation sweep investigated specifically

Impedance Evolution on Delithiation



$E_W \leq 0.7$ vs Li

$E_W = 0.8$ V vs Li

$E_W = 0.9$ V vs Li

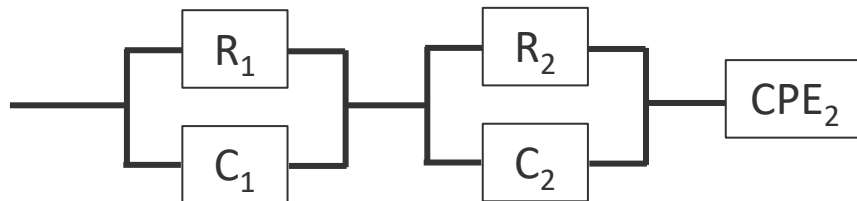
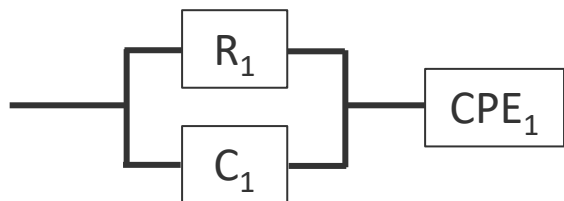
$E_W \geq 1$ V vs Li

SOC > 20%

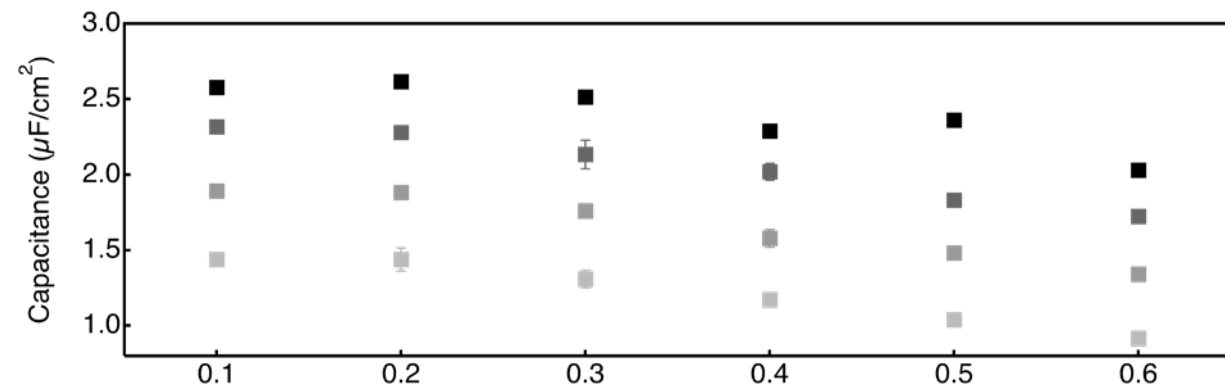
SOC ~ 20%

SOC ~ 10%

SOC < 5%



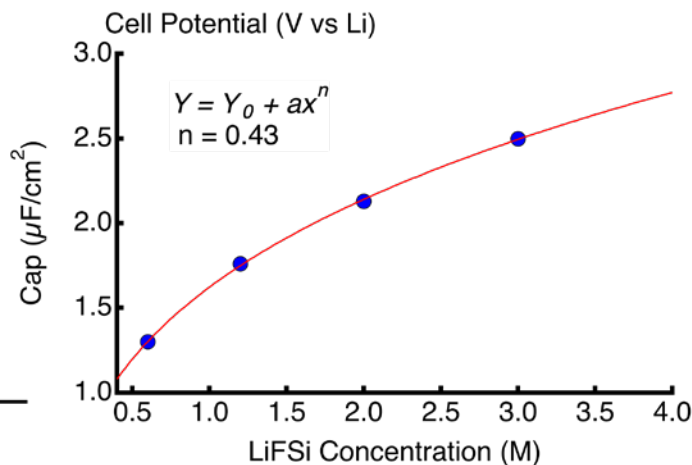
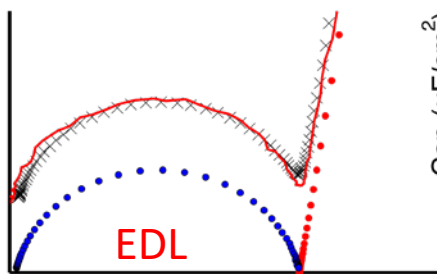
Capacitive Feature Assignment



Double layer capacitance:

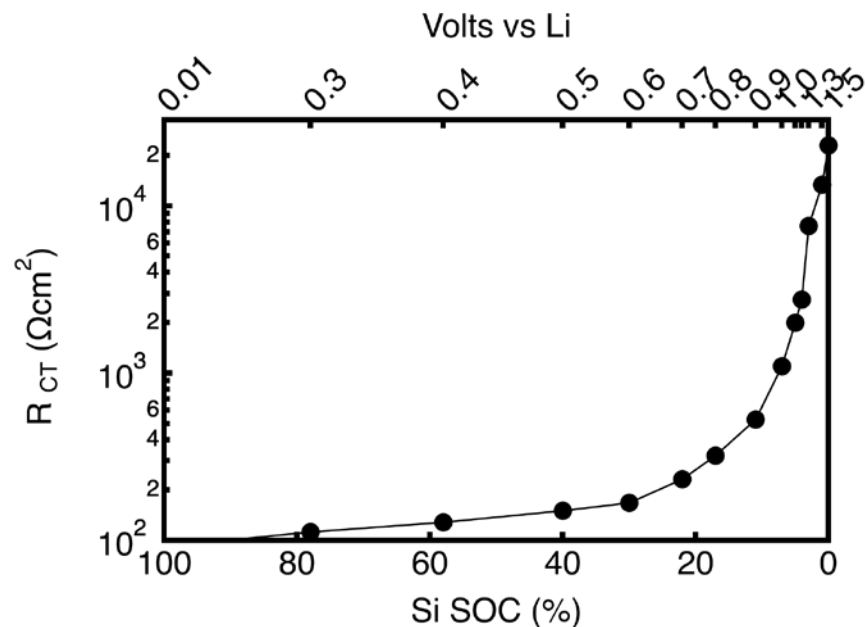
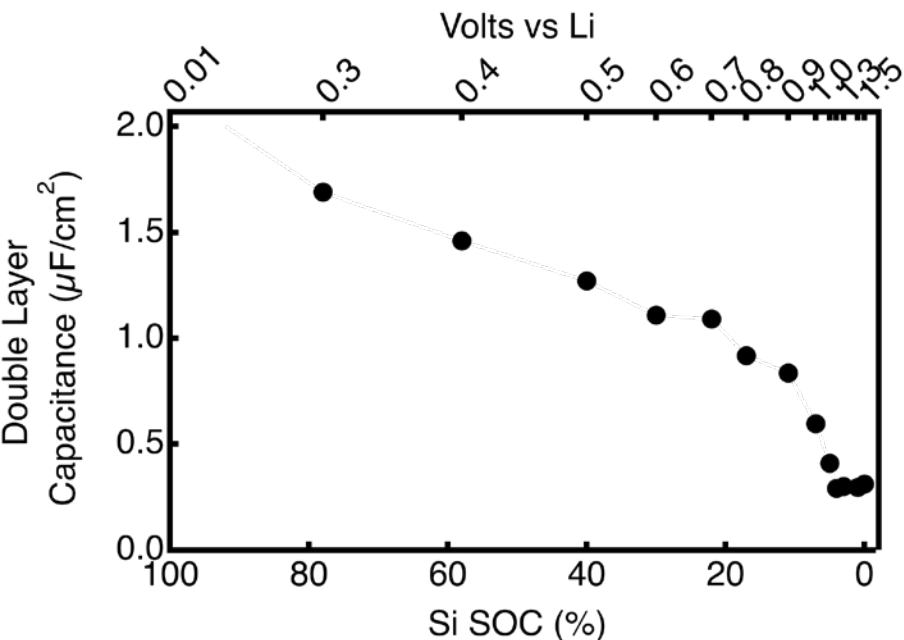
Differential capacitance is not zero

Capacitance is proportional to sq. root of ion activity ($\sim \text{concentration}^{1/2}$)



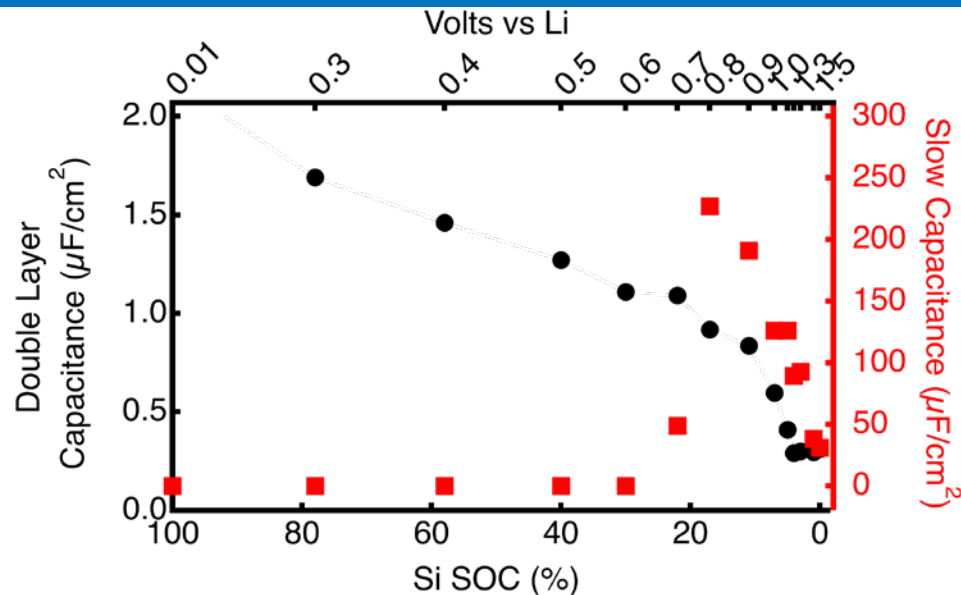
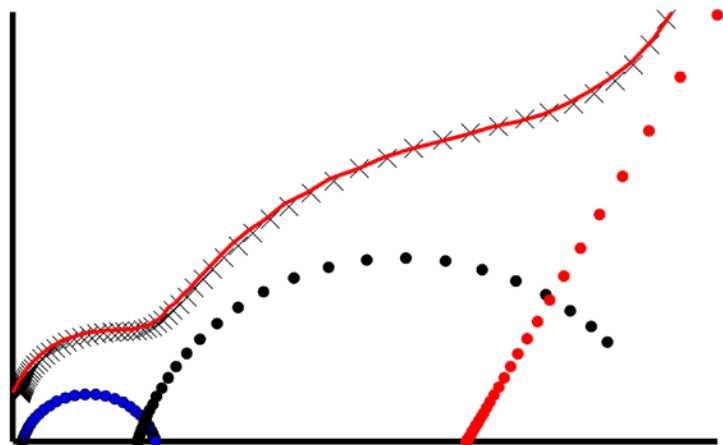
First semi circle is capacitance of EDL.

EDLC and Charge Transfer Resistance



Inverse relationship between R_{CT} and C_{DL}
Indicates change in limiting step in charge transfer

Slow Process Capacitance

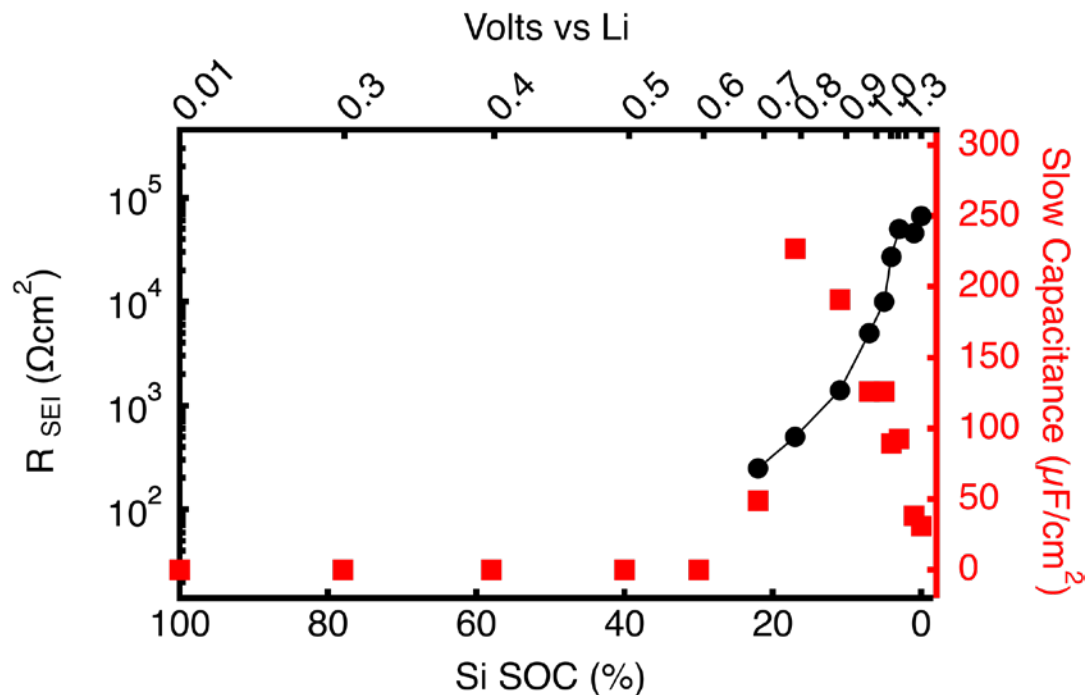


Capacitance of slow process has sharp increase at EDL drop

Slow capacitance is > 100 greater than EDL capacitance

Slow semicircle is likely SEI capacitance and resistance

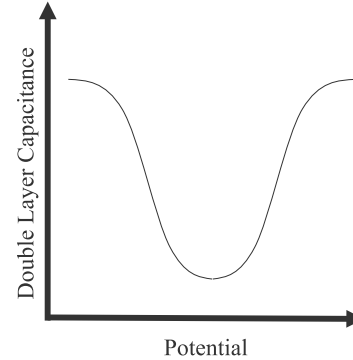
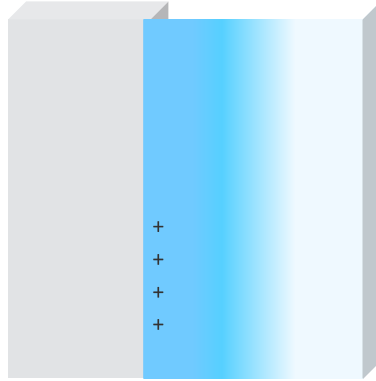
Resistance of the SEI



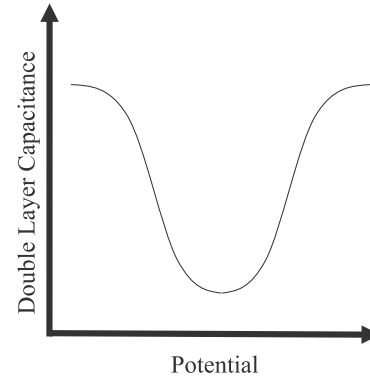
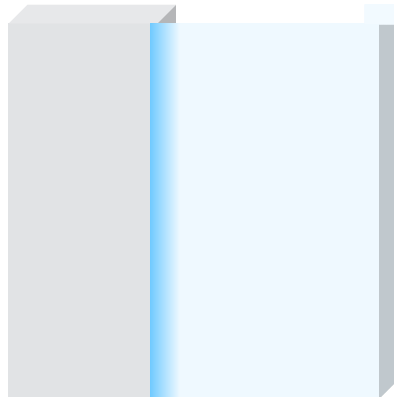
Resistance values keep climbing despite decreasing differential capacitance
Suggests SEI continues to grow

SOC-dependent EDLC

$\sim 90\% > \text{SOC} > \sim 20\%$



$\text{SOC} < \sim 20\%$



Calendar Aging in Symmetric Cells

Si@PEO || Si@PEO

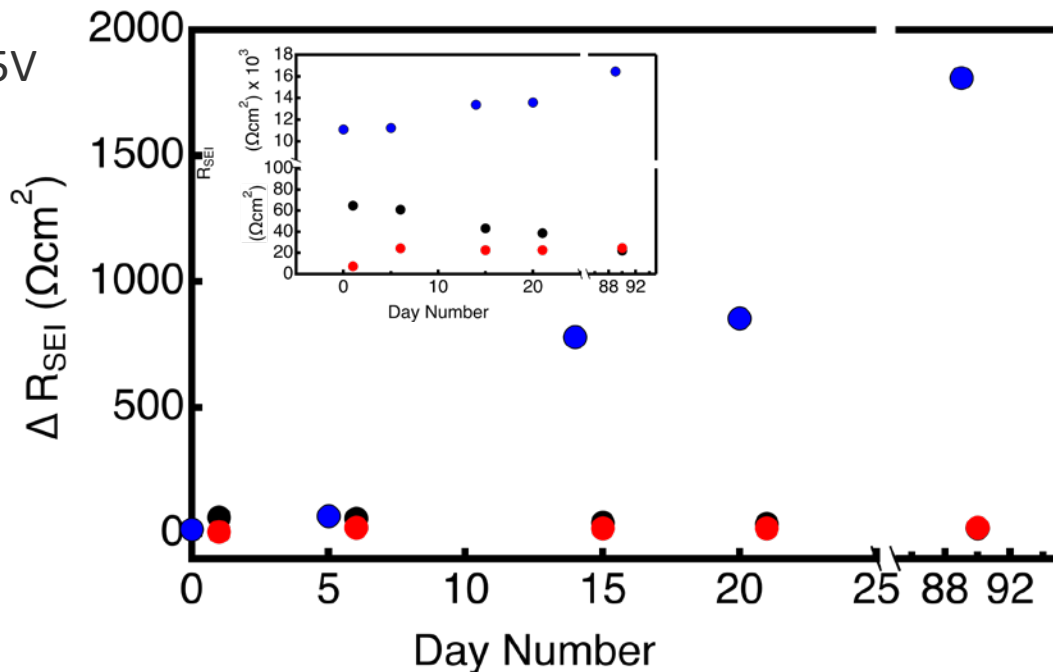
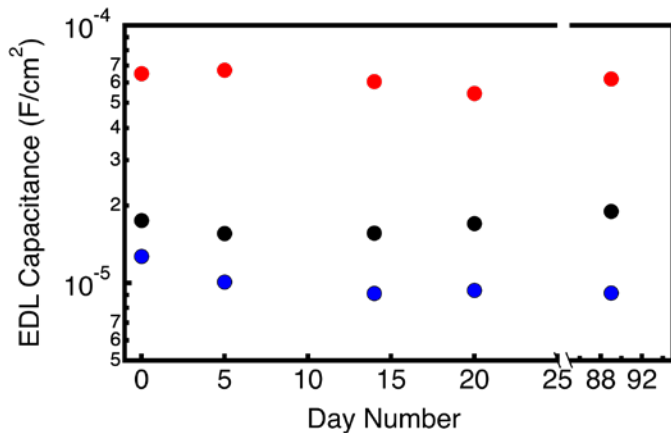
Gen2 electrolyte

Formed at C/10 between 0.01 and 1.5V

Delithiated to: 0.01 V, 0.4 V, 1 V

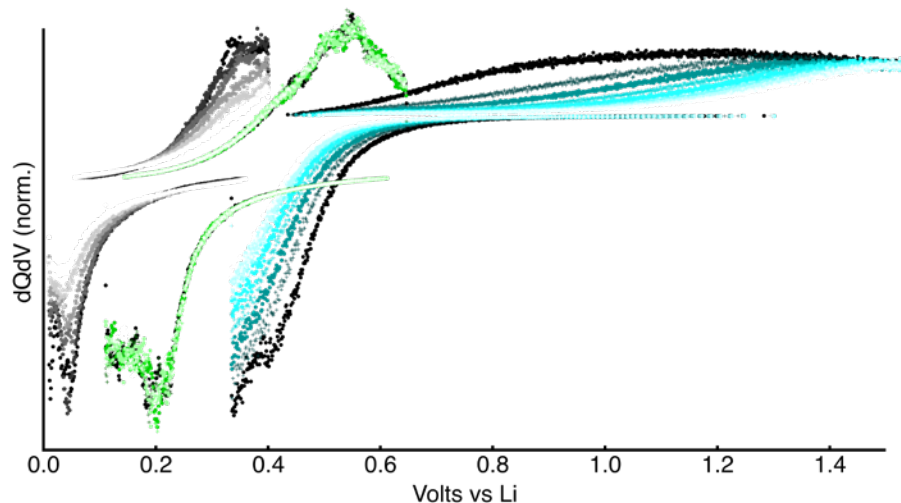
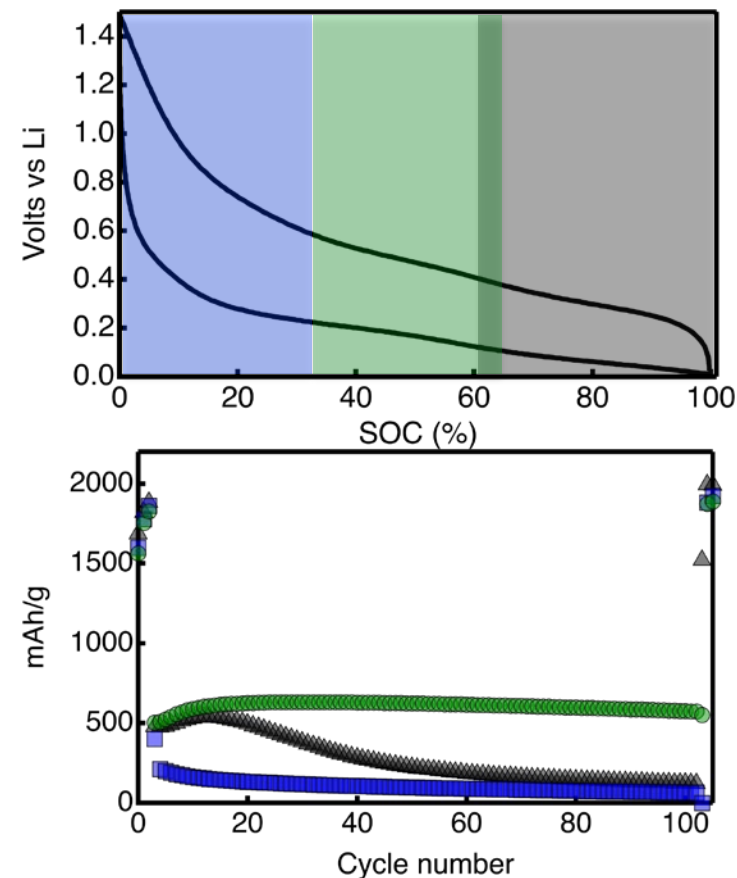
48 hr V-hold @ desired potential

Aged at 45 C at 0V vs Li_xSi



Increasing impedance at low SOC is consistent with PZC-like regime

Cycle Lifetime (Composite Half cells)

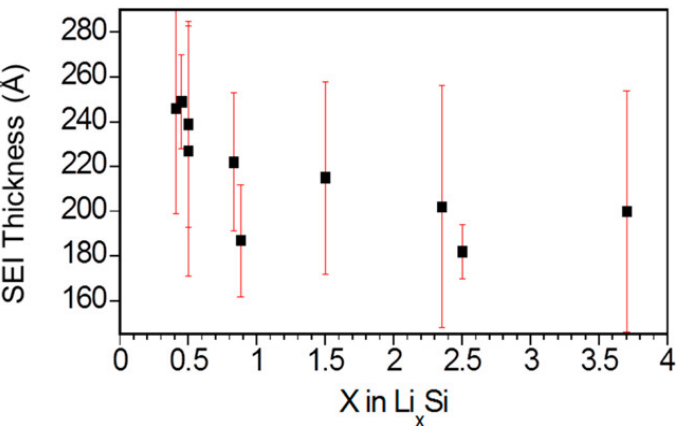


Large impedance gain in High SOC and Low SOC cycle domains

No apparent loss of active material

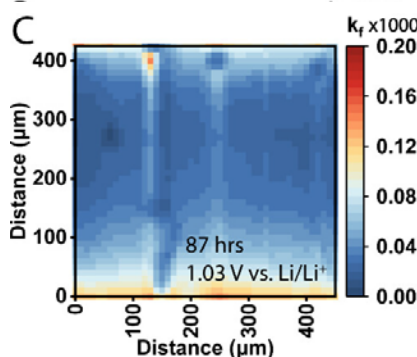
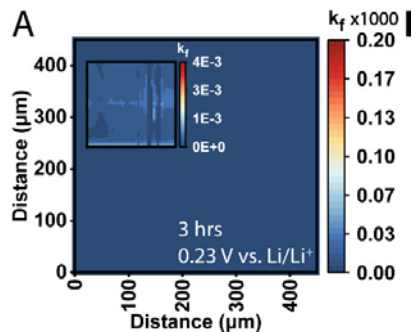
SOC-Dependent SEI Electrostatic Picture

Neutron Reflectometry Measurements of SEI thickness

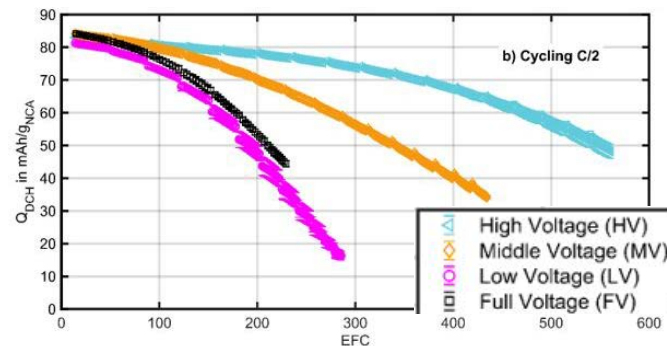


Veith, et al; *JPCC* **2019**

Scanning Electrochemical Microscopy Measurements on SEI Passivation



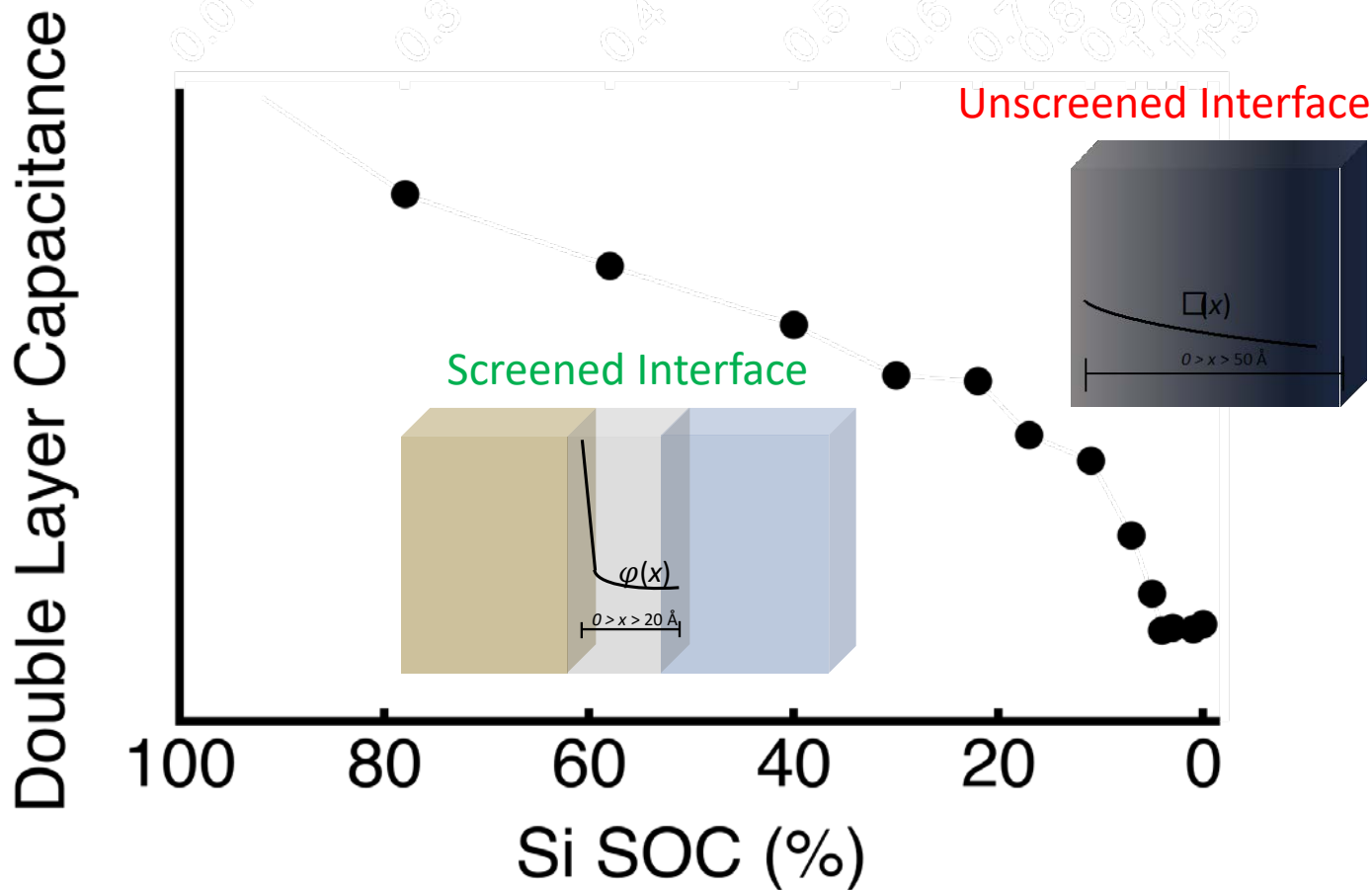
Battery Cycling



Sven Friedrich, 245th ECS
A02-0240 **2024**

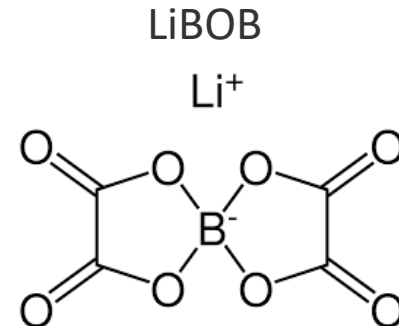
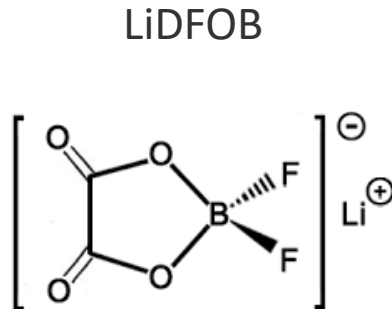
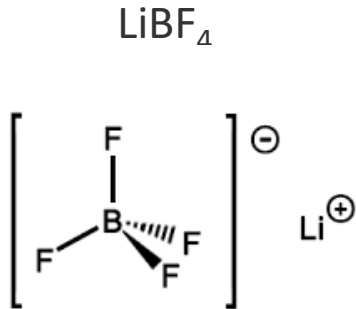
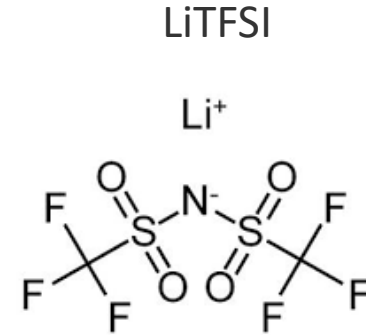
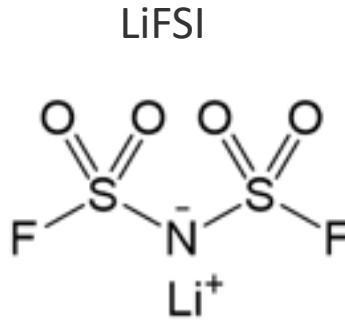
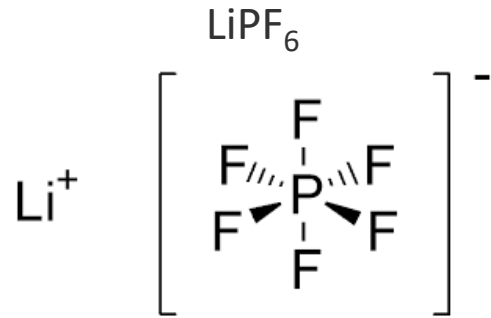
McBrayer, et al; *J. Appl. Mater. Interfaces* **2024**

SOC-Dependent SEI Electrostatic Picture

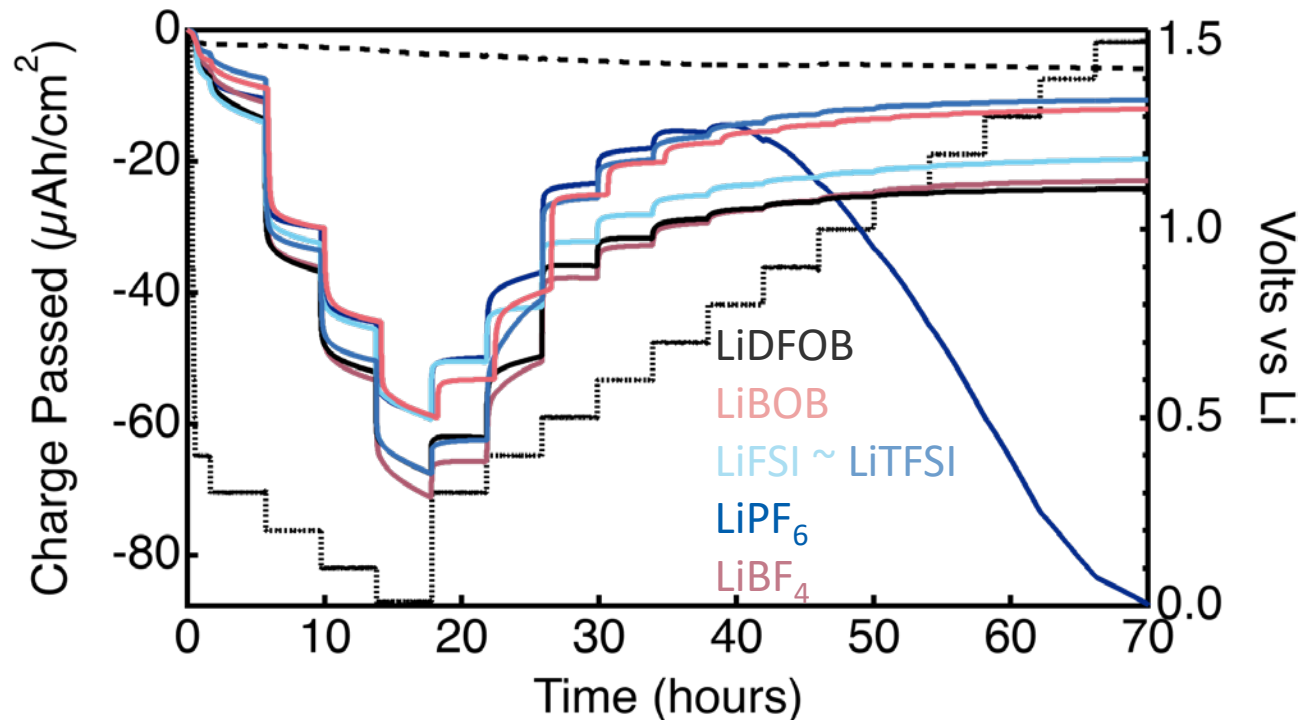


Anion Variation

1.2M LiX dissolved in 3:7 EC:EMC



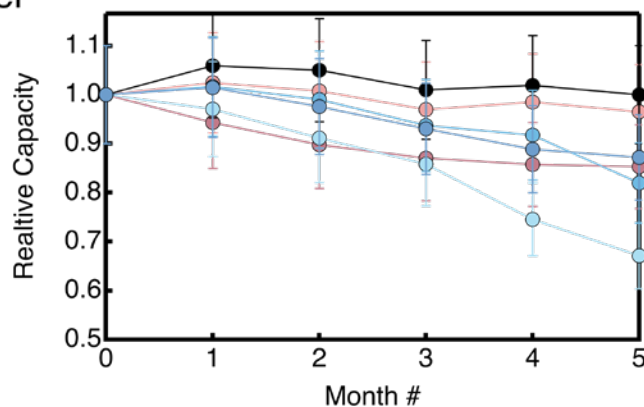
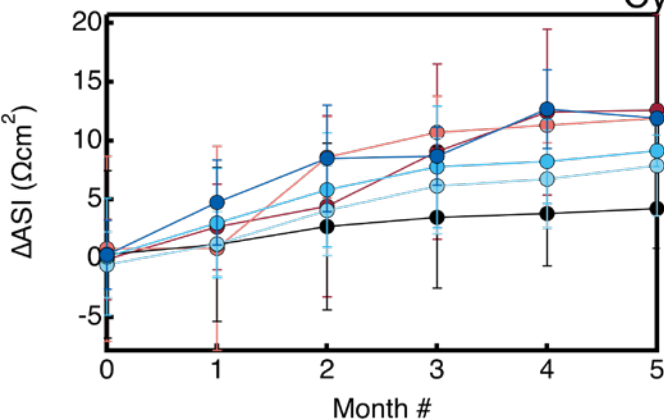
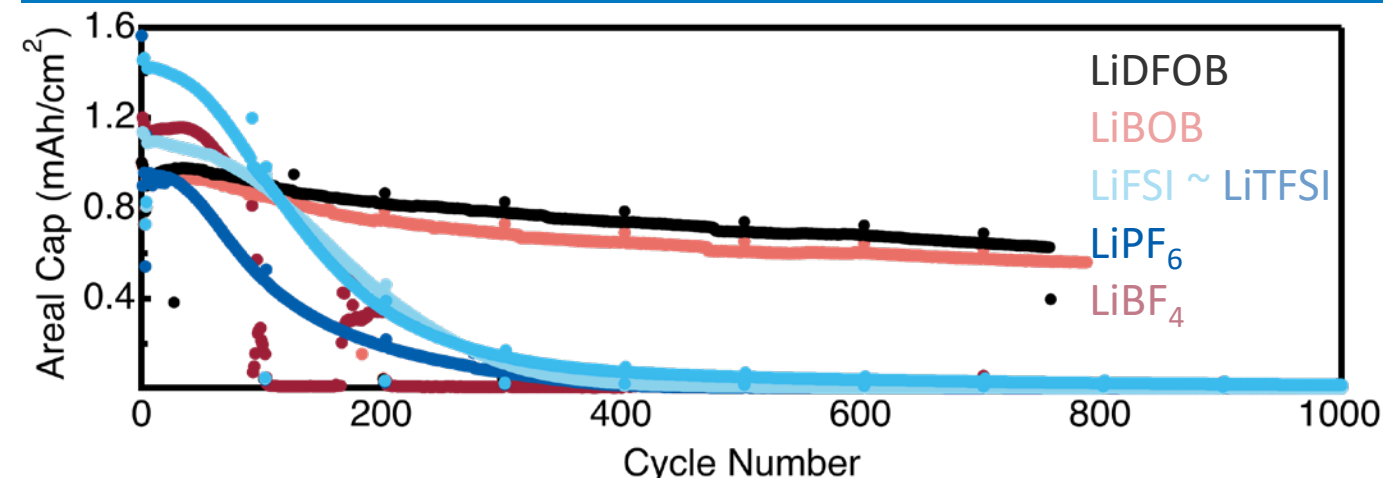
Charge Accumulation



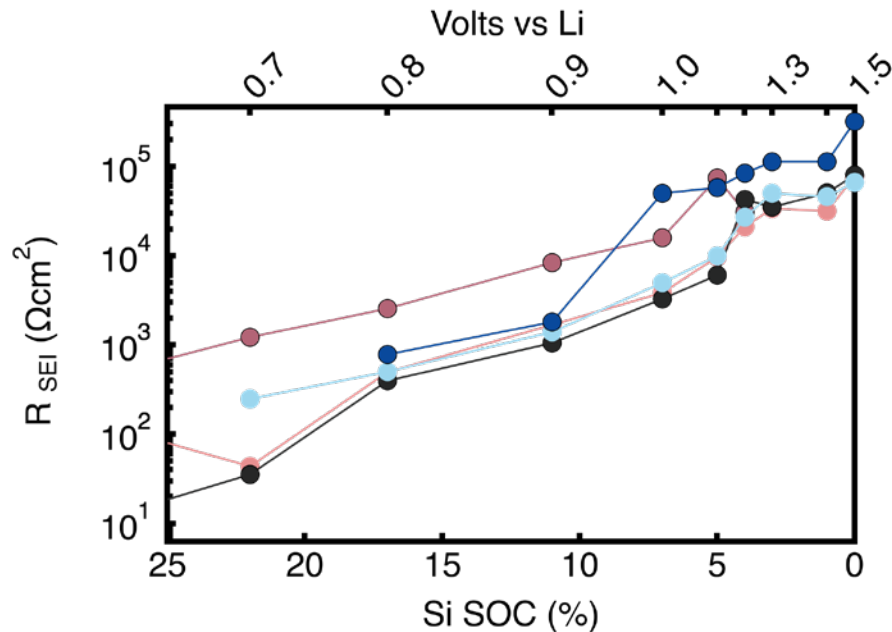
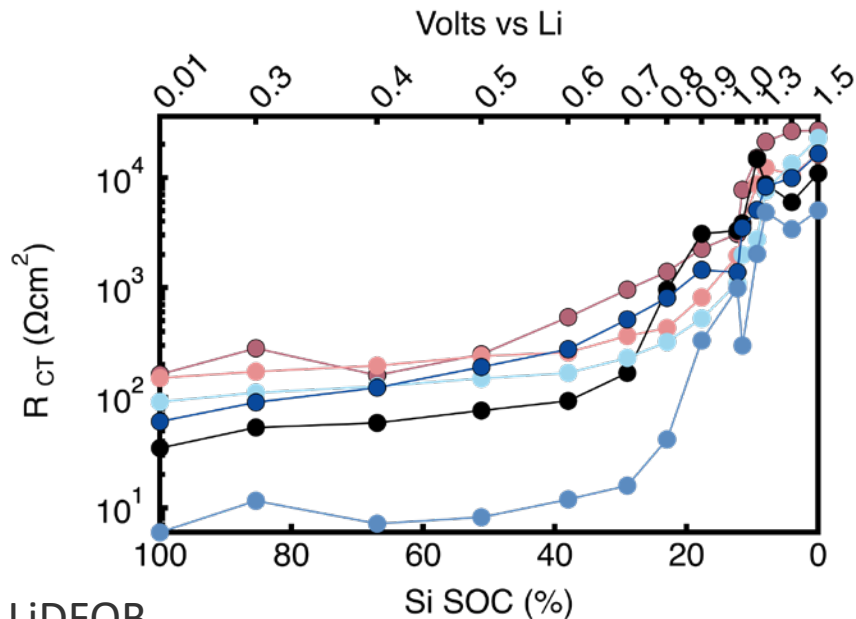
LiPF_6 coupled with EC shows continuous current at low SOC's
All other electrolytes are stable at low SOC's

Cycle/Calendar Lifetime vs LFP (n-limited)

Si@PEO || LFP
1.2 M salt
3:7 EC:EMC
coin cell
non-prelithiated



Resistances



LiDFOB

LiBOB

LiFSI ~ LiTFSI

LiPF₆

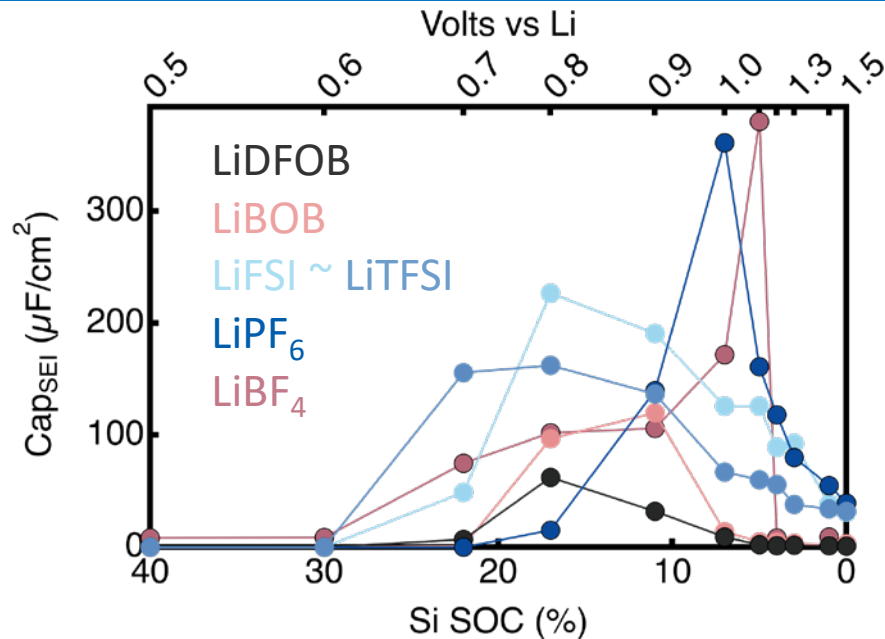
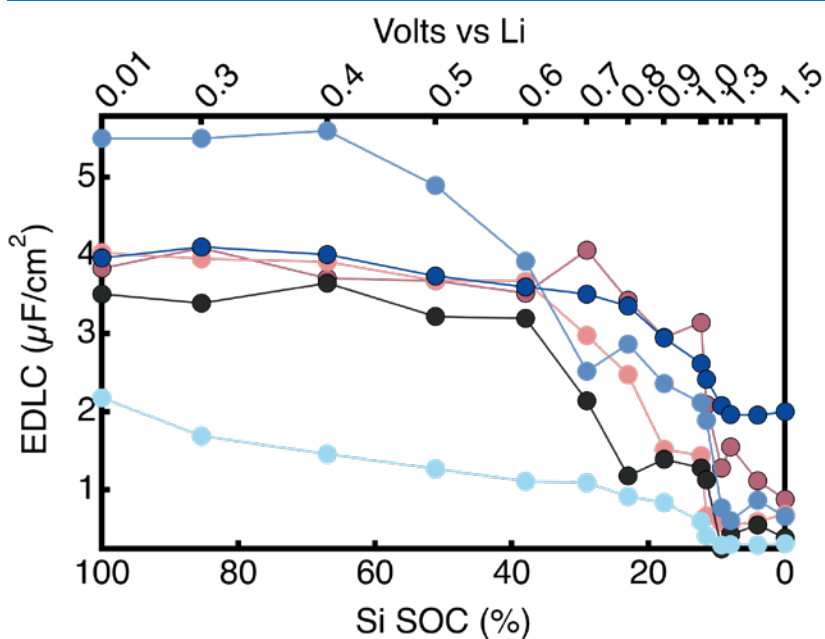
LiBF₄

R_{CT} and R_{SEI} follow the same trajectory for all salts

R_{CT} values vary significantly

R_{SEI} are generally the same

Capacitance and Passivation



Onset for SEI capacitance gain tracks with the drop in EDLC (except LiBF₄)

→ Expected as PZC is set by electrode *and* electrolyte

Capacitance in SEI roughly follows performance → lower SEI capacitance is signature of more passivated surface

Conclusions

1. Lithium silicide interface and SEI can be understood through the classical model of electrochemical interfaces
 - EIS enables measurements of relevant properties at a very non-ideal electrochemical interface
2. EDLC measurements reveal that Li_xSi have very dynamic surface electrostatics:
 - At SOC < ~20% the electrode surface resembles a structure analogous to the PZC
 - The PZC regime is least passivated--> fastest impedance gain in calendar life cells
 - The PZC regime may be appropriate for accelerating calendar life measurements and assessing the passivation of the SEI with different electrolytes
3. Measurements of different anions:
 1. The onset of PZC regime depends on anions
 2. Anions play a large role in passivation
 3. The degree of passivation (C_{SEI}) strongly correlates with cycle and calendar life

Conclusions

1. Cycle life in majority Si anodes is less relevant than calendar life
2. Lithium silicide interface and SEI can be understood through the classical model of electrochemical interfaces
 - EIS enables measurements of fundamental electrochemical interface properties of a highly non-ideal electrochemical interface
3. The interface structure and mass transport properties are strongly dependent on the electrode state of charge

If your battery has silicon in it, don't store it fully charged or discharged

The Silicon Consortium Project

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		Josey McBrayer	Lydia Meyer	Andrew Colclasure	Joseph Quinn

Questions?

www.nrel.gov

Mike Carroll

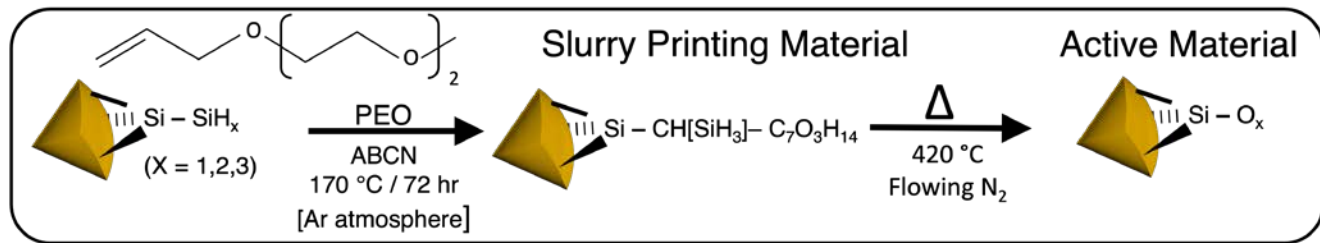
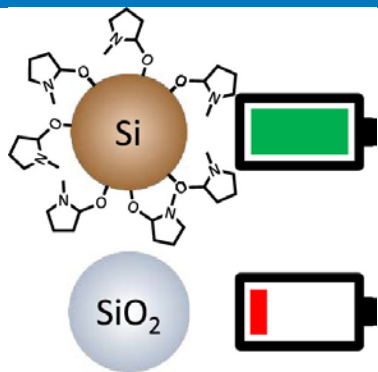
mike.carroll@nrel.gov

NREL/PR-5900-90535

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Controlling the Surface Chemistry of Si Electrodes

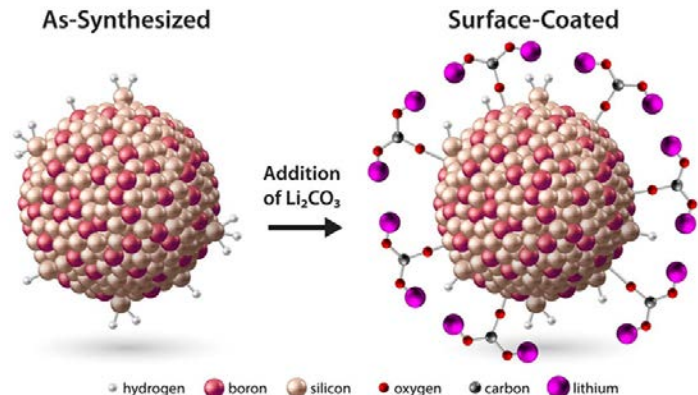


Schulze, ... Carroll; *J. Mater. Chem. A* **2023**

Carroll, ... Neale; *ACS Appl. Energy Mater.* **2020**



Schulze, et al.; *Batteries and Supercaps* **2023**



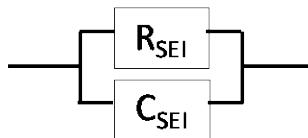
Pach, ... Carroll; *ACS Energy Letters* **2024**

EIS of a Redox Active Thin Film (SEI) at a Reflective Boundary.

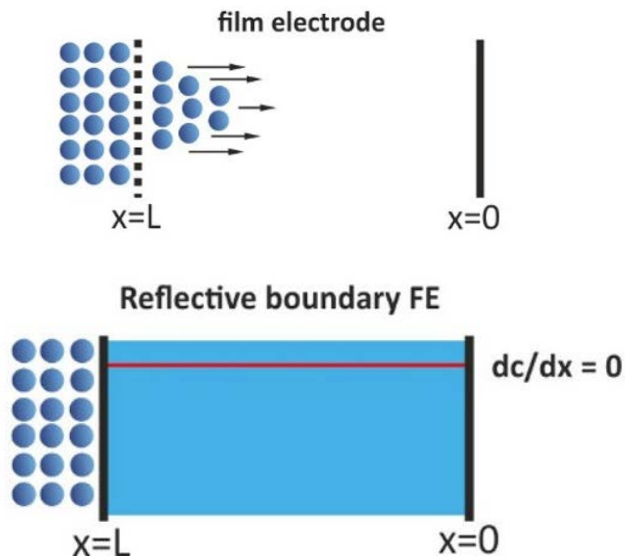
(1) Faradaic
Redox Chemistry



Randles' Circuit

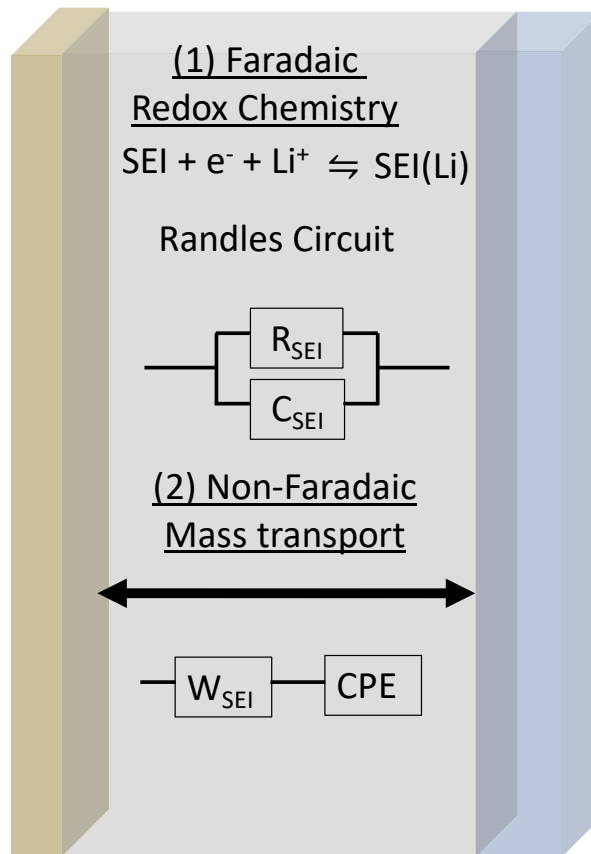


Mass Transport Through the SEI

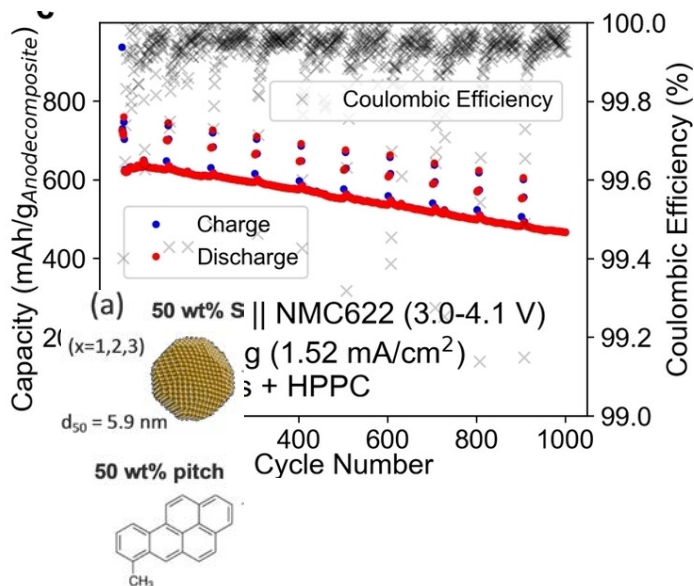


$$Z_t(\omega) = \frac{\coth(B\sqrt{j\omega})}{Y_0\sqrt{j\omega}}$$

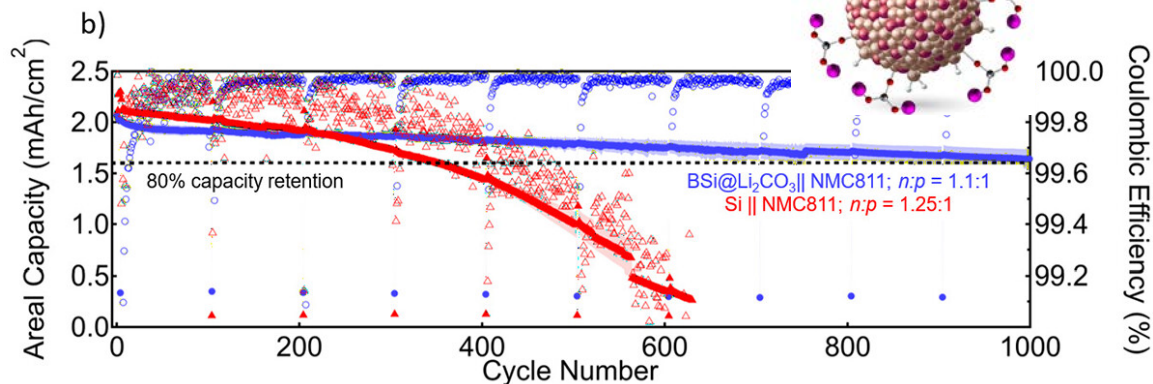
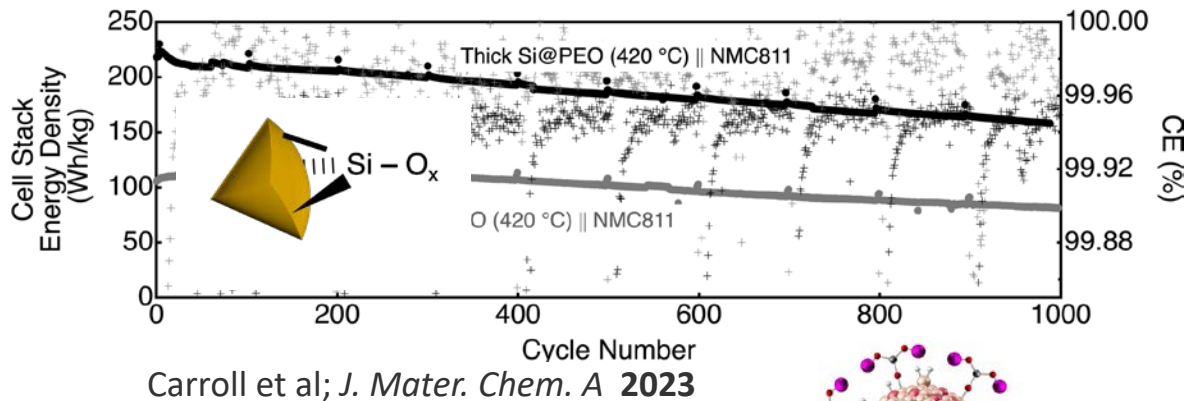
$$\beta = \frac{\text{diffusion layer thickness}}{\text{diffusion coefficient}^{1/2}}$$



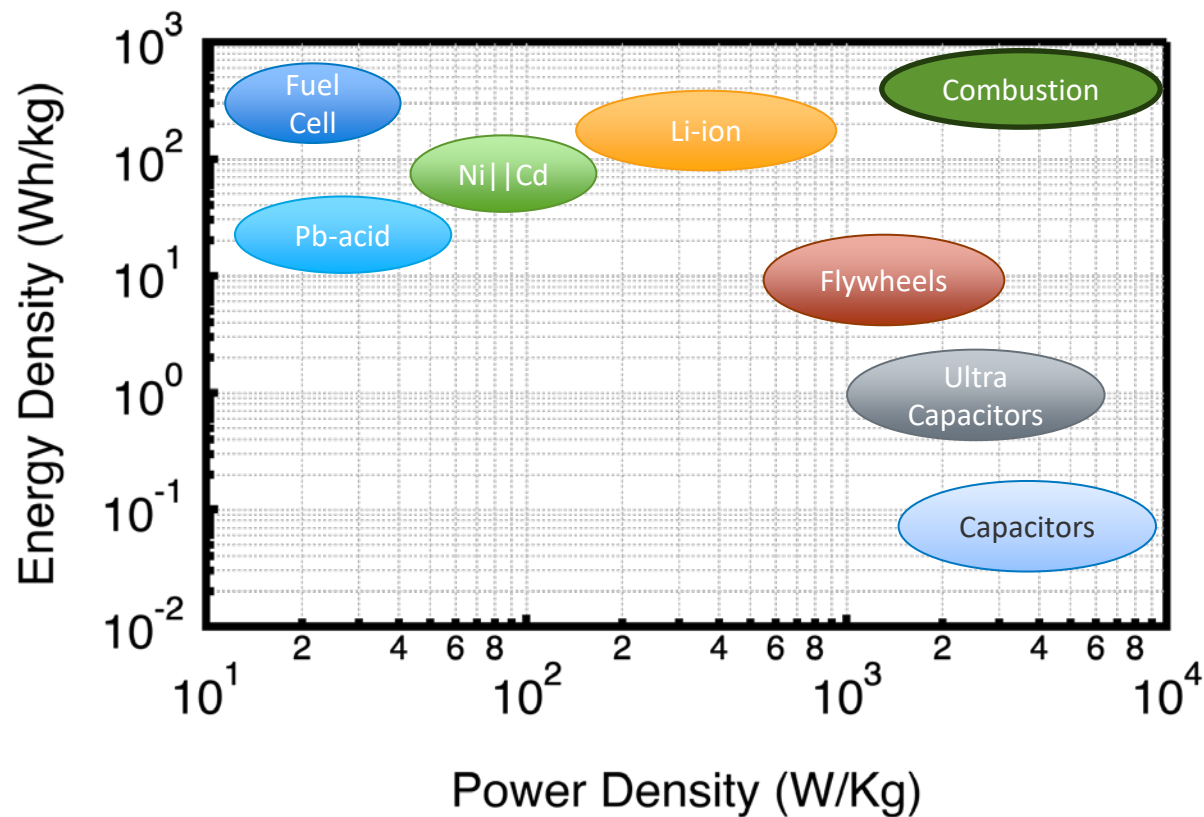
Cycle Life with Varied Surfaces



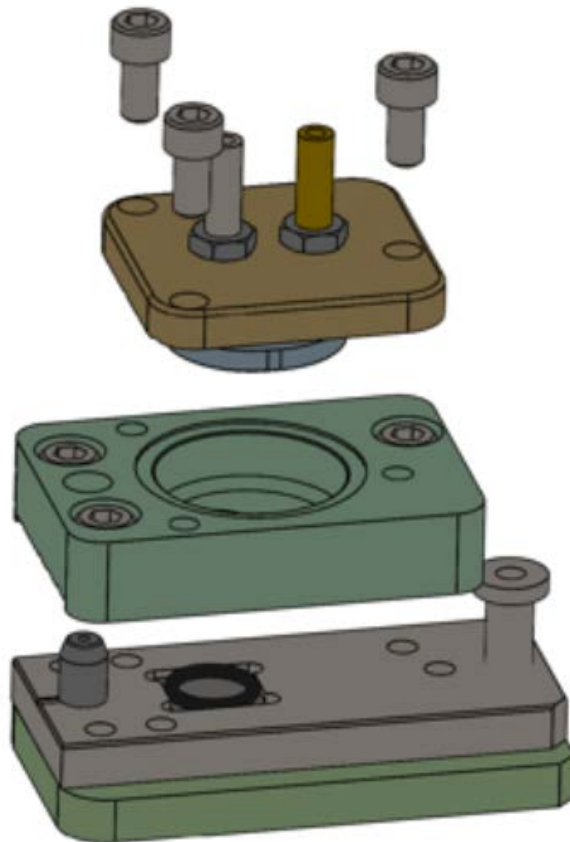
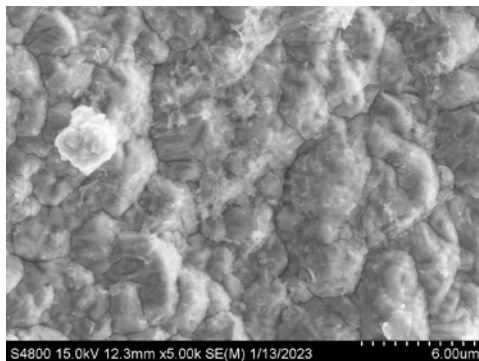
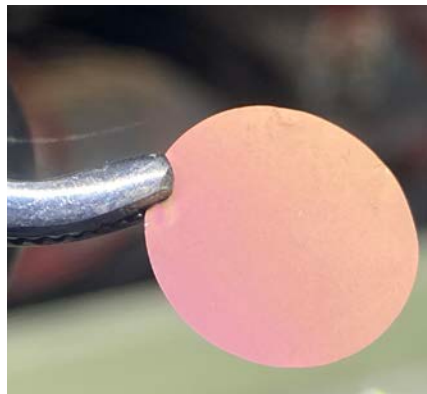
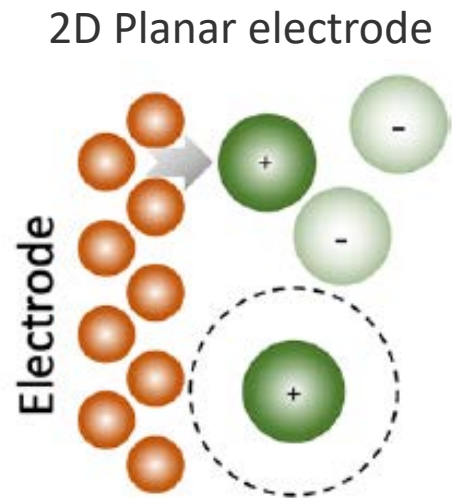
Schulze, et al.; *Batteries and Supercaps* **2023**



Energy Storage and Delivery Technology

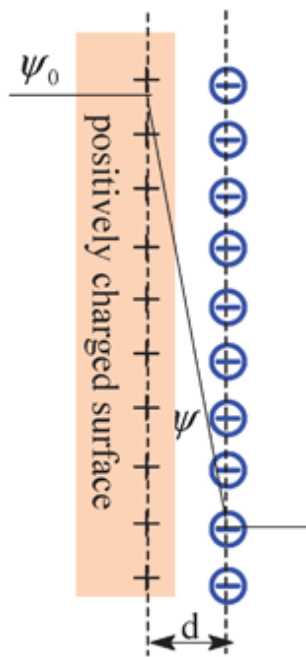


Thin Film Silicon Electrode

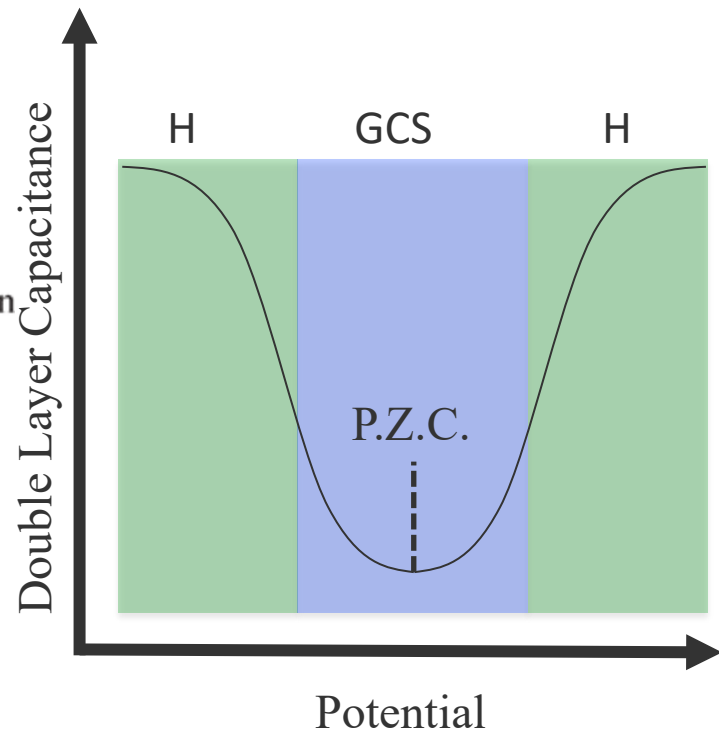
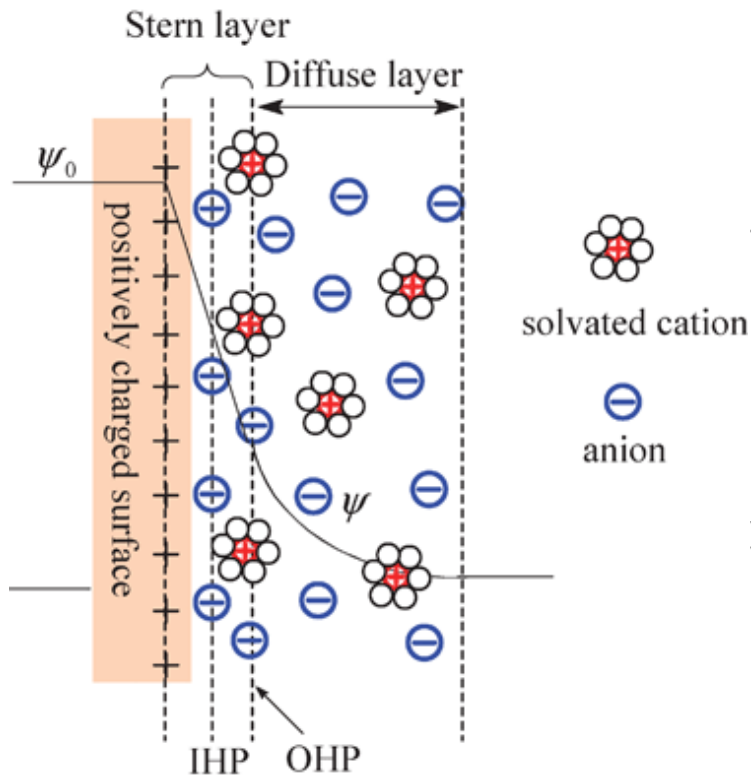


The Classical Model

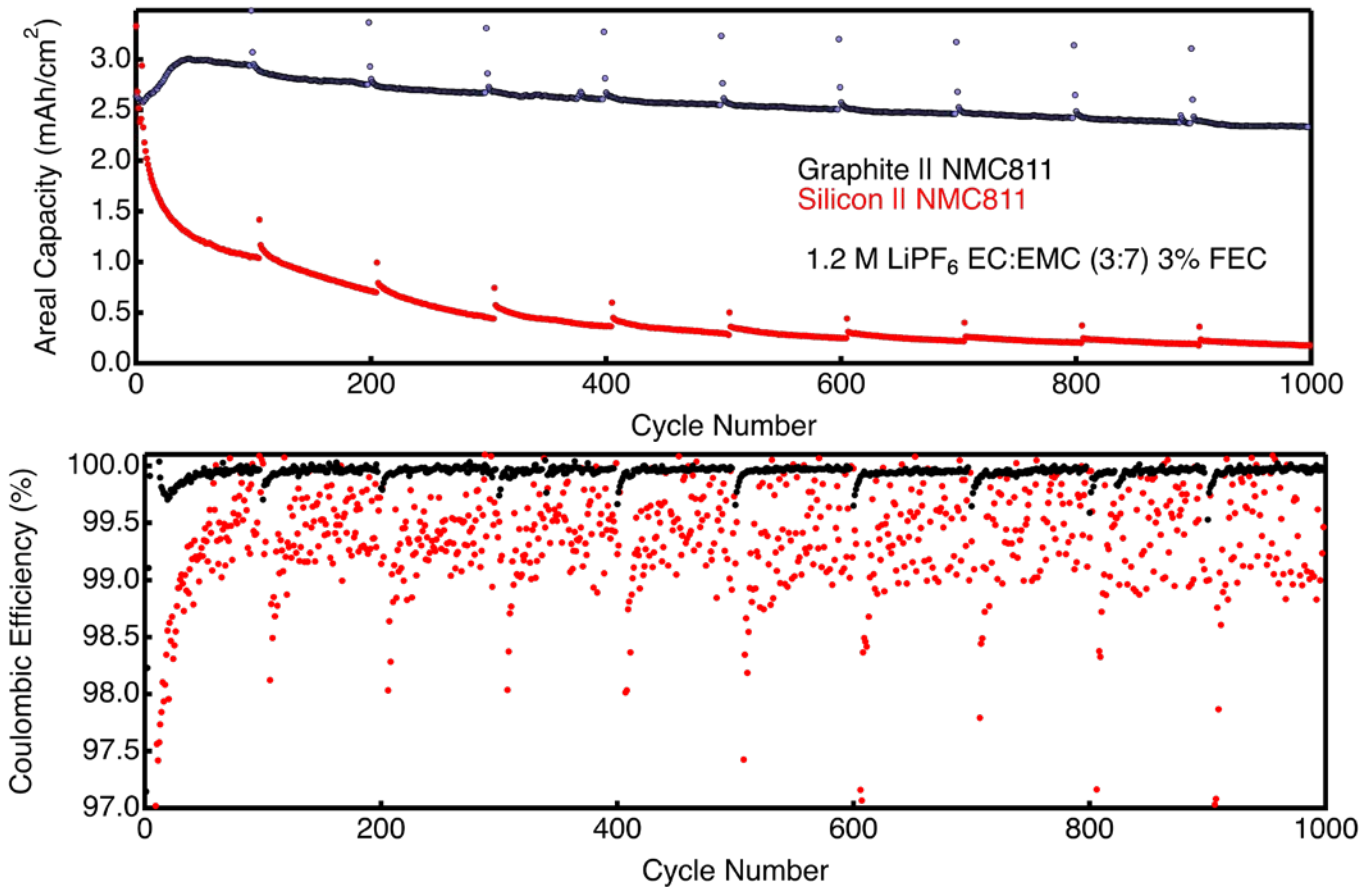
Helmholtz Model



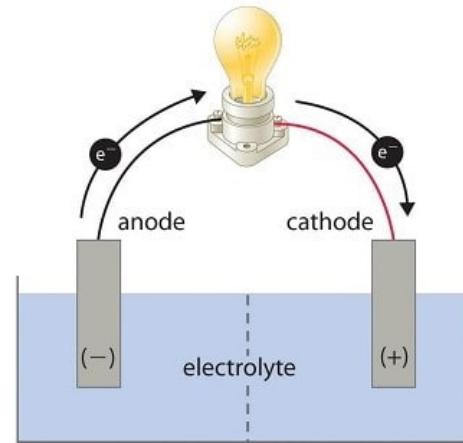
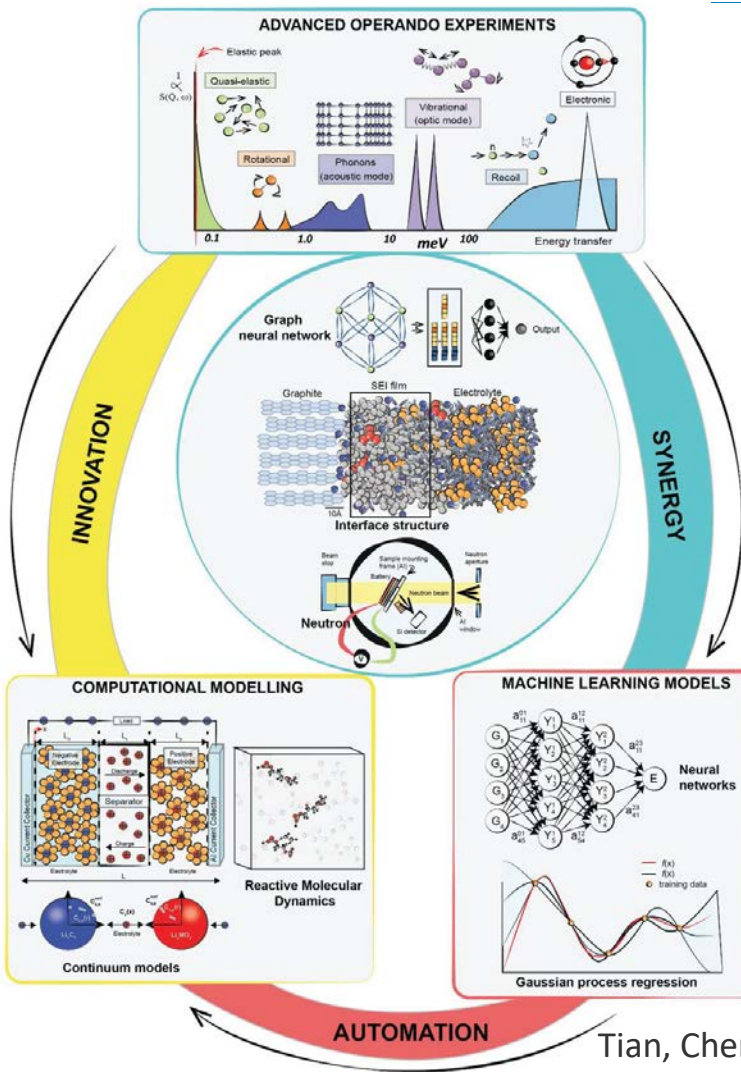
Gouy-Chapman-Stern Model



Electrochemical Cycling

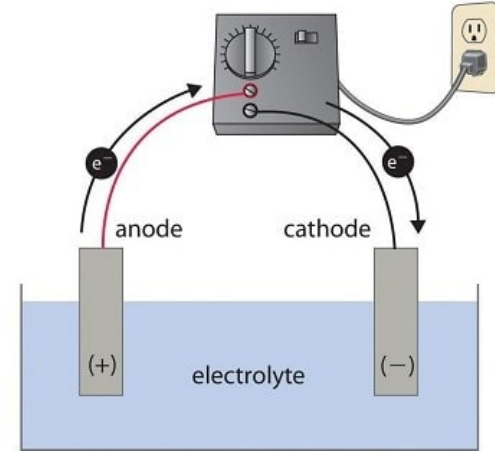


Understanding Calendar Life



GALVANIC CELL

Energy released by spontaneous redox reaction is converted to electrical energy.



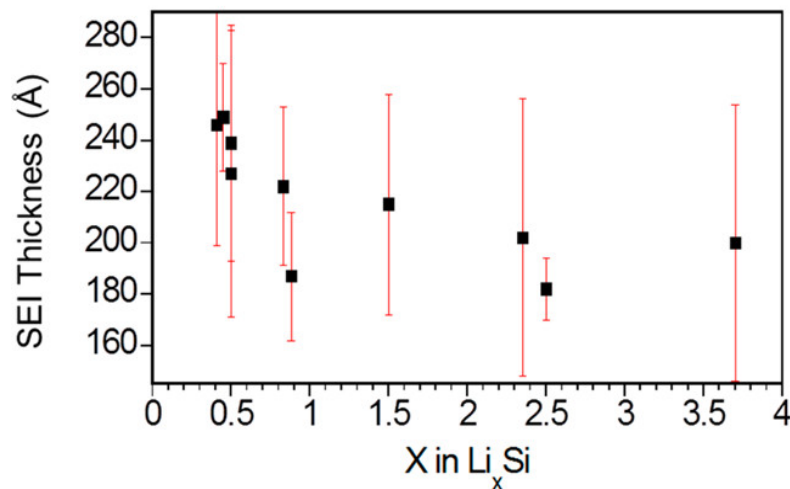
ELECTROLYTIC CELL

Electrical energy is used to drive nonspontaneous redox reaction.

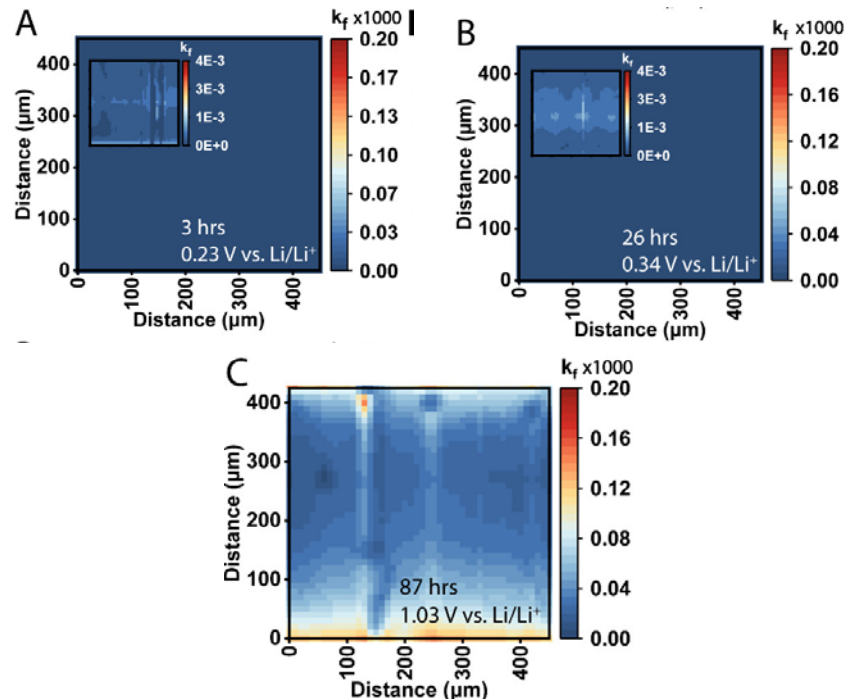
Energy Education, Univ. of Calgary

Corroborating Observations

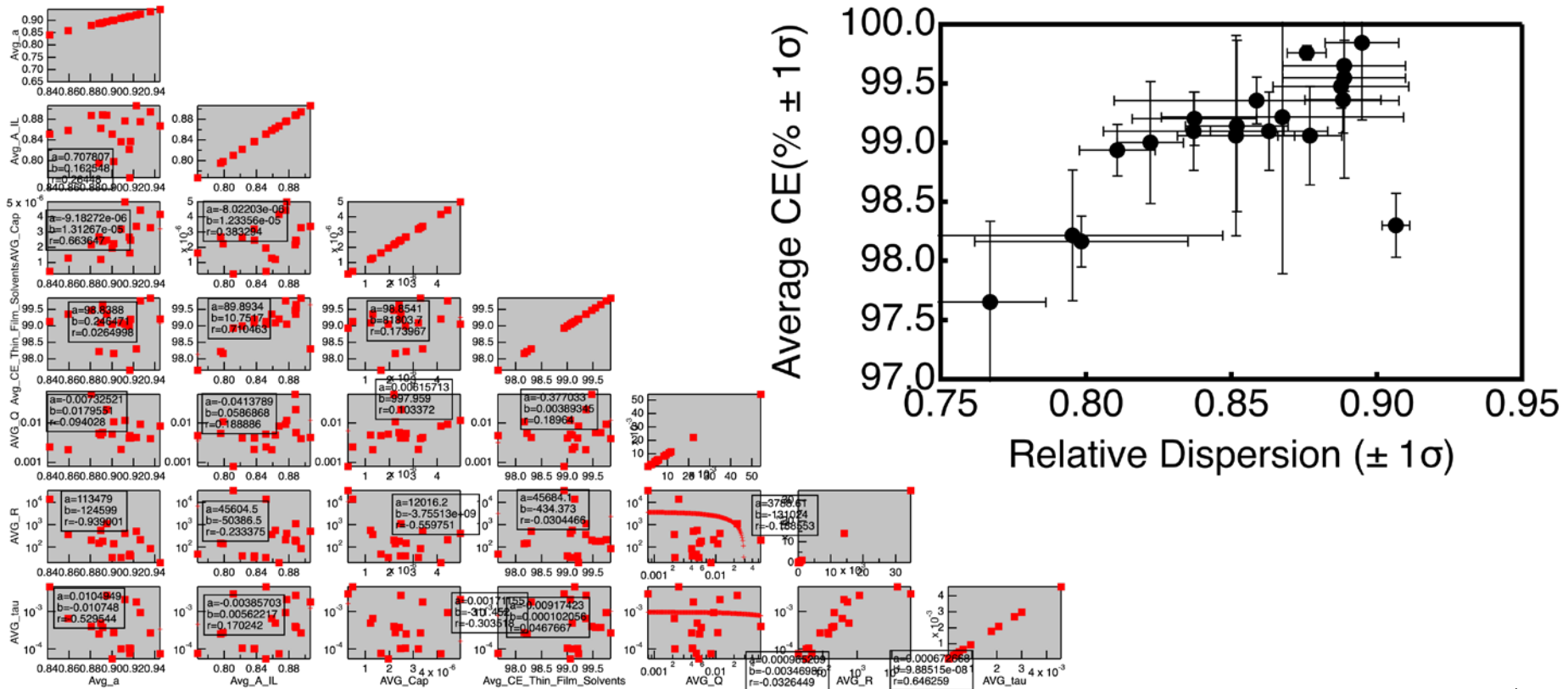
Neutron Reflectometry Measurements
of SEI thickness



Scanning Electrochemical Microscopy
Measurements on SEI Passivation

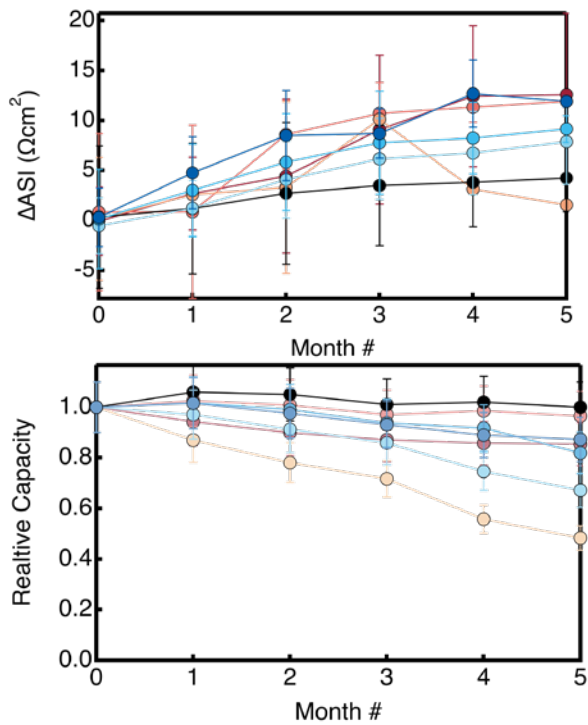


Predictive Parameters for Cycle and Calendar Life



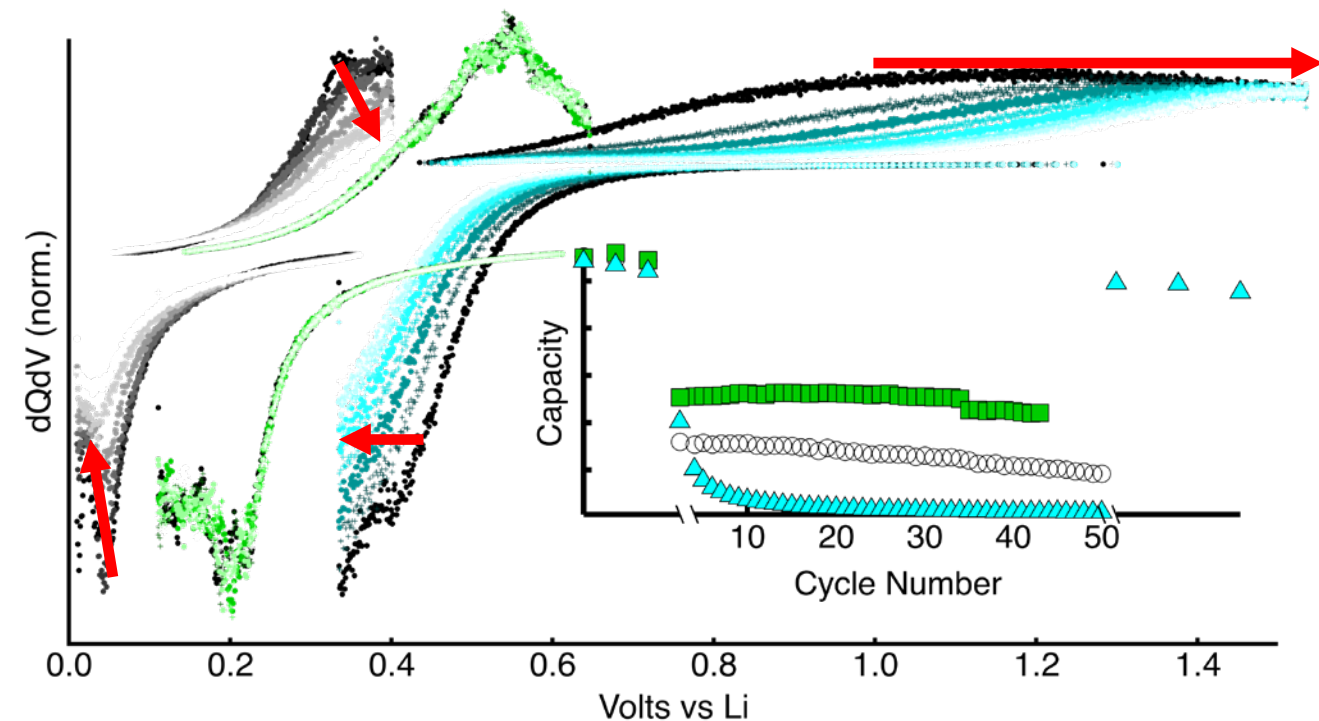
Calendar aging data in EC:EMC (Tier 2, Li⁺ limited)

LiPF₆ : LiTFSI : LiFSI : LiClO₄ :
LiBOB : LiDFOB : LiBF₄



When ranked for ASI gain and capacity retention, α shows promise as a predictor for calendar-related degradation

Impact in Real System



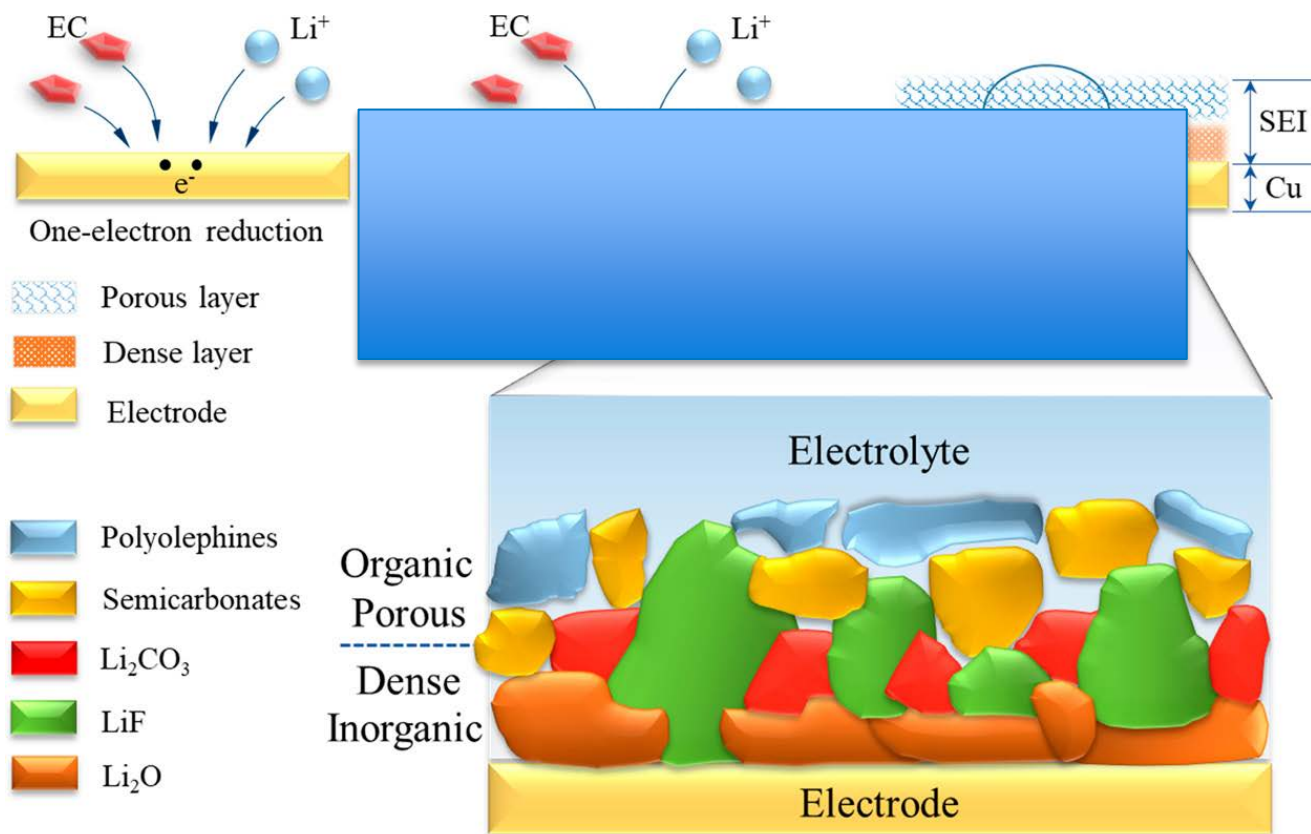
3 half cells formed between
0.01 – 1.5V

Then cycled in different
voltage windows

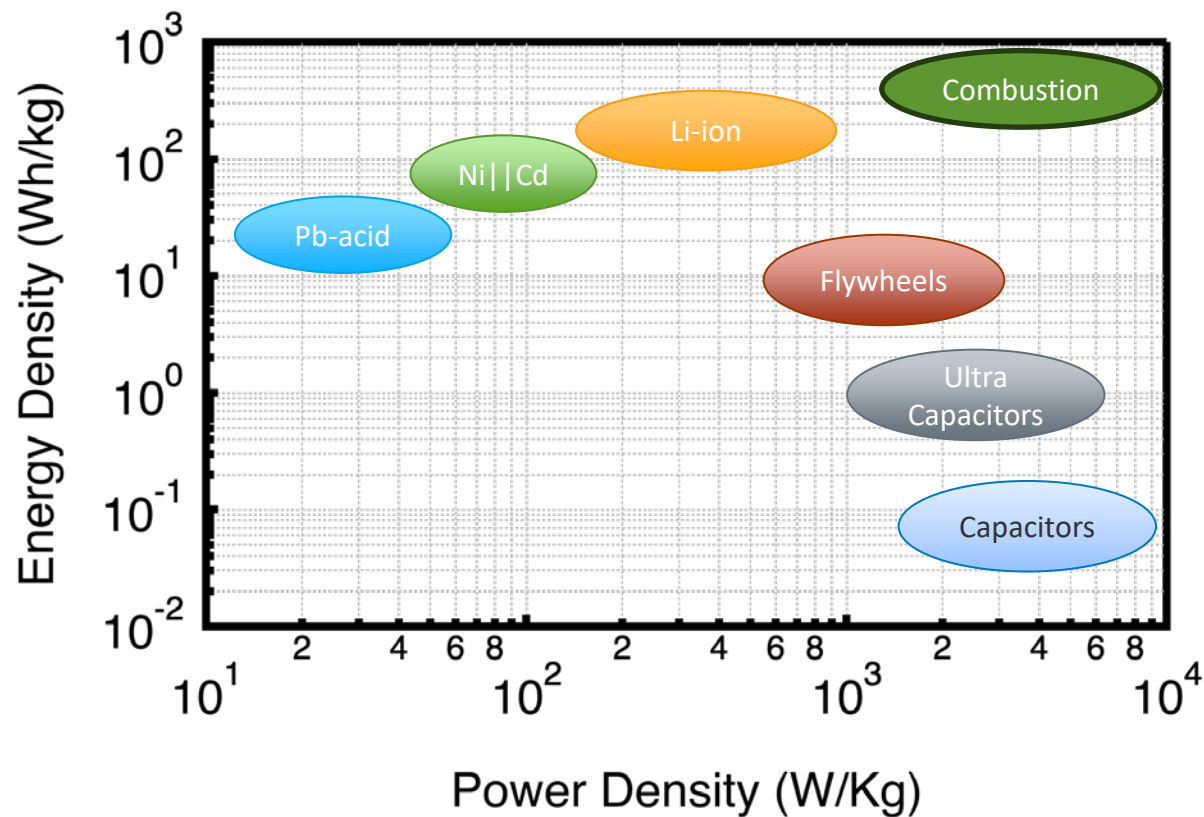
Low voltage electrode shows
increasing overpotential with
cycling consistent with
electrolyte reduction

Recovers all capacity when
cycled between 0.01 – 1.5V

SEI on Copper



Energy Storage and Delivery Technology



Electrode | Electrolyte Equilibration

