

# Characterizing Monovalent Cation Hydration Effects on Water Uptake of Ion Exchange Membranes

Elizabeth Wallace, Sara Abu-Obaid, Ngai Yin Yip  
Department of Earth and Environmental Engineering, Columbia university

YIP LAB  
SEPARATION INNOVATIONS  
FOR WATER, ENERGY, AND  
THE ENVIRONMENT



LIDLAW  
FOUNDATION

## Background

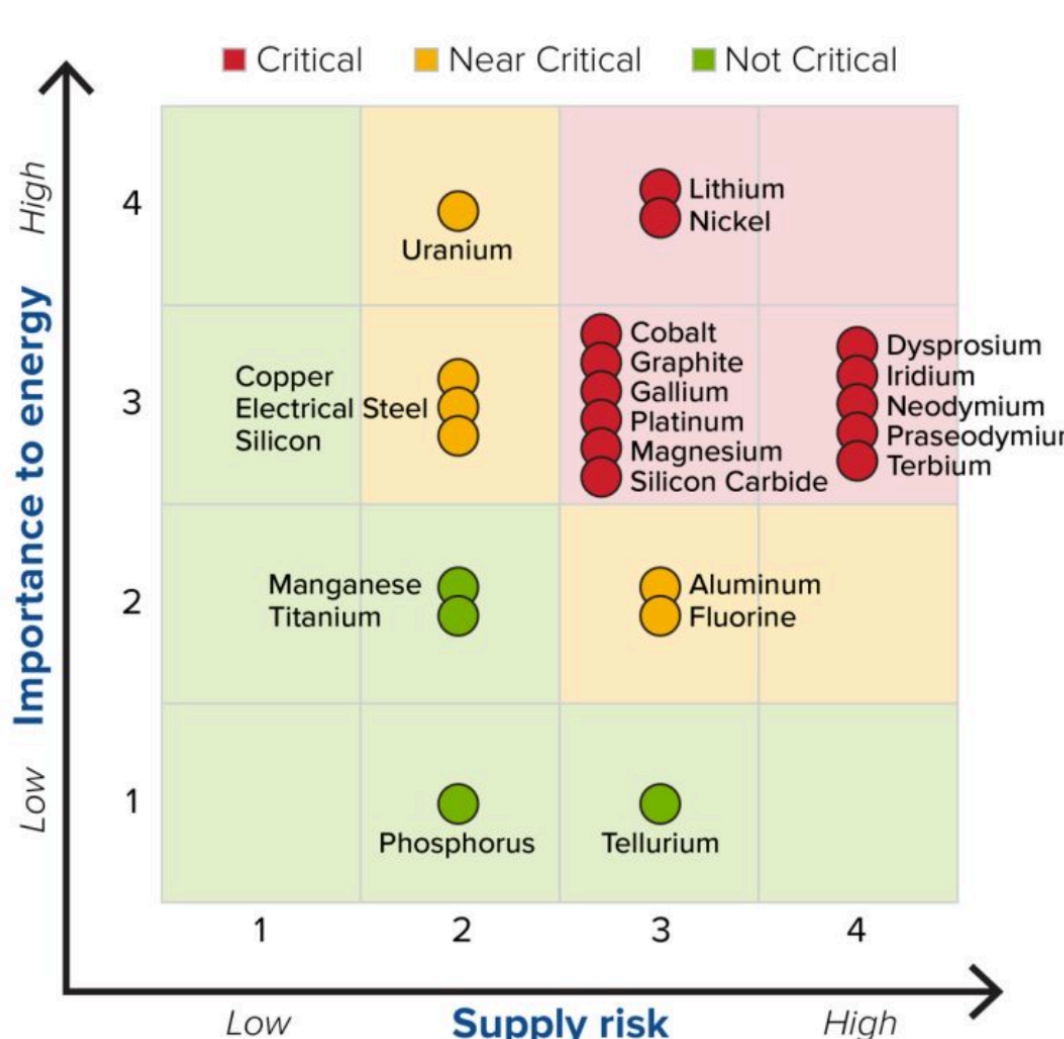


Figure 1: Supply risk vs. importance to energy of critical minerals.

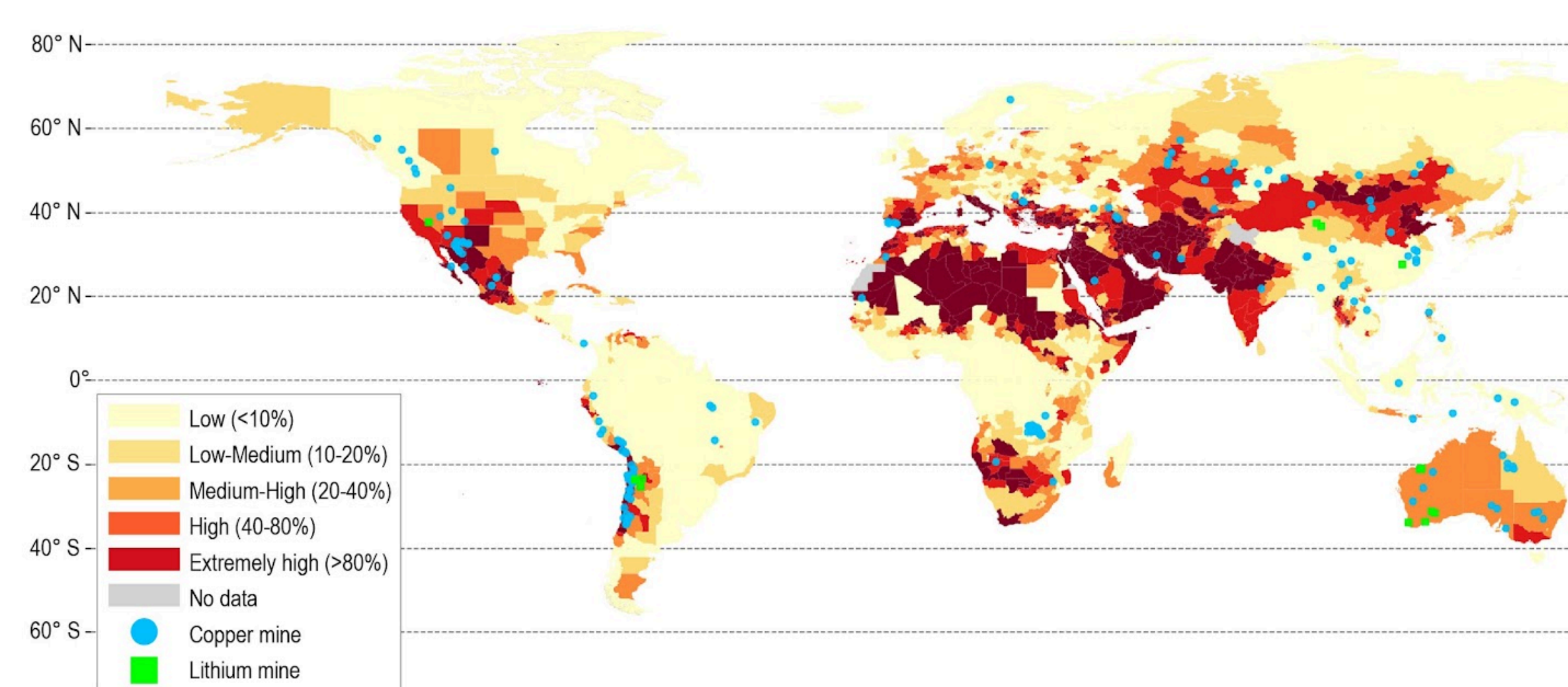


Figure 2: Global map of risk to drought events and prevalence of copper and lithium mines.

- Clean energy technologies are highly mineral-intensive.
- Lithium demand is growing fastest among critical minerals but faces high supply risk (Fig. 1).
- Risk to drought events in major lithium-producing regions, combined with water-intensive processing, highlights the need for sustainable water sourcing (Fig 2).

## Introduction

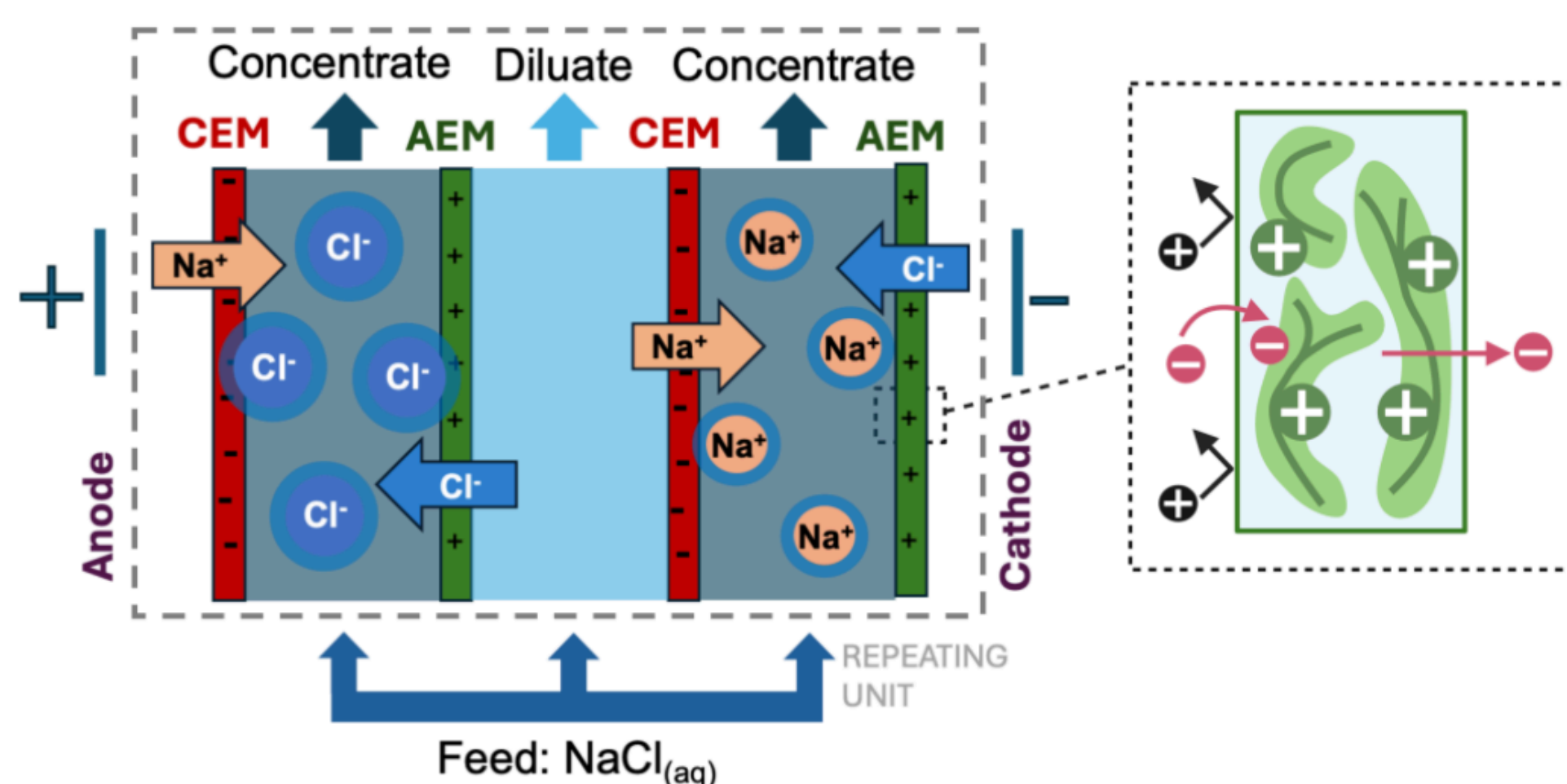


Figure 3: Schematic diagram of electrodesialysis. Ion transport begins with sorption into the membrane, then subsequent transport through.

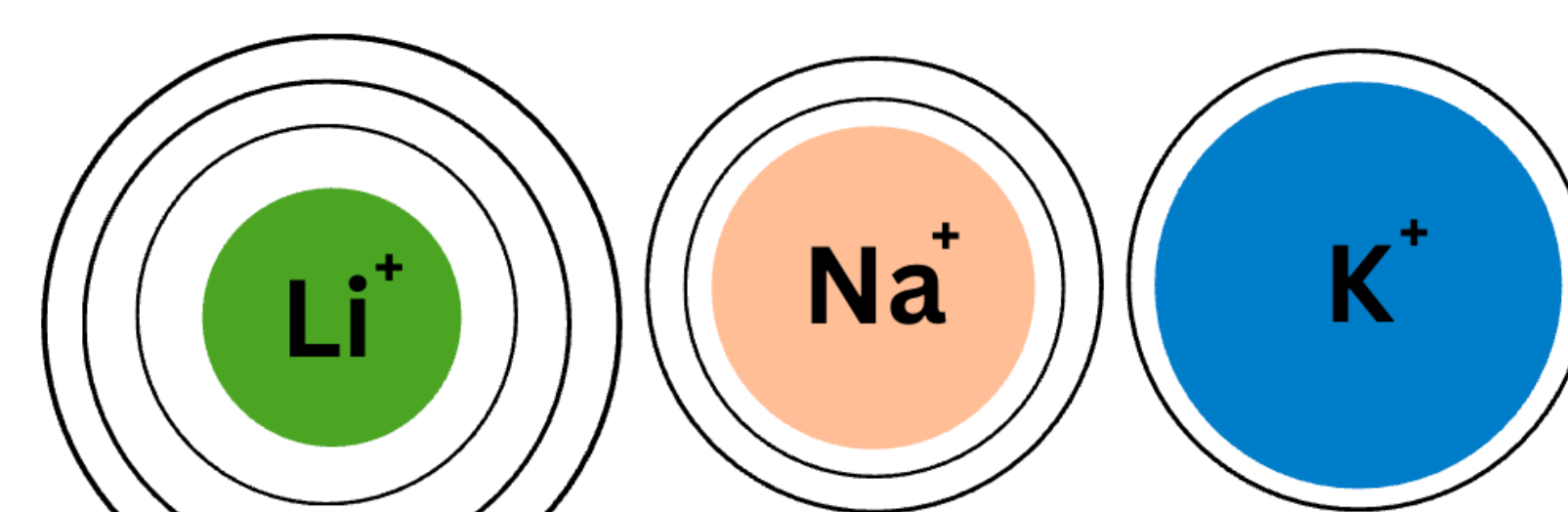
- Ion exchange membranes (IEMs) and electrodesialysis (ED) are used in various water, energy, and environmental applications.
- Membranes can be engineered for selective ion transport, enabling resource recovery.

## Problem Statement

Models describing ion exchange membranes do not distinguish between monovalent ions, although experiments demonstrated differing selectivity.

## Hypothesis

Li<sup>+</sup>, Na<sup>+</sup>, and K<sup>+</sup> have drastically different hydration energies, which could influence ion selectivity when entering the membrane.

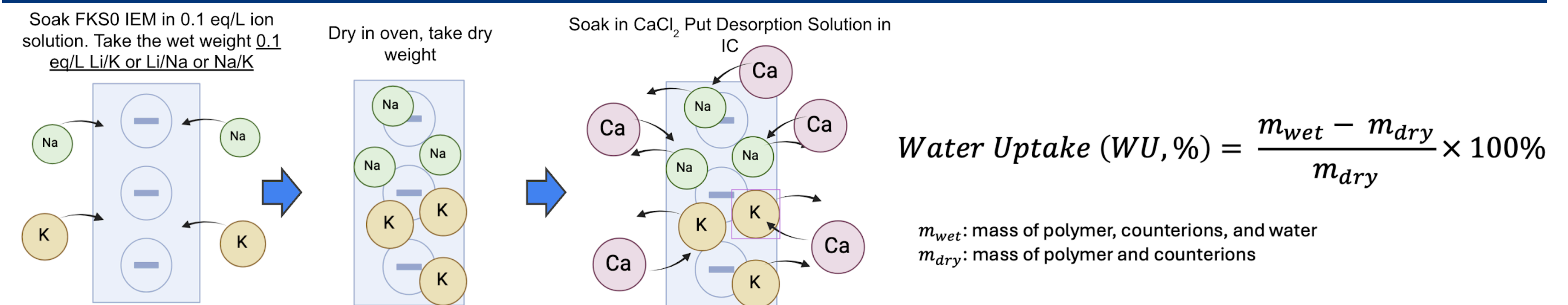


-520 kJ/mol      -406 kJ/mol      -322 kJ/mol  
Figure 4: Hydration energies of Li, Na, and K.

## Research Question

Do monovalent ions exhibit different hydrations within the membrane as they do in bulk solutions, and how does this variance affect selectivity?

## Methodology



3 Combinations of ions: Li only, Na only, K only, Li/K, Li/Na, Na/K  
5 different ratios for each combination of concentrations: 1:1, 3:1, 1:3, 9:1, 1:9

Figure 5: Schematic of water uptake procedure demonstrating the sorption and desorption process.

$$\text{Water Uptake (WU, \%)} = \frac{m_{\text{wet}} - m_{\text{dry}}}{m_{\text{dry}}} \times 100\%$$

$m_{\text{wet}}$ : mass of polymer, counterions, and water  
 $m_{\text{dry}}$ : mass of polymer and counterions

IEC of the membrane (meq/g membrane):

$$\text{IEC} = \frac{10 \times 10^{-3} \times C_d \times V_d}{m_{\text{dry}}}$$

$C_d$ : concentration (mmol/L)  
 $V_d$ : desorption solution volume (mL)

- A methodology was developed in which membranes were soaked, dried, and desorbed via soaking in CaCl<sub>2</sub> (Fig. 5).
- The first step accounts for  $m_{\text{wet}}$  value, the second step accounts for  $m_{\text{dry}}$  value, and final step accounts for Ion Exchange Capacity.

## Results

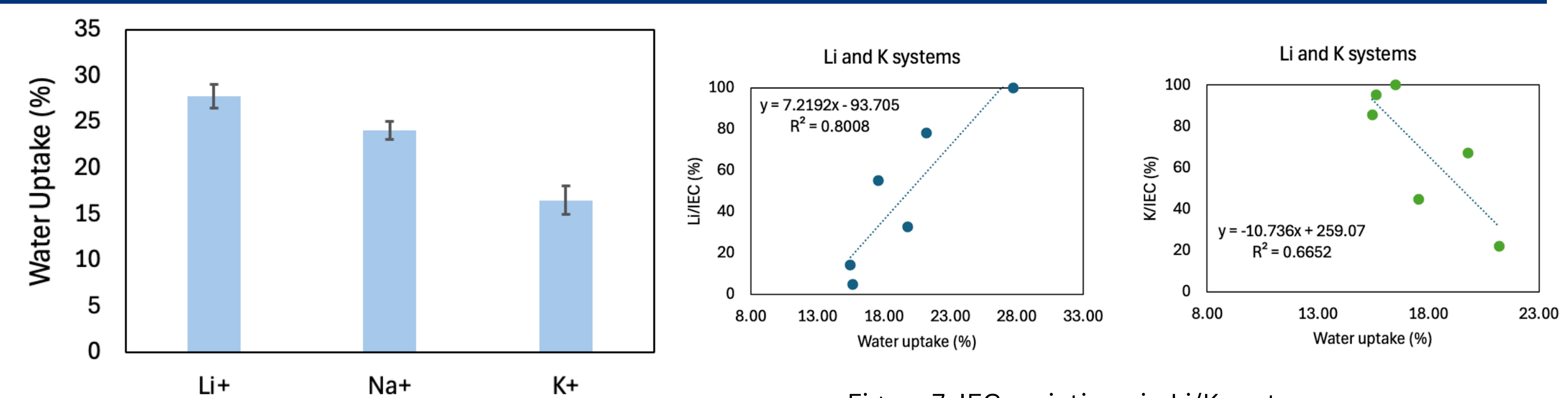


Figure 6: Water Uptake % of Li<sup>+</sup>, Na<sup>+</sup>, and K<sup>+</sup>.

- The water uptake of the membrane (i.e., water content) differs based on the type of ion within single salt systems, following the same trend as hydration energy.
- Li<sup>+</sup> presence strongly correlates with water uptake of membranes in binary systems.

Figure 7: IEC variations in Li/K systems.

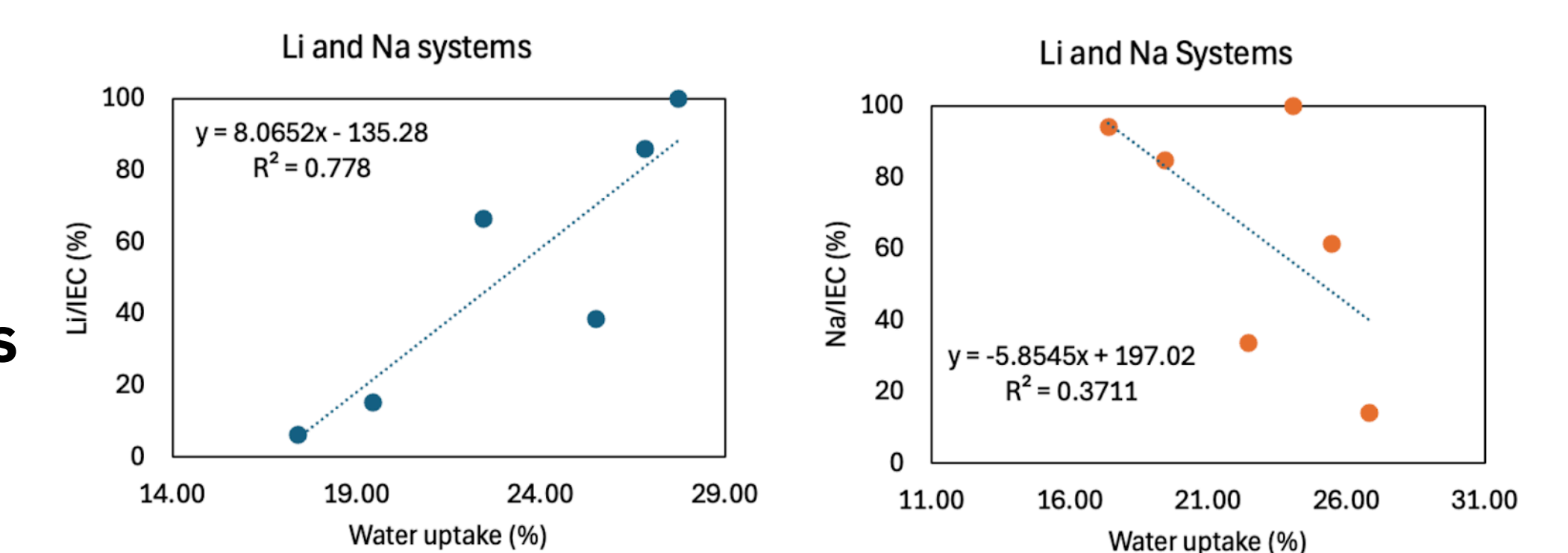


Figure 8: IEC variations in Li/Na systems.

## Discussions

There is a correlation between hydration energy and sorption selectivity that influences water uptake into the membrane.

## Future Work

Determine the quantity of water that is brought into the membrane with each type of ion, utilizing water uptake and sorption data from mixed salt systems.

## References

IEA (2021), The Role of Critical Minerals in Clean Energy Transitions, IEA, Paris  
<https://www.iea.org/reports/the-role-of-critical-minerals-in-clean-energy-transitions>,  
Licence: CC BY 4.0