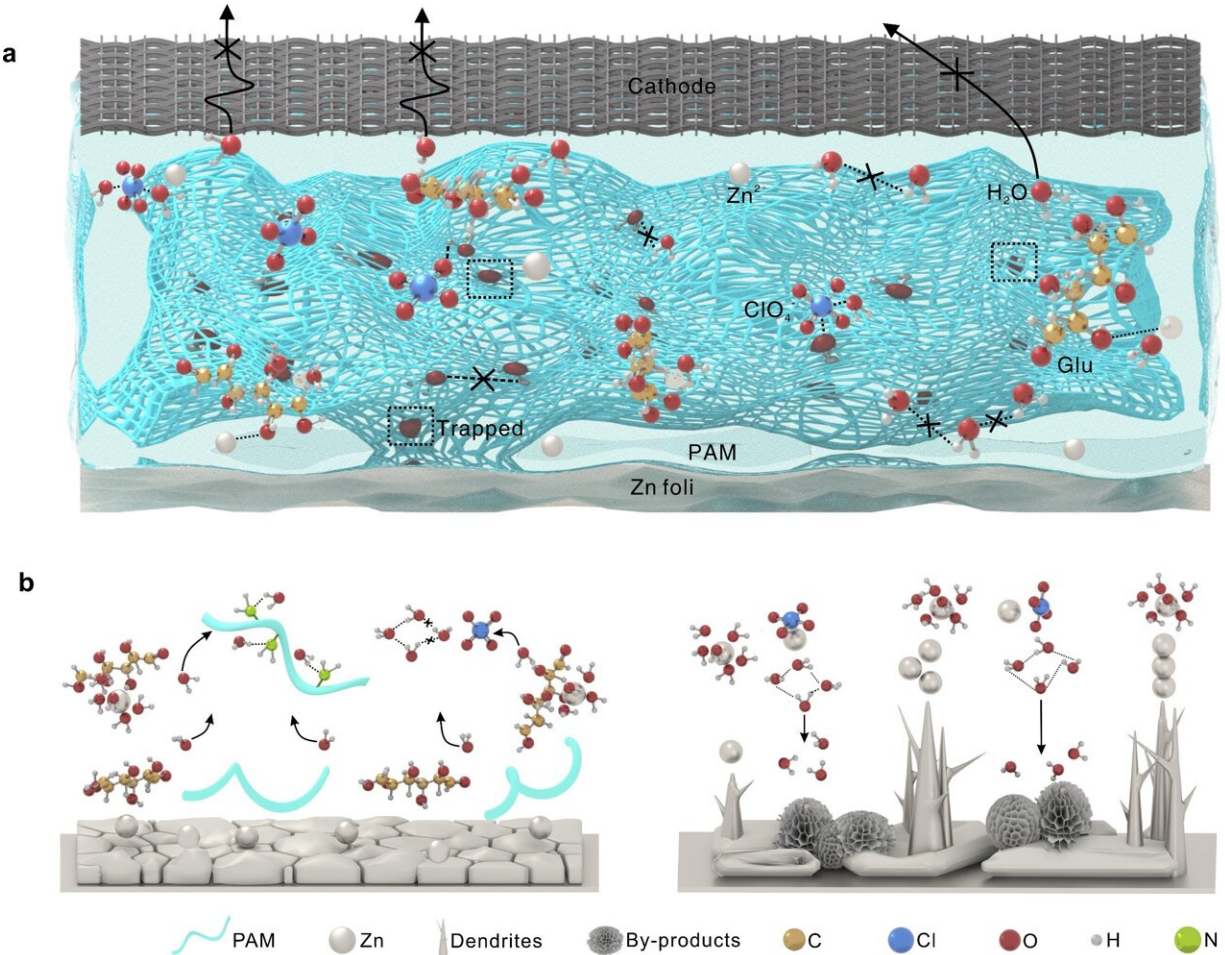


# Novel strategy proposed for all-climate zinc-ion batteries

June 14 2024, by Zhang Nannan



a. Schematic illustration of the design and construction of electrolyte structure;  
 b. Schematic illustration of Zn plating behavior in Glu/ZC/PAM (left) and pure ZC (right). Credit: Li Zhaoqian

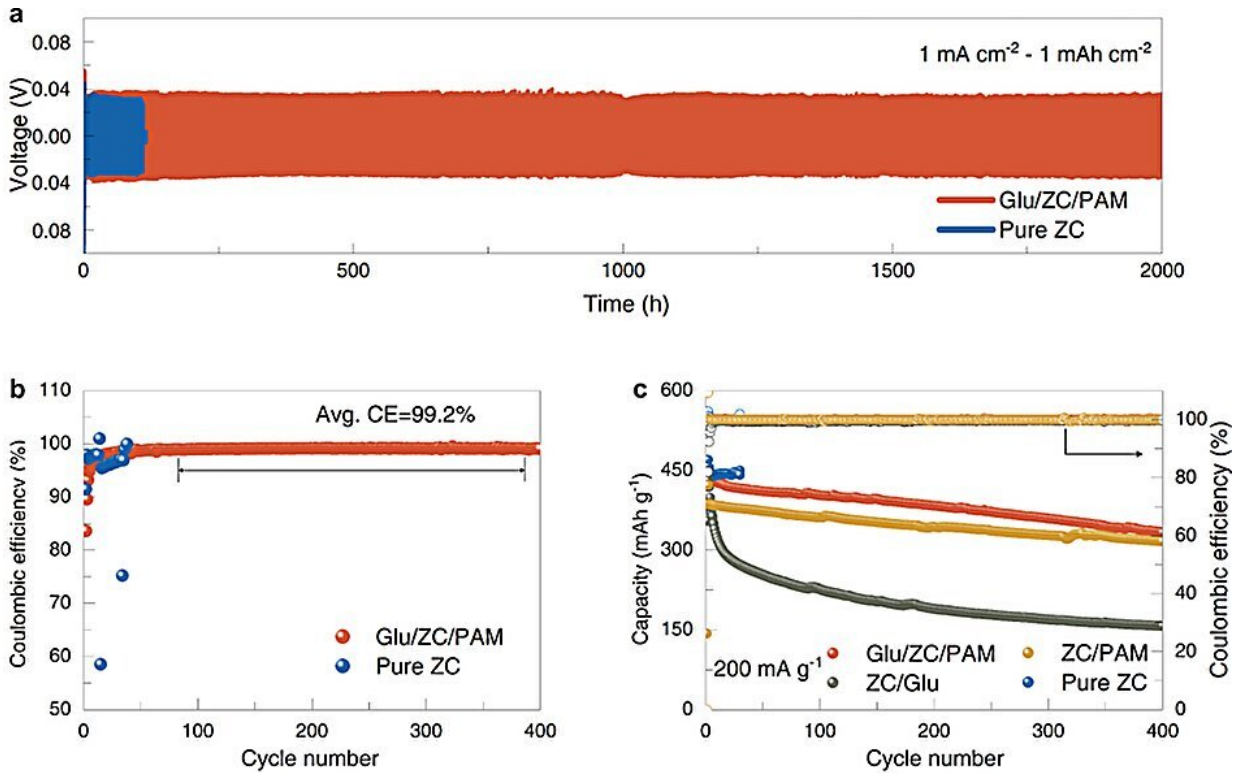
In a study [published](#) in *Advanced Energy Materials*, researchers have constructed a hydrogel electrolyte formula by using  $\text{ClO}_4^-$  anions and polyacrylamide chains to anchor water molecules, while glucose molecules preferentially regulate  $\text{Zn}^{2+}$  solvation.

Effectively interrupted [water clusters](#) and enhanced water covalency were realized, resulting in an expanded voltage stability window and stable operation over a wide temperature range.

"This means that the aqueous zinc batteries could operate stably considering the seasonal and altitude factors. Importantly, the temperature resistance mechanism in the water environment,  $\text{Zn}^{2+}$  solvation and Zn/[electrolyte](#) interface are systematically analyzed," said Li Zhaoqian, a member of the team. The research team was led by Prof. Hu Linhua from the Hefei Institutes of Physical Science of the Chinese Academy of Sciences.

Irreversible electrolyte phase transitions and an accelerated parasitic reaction greatly threaten the climate adaptability of aqueous Zn-ion batteries. Water activity affects the freezing point of the electrolyte, the voltage stability window, and interfacial Zn deposition behavior. Due to its anti-leakage property, polymer structure stability, and numerous anchoring sites for free water, the hydrogel electrolyte's rational design efficiently improves the battery's climate adaptability.

In this study, the researchers constructed a "covalency reinforced" hydrogel electrolyte with superior interfacial adhesion and strong moisture-retaining ability. Through [spectral analysis](#) and theoretical calculations, they revealed weakened bulk water activity and regulated  $\text{Zn}^{2+}$  solvation, which delayed the freezing point of the electrolyte, facilitated its moisture-retaining capacity, and inhibited water-induced side reactions.



The electrochemical properties comparison. Credit: Li Zhaoqian

COMSOL simulation and morphological evolution show the improved mechanical properties of the electrolyte and the thermodynamically stable Zn interface. These advantages resist dendrite formation and solve electrode–electrolyte contact problems, giving the batteries a wide operating range of  $-40\sim 130^{\circ}\text{C}$ .

"When the electrolyte is used in pouch batteries, it shows an impressive capacity of  $254 \text{ mAh/g}$  at  $-30^{\circ}\text{C}$  and  $438.1 \text{ mAh/g}$  at room temperature. This is a big deal because most previous batteries didn't go beyond  $200 \text{ mAh/g}$  at  $-30^{\circ}\text{C}$  or  $400 \text{ mAh/g}$  at room temperature. This work shows how effective these batteries are, both in terms of capacity and their

ability to operate over a wide range of temperatures," said Dr. Li.

They also assembled the Zn//Zn and Zn//Cu batteries to evaluate stable lifespan and Zn plating/stripping reversibility. At low current density, the lifetime of the Zn anode exceeds 2,000 hours, which is better than that of the liquid electrolyte. Even at high current density, the battery with Glu/ZC/PAM can work steadily for more than 500 hours.

The Zn//Cu batteries could work steadily for more than 800 hours with a high average Coulomb efficiency of 99.2%, highly competitive with previous hydrogel electrolytes.

This study modulates the coordination structure and tailors thermodynamic activity between the electrolyte/Zn interface by employing a multifunctional hydrogel electrolyte, which degenerates detrimental parasitic reactions and extends the operating temperature range. It provides a safe and highly efficient strategy to realize all-climate aqueous zinc-ion devices.

**More information:** Yifan Wang et al, Regulating Water Activity for All-Climate Aqueous Zinc-Ion Batteries, *Advanced Energy Materials* (2024). [DOI: 10.1002/aenm.202402041](https://doi.org/10.1002/aenm.202402041)

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