



Quest for Grid Energy Storage: Case for the Performance of Iron-Ion/Hydrogen Redox Flow Battery Mixed Electrolytes

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FLORIDA ENERGY SYSTEMS CONSORTIUM

May 20th, 2015

RFB PRACTICAL APPLICATIONS

Iron-Ion/Hydrogen Redox Flow Battery

Flow batteries are just one technology that can store electricity but they could be amongst the cheapest and most versatile for large scale storage

CATCHING THE WIND

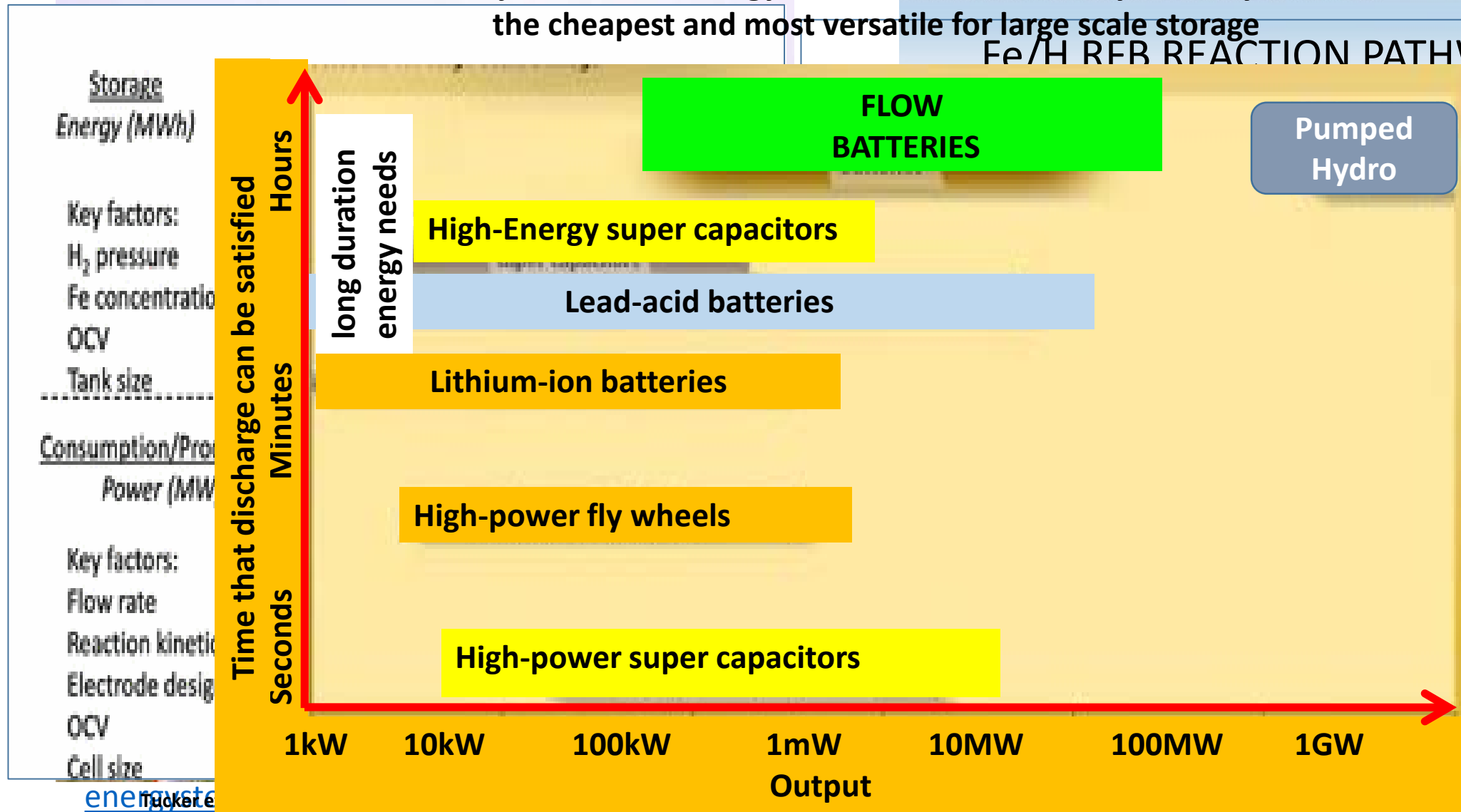
Fe/H RFB REACTION PATHWAY

When wind power alone cannot generate enough electricity to supply demand, the pumps on the flow cell switch on, pumping the two vanadium solutions through the reactor. Electrons are released at the negative electrode, and hydrogen ions move across the membrane to balance. The electricity that is generated is fed to the power grid



1.00 V vs. SHE

1.77 V



energy storage

OBJECTIVE

- Determine the influence of electrolyte on Iron-Ion/Hydrogen Redox Flow Battery using Cyclic voltammetry (CV) techniques

ELECTRON EFFICIENCY

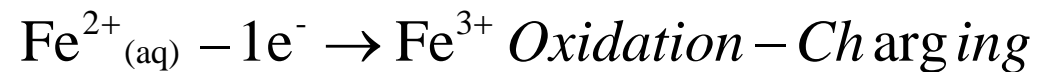
$$\text{Electron efficiency } (\eta) = \left(\frac{\text{Number of electrons from Randles – Sevcik Eqn.}}{\text{Number of electrons in balanced Redox Eqn.}} \right)$$

- Randles-Sevcik equation at room conditions:

$$I_p = 2686000 N^{3/2} A D^{1/2} C v^{1/2} \quad \longrightarrow \quad N = \left(\frac{I_p}{2686000 A D^{1/2} C v^{1/2}} \right)^{2/3}$$

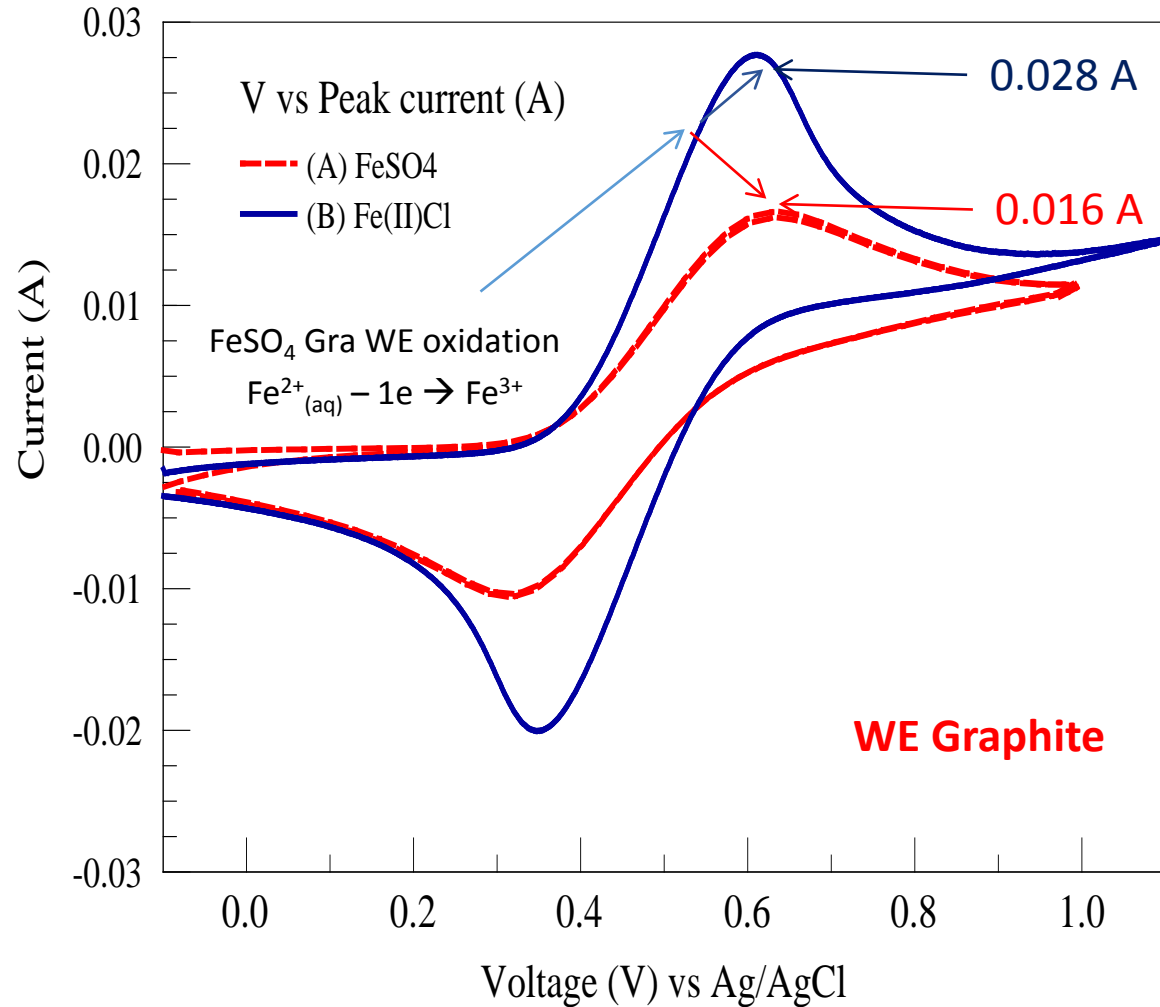
Where;

- N = Number of electron
- A = Area of electrode (cm²)
- C = Species concentration (mol/cm³)
- D = Diffusion coefficient (cm²/s)
- V = Scan rate (V/s)

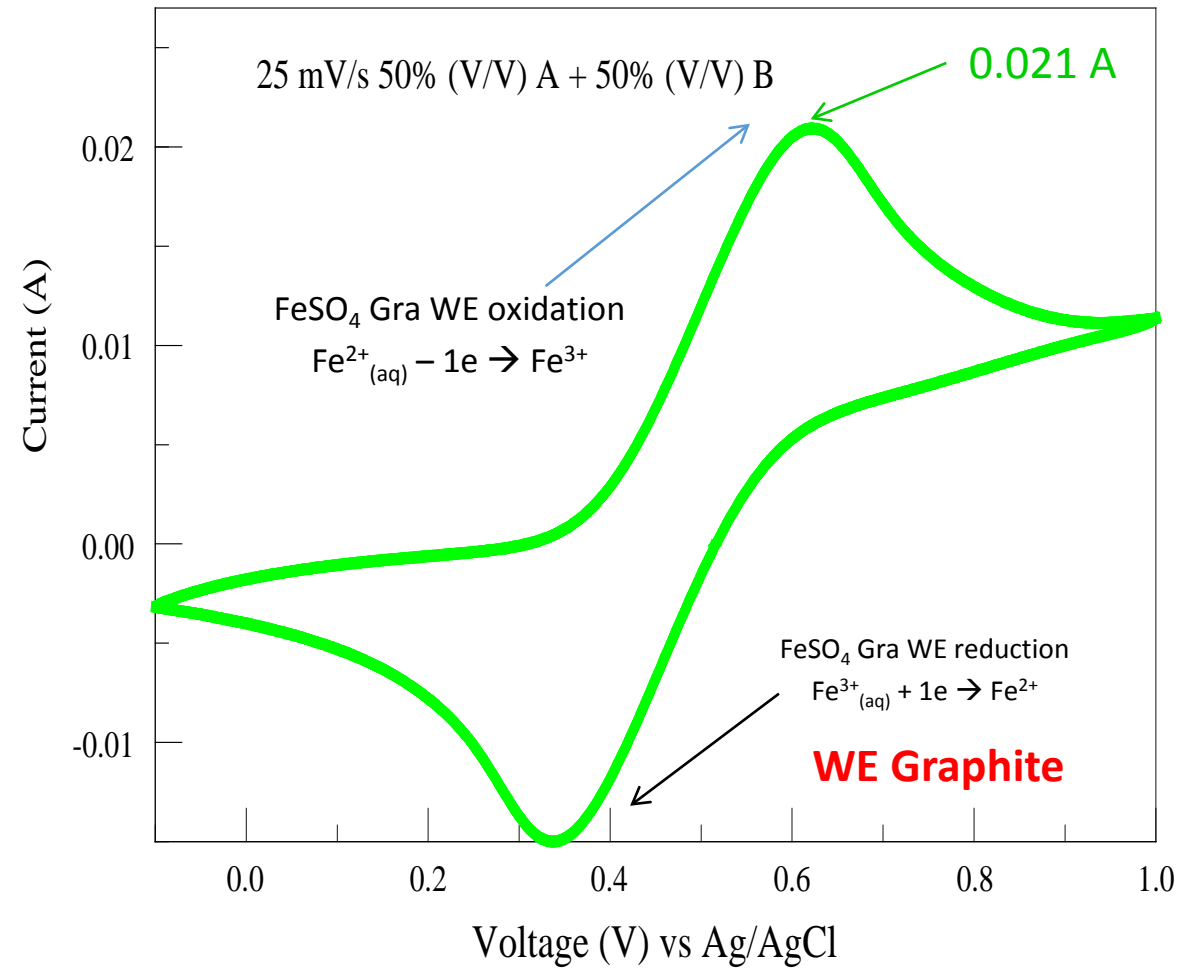


Mixed Electrolyte: Improve charge/discharge of Fe-H RFB electrolyte

BEFORE MIXING



AFTER MIXING





Influence of Mixed Electrolyte on the Performance of Iron-Ion/Hydrogen Redox Flow Battery

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Improved charge/discharge performance of Iron-ion/Hydrogen redox flow battery (RFB) electrolyte with a mixed FeSO₄ and FeCl₂ is reported. Addition of Cl⁻ ions into a sulfate electrolyte changes the charge/discharge behavior of the sulfate electrolyte leading to a reduction in charging potential for a mixed FeSO₄ and FeCl₂ electrolyte system. This suggests that a sulfate/chloride electrolyte system can lead to improved charge/discharge of the Fe-ion/H₂ RFB. Reverse addition of FeSO₄ to FeCl₂ showed a decrease in the mixed electron transfer efficiency (experimental current relative to theoretical) equivalent to a decrease in electrolyte performance. We deduce that 0.8 M FeCl₂ corrosive electrolyte can be replaced by less corrosive mixture of 46 mol % Cl⁻ in 0.8 M FeSO₄ to achieve the same performance that can be obtained using an all chloride system.

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Manuscript submitted January 7, 2015; revised manuscript received April 27, 2015. Published May 15, 2015. This was Paper 965 presented at the Cancun, Mexico, Meeting of the Society, October 5–9, 2014.