

Hydrogen production efficiency in SO₂ Depolarized Electrolyser - Impact of proton electrolyte membrane



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Introduction

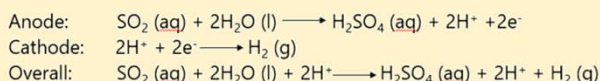
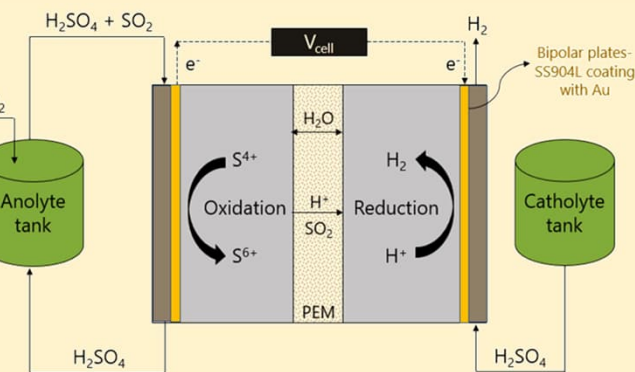
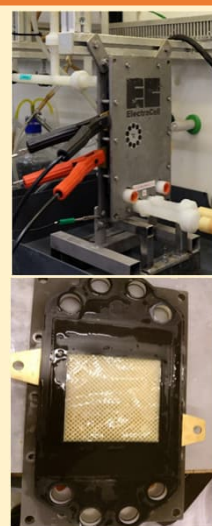
The SO₂ Depolarized Electrolyser (SDE) is a breakthrough in the production of hydrogen on a large scale. The working of the SDE can be explained as:

- Oxidation of SO₂ from +4 to +6 oxidation state to produce Protons (H⁺) and Sulfate ions (SO₄²⁻) at the anode^[1]
- The protons flow through a polymer electrolyte membrane (PEM) and reduce at the cathode to produce hydrogen.
- Along with protons, PEM also allows SO₂ to pass through leading to parasitic reactions at the cathode.
- SDE has lower standard reversible voltage, E_{0,SDE} = 0.158V, compared to PEM water electrolyser with E_{0,PEM} = 1.23V^[3].

The goal is to determine the effect of using different membranes in SDE to make it a viable technology for producing renewable hydrogen on a large scale by preventing parasitic reactions at the cathode.

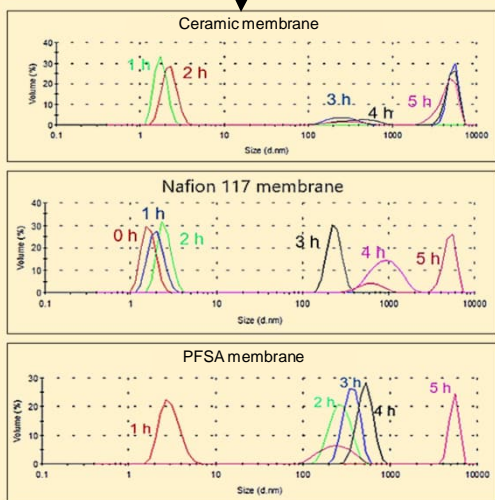
Methodology

- Bench-scale electrolyser setup with Au electrocatalyst^[2] was used
- Three membranes tested: Nafion 117, Ceramic, and PFSA – pre-commercial membrane
- Initial operation without SO₂; introduced after 500 s
- SO₂ feed halted after 1 hour, operation continued with 350 mM SO₂ for 5 hours
- Operating conditions: 15 wt% H₂SO₄ as electrolyte, 1.7 V operating potential
- Photon Correlation Spectroscopy^[1] (PCS) used to analyze sulfur particle formation in the catholyte.

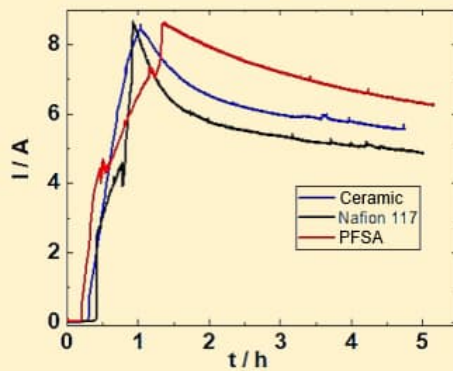


Sulphur particles observed in catholyte

- Small sulfur particles: 1-10 nm
- Large sulfur particles: 100-10000 nm
- Particle growth begins after 2 hours with PFSA, 3 hours with Nafion, and 5 hours with Ceramic.



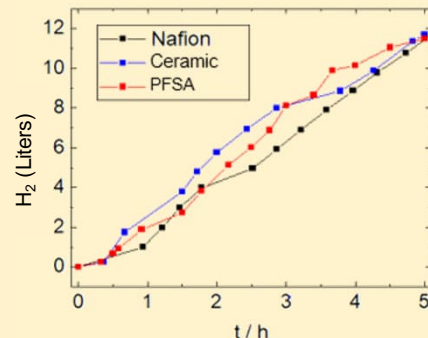
Results and Discussion



Hydrogen produced with time
Cumulative hydrogen production after 5 hours is similar for all three membranes

Performance of SDE based on current produced

- Similar maximum current density observed for all separators.
- Nafion 117 and ceramic membranes exhibit faster current decrease compared to PFSA; after 5 hours, the current is 5 A for ceramic, 6 A for Nafion, and 6.5 A for PFSA.



Conclusion

- The pre-commercial PFSA membrane shows steady performance in hydrogen production efficiency
- Additional testing needed to investigate SO₂ crossover using diffusion cell experiments, to understand the differences in different separator materials
- **Next steps:** Identify and validate different strategies to prevent SO₂ crossover, such as membrane coating or electrolyte solution modification.

References

- (1) Santasalo-Aarnio, A.; Virtanen, J.; Gasik, M., SO₂ Carry-over and sulphur formation in a SO₂-depolarized electrolyser. Journal of Solid-State Electrochemistry 2016, 20, 1655-1663. <https://doi.org/10.1007/s10008-016-3169-8>
- (2) Santasalo-Aarnio A, Lokkila A, Virtanen J, Gasik MM. Performance of electrocatalytic gold coating on bipolar plates for SO₂ depolarized electrolyser. Journal of Power Sources. 2016 Feb 29;306:1-7.
- (3) Gorensek, M.B., et al., A thermodynamic analysis of the SO₂/H₂SO₄ system in SO₂-depolarized electrolysis. international journal of hydrogen energy, 2009. 34(15): p. 6089-6095.

