



CHANNEL

Thulile Khoza, AEM webinar, 6th July 2021

Development of the most Cost-efficient Hydrogen production unit based on ANioN exchange membrane ELectrolysis

11.2 Joint Undertaking under grant agreement No 875088. This Joint undertaking receives support from the and Hydrogen Europe and Hydrogen Europe Research





AEMWE an emerging electrolyser technology and has the potential to combine the best of both worlds.....



European Union's Horizon 2020 research innovation programme and Hydrogen Europe and Hydrogen Europe Research

S. Gottesfeld, D. R. Dekel, M. Page, C. Bae, Y. Yan, P. Zelenay, Y. S. Kim, Journal of Power Sources 375 (2018) 170-184



Challenges for AEMWE

- Anion exchange membrane, conductivity, chemical and mechanical stability
- Active and durable PGM-free catalysts
- Electrode design and three phase boundary
- Current collector and BPP materials and design







- QA-based polymers cationic groups are susceptibility to hydroxide attack, leading to reduced IEC. Low KOH concentrations, DI water operation preferred.
- Ni and Ni based alloy nanostructures with high surface area may become unstable over time, loss of active area/dissolution





• AEI conductivity and stability, binding properties, catalyst poisoning and membrane interface



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CREATE Enapter

SINTEF

• Topic: FCH-02-4-2019: New Anion Exchange Membrane Electrolysers

- Duration: 2020-2022
- Total budget: 2 M€
- From TRL 2 to TRL3

The aim of CHANNEL is to design, construct and test a cost-efficient, 2 kW AEM water electrolyser stack and balance of plant able to operate at differential pressure.

The electrolyser will be based on low-cost materials, including non-PGM electrocatalysts, porous transport layers and bi-polar plates, performing at < 1.85 V per cell at 1 A cm⁻², using diluted KOH electrolyte at a system capital cost of < 600 ϵ/kW







Specific Objectives

- To further develop best-in-class EVONIK polymer materials and deliver > 40 units of large area AEM membranes > 500 cm² fulfilling the membrane and ionomer KPI's according to the FCHJU objective 2.4-2019;
- 2. To further optimise nanostructured Ni-based electrocatalysts with respect to activity and durability for the HER and OER;
- To optimise coating methods, catalyst loading, as well as ionomer type and loading in order to obtain the single cell performance of < 1,85 V per cell at 1 A cm⁻² and outstanding durability, also including studies focused to understand the interaction between catalyst and ionomer, as well as the electrode-membrane interface;
- 4. To design and integrate the newly developed components in a 100 cm² active area, 10 cell, 2 kW stack platform, with cell voltages < 1,85 V per cell at 1 A cm⁻² below 50°C, using diluted electrolytes (≤ 1 M KOH) and an operating differential pressure of 30 bar.
- 5. To develop a low-cost electrolyser unit with a CAPEX equal to or below current classical alkaline electrolyser.







As part of the CHANNEL consortium, Evonik is working on the best-in-class polymer & membrane

	Evonik be class pol M3, D	est-in- lymer 22.1 testing best-in-class feedback loops GEN1 Dev		testing GEN1 feedback loops GEN2 Dev	÷		testing GEN2 feedback loops GEN3 Dev		2 kW AEM demonstrator
	MO	M10, D2.2			M1	8, D2.3		M24, D2.4	
		KPI				UNIT	OBJECTIVE FCHJU 2.4-2019	OBJECTIVE CHANNEL	
			Are	ea specifc resitance	ASR, T = RT	Ωcm^2	< 0,07	< 0,06	
				OH conduct	ivity, T = RT	mS/cm	50	> 50	
				OH conductivi	ty, T = 60°C	mS/cm	not specified	> 90	
		Ex-situ stab	oility (AST proto	col, 1 M KOH, T = 60) °C, 600 hr)	mS/cm	not specified	> 80	
				hydrogen crossove	er (T = 60°C)	[mol/m.s.Pa]x10 ⁻¹⁵	not specified	< 15	
and the second se				water up	take, T = RT	w-%	not specified	< 10	
And		Dry/wet swelling machine Direction (MD)				%	< 1	< 1	
		Dry/wet swelling traverse Direction (TD)				%	< 4	< 4	
		Mechanical strength (in dry conditions, T = RT, RH = 50%)				MPa	15	15	
	1	Elongation at break (in dry conditions, T = RT, RH = 50%)				%	100	100	
		Mechanical strength (DMTA, in fully hydrated, swollen conditions, T = 30°C)				MPa	not specified	> 0,1	
		Mechanical strength (DMTA, i	in fully hydrated	l, swollen condition	s, T = 60°C)	MPa	not specified	> 0,1	
			lone	omer OH conductivi	ty, T = 60°C	mS/cm	20	> 60	
	-0.			In-situ stability A	SR remains	h	2000	> 5000	







Evonik AEM outperforms PEM benchmark Nafion N-115



- Evonik AEM outperforms in 1M KOH electrolyte benchmark PEM membrane Nafion N-115 in DI-H₂O @ 60°C by 0.186 V @ 2 A/cm² (lab scale, single cell, 25 cm² active area)
- Application of Evonik AEM in 1M KOH electrolyte @ 60°C
 & @ 2 A/cm² can enable reduction of operational costs up to 9.5% in comparison to PEM water electrolysis





OER Development Electrochemical performance

Ex situ testing- RDE

• OER-CHANNEL catalyst present an

overpotential < 300 mV at 10 mA/cm²_{geo}



• OER-CHANNEL catalyst is

stable for more than 500

hours at 1.6 V – degradation

of less than 20 mV

 OER-CHANNEL catalyst showed higher

performance

compared to Ni-based

catalysts presented in

the literature.







OER catalysts development Electrochemical performance

In situ testing- single cell performance

- OER- CHANNEL catalyst loading: 5mg cm⁻²
- OER-CHANNEL catalyst present cell voltage of < 1.8 V at 1 A cm⁻²
- 500h cell testing present stable and robust catalyst and electrode structure with insignificant degradation rates







HER catalyst development

- HER-CHANNEL catalyst achieves <150 mV overpotential at 10 mA cm⁻²_{geo} in <1 MKOH.
- The catalyst shows only 26 μ V/hr degradation rate.
- HER catalysts achieve performance comparable to Pt/C in alkaline







CHANNEL Full non-PGM electrolyzer

Performance of 1 A cm⁻² at 1.9 V with good stability.



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Development of 1-D transient AEM electrolyzer model

- Unique challenges for AEM electrolysis include:
 - Parallel ion conduction pathways
 - Liquid electrolyte (I, VI)
 - Solid ionomer (II, V)
 - Ion exchange (III)
 - Water absorption, desorption, and transport in AEM (IV)
 - Evaporation and condensation (VII)
- Coupled equations are solved using custom MATLAB scripts









- 100 cm² active area stack
- Cell voltages of <1.85 V per cell at 1 A cm⁻² below 50°C, using diluted electrolytes (≤ 1 M KOH)
- Operating differential pressure 30 bar
- Maintain stable performance for 2,000 h with a degradation gap of less than 50 mV

In addition, all electronic conductive components, PTLs, foams, mesh and BPPs, must maintain low ICR

• < 20 $m\Omega$ cm² after ex-situ AST test in KOH and validated in 25 cm² AEM single cell













Bipolar plates / PTLs / Flow field

- Bipolar materials
 - Ti
 - SS AISI 304L
 - Inconel 625
 - SS AISI 316L
 - Nickel
- PTL materials
 - SS AISI 316L
 - Nickel
 - Titanium







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Solution

H2



Summary

- OER and HER CHANNEL catalysts exhibit excellent performance in *ex situ* and in *in situ* electrochemical testing
- Production of both electrocatalysts is cost efficient and simple
- Development of 1-D transient model to predict durability of catalyst layers over time, and probing local effects
- Preliminary stack design concluded and validation of PTLs and Bipolar plate materials, including sealings
- Construct prelimanary stack underway



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Future work

- Further optimisation of membranes and ionomers to reach the KPIs
- Further optimization of non-PGM electrocatalysts and catalyst layers to achieve the performance of < 1,85 V per cell at 1 A cm⁻², while testing with non-PGMs.
- Scaling-up of synthesis batches to larger quantities without compromising electrocatalytic performance
- Understanding catalyst and ionomer interaction to reach better chemical compatibilities
- Testing of the preliminary short stack before end of 2021, and finalising the final 2kW stack design
- Develop a beyond the state-of-the-art AEM electrolyser system including power supply, system control, gas drying unit achieving:
 - ▶ An electrolyser cost $< 600 \notin W$ at 500 kW system level
 - An energy consumption < 4.7 kWh/Nm³ at a system level
 - A 100% EU supply chain and increased EU competitiveness in production of green hydrogen from renewable energy sources.
 - Explore the upscale and commercialisation of the newly developed technology.







Acknowledgements



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