



**HydroGEN**  
Advanced Water Splitting Materials

# Advanced Water-Splitting Technology Pathways Benchmarking & Protocols Workshop

## **Breakout Session Supplemental Slides** *Low Temperature Electrolysis (LTE)*

**March 2 – 3, 2021**

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# LTE Breakout Sessions

Session ID	Topic	Lead
LTE-1	LTE Technology Roadmap Review & Discussion- Catalysts	Shannon Boettcher (Univ of Oregon)
LTE-2	Technology Roadmap Review & Discussion- Porous Transport Layer (PTL) Tech	Nemanja Danilovic (LBNL)
LTE-3	Techno-Economic Analysis - LTE	Brian James (Strategic Analysis, Inc)
LTE-5	LTE Cell Test Methods & Reference Cell	Marcelo Carmo (Juelich)
LTE-7	Technology Roadmap Review & Discussion - Membranes	Andrew Motz (Nel Hydrogen)



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## Techno-Economic Analysis Breakout Session Low Temperature Electrolysis (LTE)

Session ID: LTE-3

Session Chair: Brian James

Affiliation: Strategic Analysis Inc.

Date: March 2, 2021

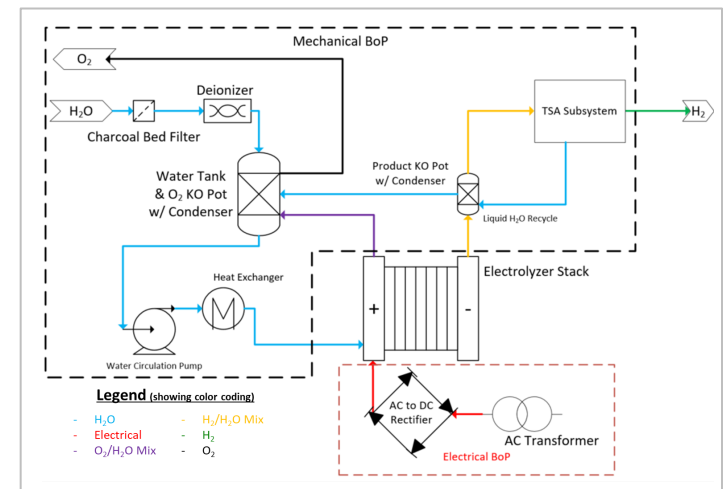
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# Overview of Techno-Economic Analysis Methodology

- The objective of techno-economic analysis (TEA) is to **evaluate** and **compare** competing technologies and **chart progress** on the basis of cost and technical performance
- **TEA Method Steps**
  - Define system: develop flow schematic and bill of materials
  - Perform system mass & heat balance modeling to identify critical design parameters
  - Enumerate H<sub>2</sub> production plant capital cost
  - Investigate and input technical and financial values into discounted cash flow analysis model H2A to evaluate the levelized cost of hydrogen (\$/kgH<sub>2</sub>)
- **Results and Post-Analysis**
  - Perform sensitivity analyses to identify components with greatest impact on cost
    - Tornados and Monte Carlo
  - Obtain external review and feedback
  - Use feedback to update models





## LTE Case Studies

- Alkaline is industry electrolysis standard
- PEM (2019) – Results and assumptions included in [DOE Record](#)
  - Distributed: 1,500 kgH<sub>2</sub>/day
  - Central: 50 TPD
  - Two time-frames, both at ~600MW/year
    - “Projected Current” SOA in 2019 (Not “Existing” or “Commercial”!)
    - “Projected Future” SOA in 2035
- AEM (2020) – Preliminary analysis in 2020, however, there has been significant progress in performance and durability in the last year

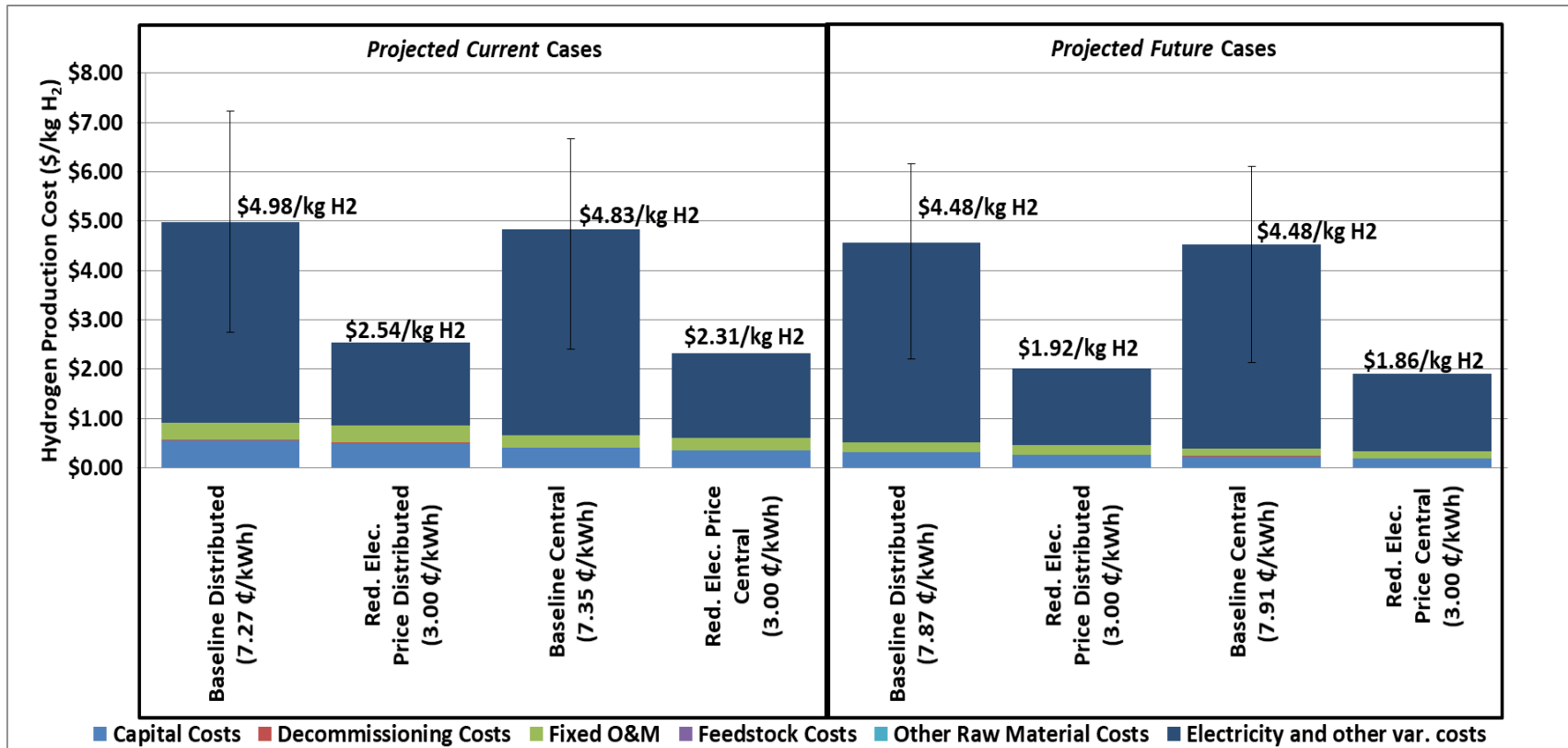


# Hierarchy of LTE Cost Drivers

- **Electricity Cost**
  - Electricity price (cents/kWh)
    - Also (possibly by) time-of-day, interruptible, etc.
  - System Electrical Efficiency
    - Primarily Cell Operating Voltage
      - Operating choice influenced by current density (and stack cost)
      - **Main focus has been to improve Polarization Curve**
- **Stack Cost**
  - \$/cm<sup>2</sup>
    - Material/Manufacturing Costs
      - Catalyst/Loading, Plate Material/Coating, PTL, MEA
      - **Main focus has been to reduce materials/manufac. costs**
  - Current Density (at cell voltage, see above)
- **Stack Durability**
  - Stack degradation rate, mW/1,000 h
    - Desire 90kh to match Alkaline stacks, PEM currently modeled as 1.5mW/1kh
    - **Efforts to improve lifetime**
- **All other cost elements are down in the weeds**



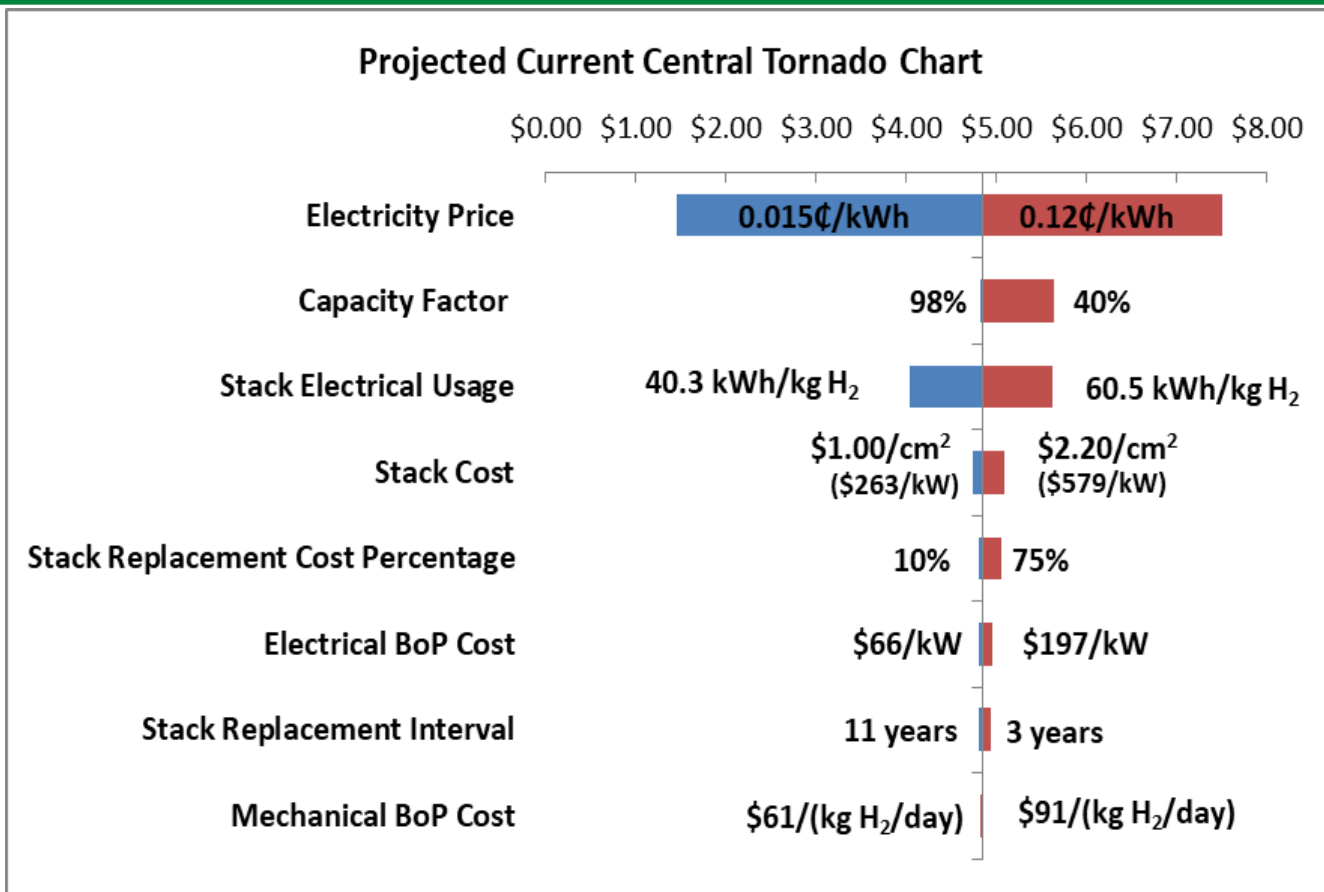
# PEM Electrolyzer Results and Sensitivity Study



- Electricity price most impactful but have least control over from stack technology point of view
- Stack electrical efficiency (kWh/kg) is a key parameter design feature



# PEM Electrolyzer Results and Sensitivity Study



Central	Existing	Proj. Current	Proj. Future	HydroGEN
Stack	-	\$342/kW	\$143/kW	\$100/kW
BOP	-	\$118/kW	\$91/kW	
Total	~\$1,500/kW	\$460/kW	\$233/kW	





# PEM Electrolyzer TEA Model Assumptions

Parameter	Current Distributed 1,500 kg/day	Future Distributed 1,500 kg/day	Current Central 50,000 kg/day	Future Central 50,000 kg/day
Technology Year	2019	2035	2019	2035
Start-up Year	2015	2040	2015	2040
Total Uninstalled Capital (2016\$/kW)	\$599	\$379	\$460	\$233
Stack Capital Cost (2016\$/kW)	\$342	\$143	\$342	\$143
BoP CapEx (2016\$/kW)	\$257	\$236	\$118	\$91
Mechanical BoP Cost (2016\$/kW)	\$136	\$140	\$36	\$23
Electrical BoP Cost (2016\$/kW)	\$121	\$97	\$82	\$68
Total Electrical Usage (kWh/kg)	<b>55.8</b>	<b>51.4</b>	<b>55.5</b>	<b>51.3</b>
[% LHV] (% HHV)	<b>[59.7%] (70.6%)</b>	<b>[64.8%] (76.6%)</b>	<b>[60.1%] (71.0%)</b>	<b>[65.0%] (76.8%)</b>
Stack Electrical Usage (kWh/kg)	50.4	47.8	50.4	47.8
[% LHV] (% HHV)	[66.1%] (78.2%)	[69.8%] (82.4%)	[66.1%] (78.2%)	[69.8%] (82.4%)
BoP Electrical Usage (kWh/kg)	5.4	3.66	5.04	3.54
Stack Current Density (A/cm <sup>2</sup> )	<b>2.0</b>	<b>3.0</b>	<b>2.0</b>	<b>3.0</b>
Cell Voltage (V)	<b>1.9</b>	<b>1.8</b>	<b>1.9</b>	<b>1.8</b>
Electrolyzer Power Consumption at Peak Production (MW)	3.56	3.53	119	118
Effective Electricity Price over Life of Plant (2016¢/kWh)	7.27	7.87	7.35	7.91
Outlet Pressure from Electrolyzer (psi)	300	700	300	700
Installation Cost (% of uninstalled capital cost)	12%	10%	12%	10%
Stack Replacement Interval (years)	7	10	7	10
Stack Replacement Cost Percentage (% of installed capital cost)	15%	15%	15%	15%
Plant Life (years)	20	20	40	40
Stack Degradation Rate (mV/khrs)	1.5	1	1.5	1
Cell Active Area (cm <sup>2</sup> )	700	700	1,500	1,500
Capacity Factor (%)	97%	97%	97%	97%



# Common Stack Materials

	Alkaline	PEM	AEM
Electrolyte	30 wt% KOH	Polyfluorosulfonic acid (PFSA) membrane	$N_4^+/P_4^+$ membrane May have dilute KOH on O.E.
Separator	Porous polyphenylene sulfide w/ ZrO <sub>2</sub> & polymer coatings (Zirfon)	--	
Hydrogen Electrode	Porous nickel or nickel-coated stainless steel	Platinum on carbon	Platinum on carbon (non-PGM in future)
Oxygen Electrode	Porous nickel or nickel-coated stainless steel	Iridium oxide (in Pt/Ru alloys)	Non-PGM metal alloys (Fe/Ni common)
Transport Layers	Nickel mesh	Porous, coated Ti (HE) Graphite (OE)	Ni foam (OE) Carbon GDL (HE)
Bipolar Plates	Nickel-coated stainless steel	Pt-Coated titanium	Stainless Steel
Frames/Sealing	Polymer	Polymer	Polymer



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# Advanced Water-Splitting Technology Pathways Benchmarking & Protocols Workshop

## Technology Roadmap Review & Discussion - Membranes Technology: LTE

**Session ID: LTE-7**

**Session Chair: Andrew Motz**

**Affiliation: Nel Hydrogen**

**Date: March 3, 2021**

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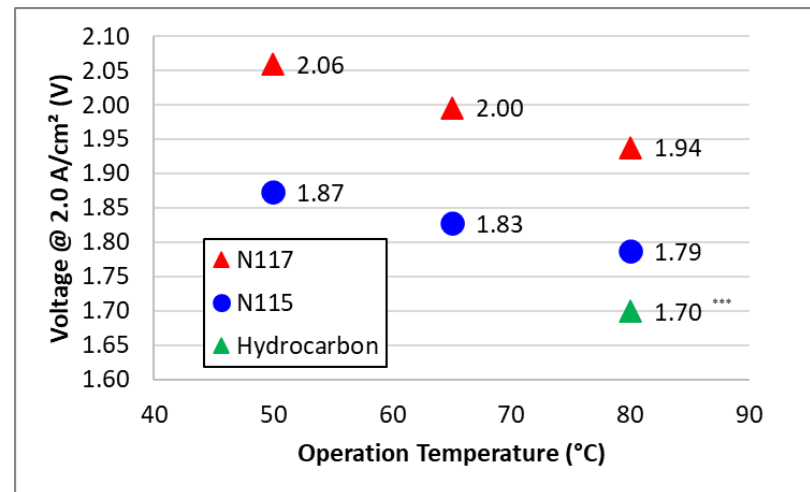


# PEM - Temperature and Membrane Thickness

- Starting with a base case of N117 at 50° C
  - Increasing temperature to 80° C can result in an efficiency improvement of about 120 mV
  - Alternatively moving to N115 membrane can result in an efficiency improvement of about 187 mV
  - With thinner membranes, there are diminishing returns on increasing operating temperature
  - Alternative proton exchange membrane chemistries have received minimal investigation

**How much further can we reduce operating potential with membranes alone (50, 100, 150 mV)?**

**Where should the community focus efforts? Membrane thickness, operating temperature, ion transport?**



\*\*\*Hydrocarbon data from reference and had poor durability (C. Klose et. al. 2020 Adv. Energy Mater. 10 pg. 1903995)

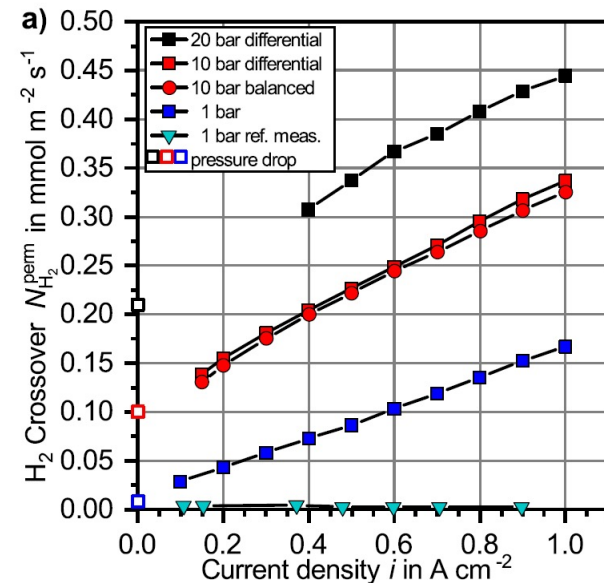


# PEM – Thin membrane challenges

- Gas crossover at differential pressure will be greater based on Fickian diffusion
- To add another challenge for thin membranes, there have been reports that at higher current densities and a fixed  $H_2$  pressure, the flux of  $H_2$  across the membrane increases.

**What is the mechanism for this phenomena?**

**Do we need to account for these losses in the reported efficiency?**





# PEM – Failure Modes

- Chemical Degradation

- There is some evidence of membrane thinning (S. Grigoriev et al 2014 Int. J. Hydrog. Energy 39 pg. 20440)– 90°C / voltage ~2.3 V / cycling / was still stable for 4000 h before exponential failure. **Is this more extreme than we would expect in the field over 80,000 h?**
- Fluoride release rate has also been reported (M. Chandesris et al 2015 Int. J. Hydrog. Energy 40 pg. 1353) – Water fed anode had minimal fluoride release, cathode water (flux across the membrane) had higher fluoride levels detected – **How accurate is measuring the water on the cathode? Rate varies with current density and there will be loss of water vapor in the H<sub>2</sub> product.**
- How does metal dissolution into the membrane impact degradation?
- **Are there better approaches to accelerate chemical degradation?**



## PEM – Failure Modes

- Chemical Degradation - Are there better approaches to accelerate chemical degradation?
- Mechanical Degradation
  - Electronic shorting is a common failure mechanism for many low resistance membranes (i.e. low EW or sub 127  $\mu\text{m}$ )
  - Fully hydrated mechanical properties are rarely reported
  - Is there a valuable ex-situ metric that can come from a fully hydrated tensile or burst strength to avoid failures due to electronic shorting?



## PEM – Failure Modes

- Chemical Degradation - Are there better approaches to accelerate chemical degradation?
- Mechanical Degradation - Is there a valuable ex-situ metric that can come from a fully hydrated tensile or burst strength to avoid failures due to electronic shorting?
- Thermal Degradation
  - Increased temperature can accelerate chemical and mechanical degradation through more facile kinetics or softening of the polymer
  - Delamination through freeze thaw cycles or typical on/off cycling is also possible – What level of impact / importance does this have on actual devices?





## PEM – Summary

- Chemical Degradation – Are there better approaches to accelerate chemical degradation?
- Mechanical Degradation – Is there a valuable ex-situ metric that can come from a fully hydrated tensile or burst strength to avoid failures due to electronic shorting?
- Thermal Degradation – What level of impact / importance does this have on actual devices?
- Thinner membranes and higher operation temperatures are critical to achieving the roadmap efficiency targets – **how do we best advance the field without sacrificing reliability?**



# AEM – New Materials

- In the last few years, numerous AEMs and ionomers have become commercially available



More options is great for improving chances for success, but makes standardization difficult



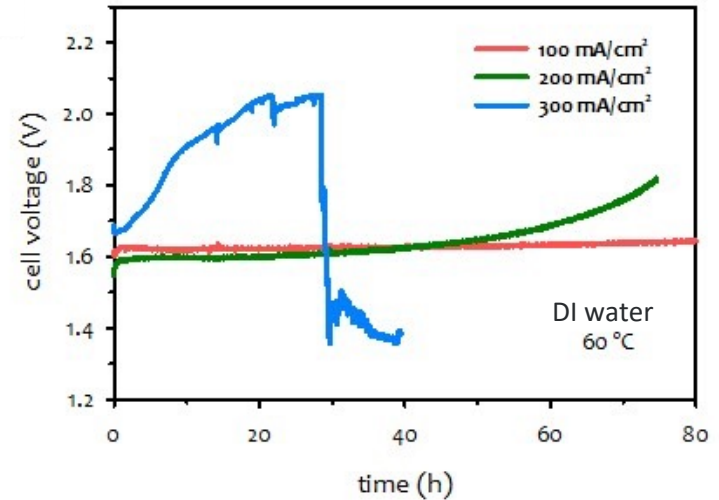
# AEM – Cell Operation / Degradation

- Mode of operation

- Should we run these cells in DI water,  $K_2CO_3$ , KOH? Should there be a standard concentration?

- Degradation

- Most AEMWE degradation rates are over an order of magnitude higher than comparable PEMWE – is it feasible to close this gap? How soon? Is it a requirement for the technology?
- Mechanical degradation – AEMs used in literature are often much thinner than standard PEMs . Is the answer to simply use thicker AEMs? When does water flux become a challenge?
- Chemical degradation – Is the able to be overcome to achieve
- Thermal Degradation – is high temperature required to take advantage of non-PGM catalysts?



D. Li, et al, Nature Energy, 2020, 5, 378-385