







# Electrochemical Mitigation of Corrosion in Molten Chloride Salts During CSP Plant Operation

Dr. Kerry Rippy ES2021, Virtual, Online June 16-18, 2021

#### Molten Chloride Salt

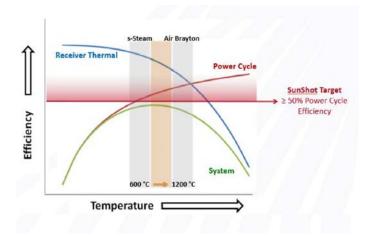
# Molten chloride salts are stable at higher temperatures than other molten salts

- Could enable more efficient operation of CSP plants, leading to lower LCOE
- Also useful for other reactors types (nuclear, electrochemical synthesis, etc.)

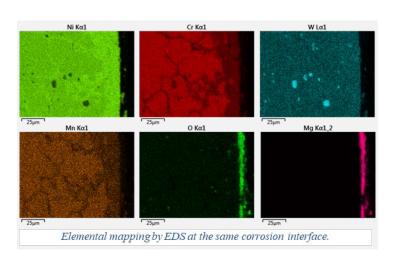
# For this work, we are using a ternary NaCl/KCl/MgCl<sub>2</sub> salt

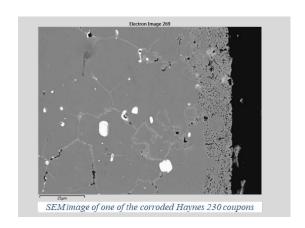
- Has favorable properties including low melting point
- Can be inexpensively sourced



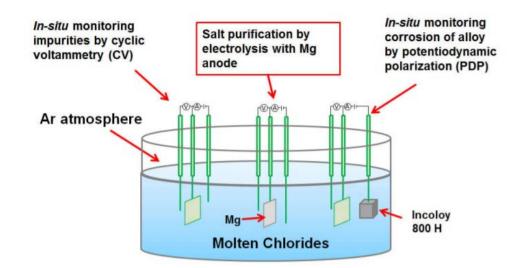


**Problem:** Molten chloride salts are corrosive! The corrosive impurity MgOHCl forms in salt when exposed to H<sub>2</sub>O.





Alloy	OCP vs. pseudo-RE $[mV]$	$E_{corr}$ vs. pseudo-RE $[mV]$	$j_{\rm corr}[\mu A/cm^2]$	CR [mm/year]	
650 °C					
SS347	-784±113	-847±91	713.00±30.74	7.49±0.32	
SS310	-888±170	-938±218	626.16±38.72	6.42±0.40	
In800H	-876±32	-910±22	573.79±34.19	5.94±0.33	
IN625	-849±73	-856±57	233.00±94.68	2.80±0.38	
700 °C					
SS310	-506±70	-571±75	1213.61±148.10	12.45±1.52	
In800H	-453±5	-474±3	1387.79±131.96	14.31±1.36	
SS310					



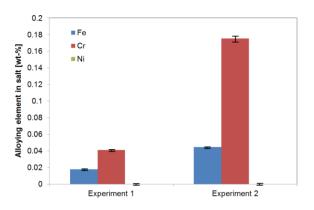


Figure 13: Alloying elements Cr. Fe and Ni dissolved in the chloride salts after 450 h (Experiment 1, purified salt) and 340 h (Experiment 2, unpurified salt) exposure, measured with AAS. Ni Content is below the detection limit of AAS (10 ppm in weight). The error bar represents the standard deviation of three measurements

Ding, W.; et. Al. Solar Energy Materials and Solar Cells 2019.

## MgOHCl can be removed using electrolysis!

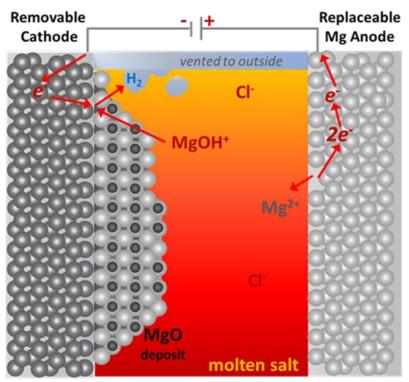
$2MgOHCl + Mg(s) \rightarrow MgCl_2 + 2MgO + H_2(g)$	) [overall reaction]
$Mg(s) \rightarrow Mg^{2+} + 2e^-$	_[anode]
$2MgOH^+ + 2e^-$ → $2MgO + H_2(g)$	[cathode]
$Mg^{2+} + 2CI^- \rightarrow MgCI_2$	[molten salt]
2MgOHCl → 2MgOH+ + 2Cl-	[molten salt]

### Goal: Design a purification cell for a Gen3 CSP Plant



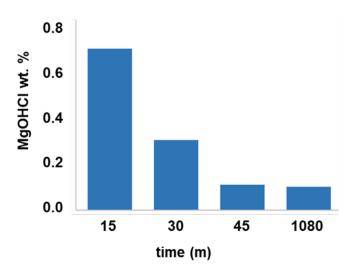
#### overall reaction:

$$MgOH^+ + e^- \rightarrow MgO_{(s)} + \frac{1}{2}H_{2(g)}$$



Lattice strain minimized at interface





#### **Lab Scale Results**

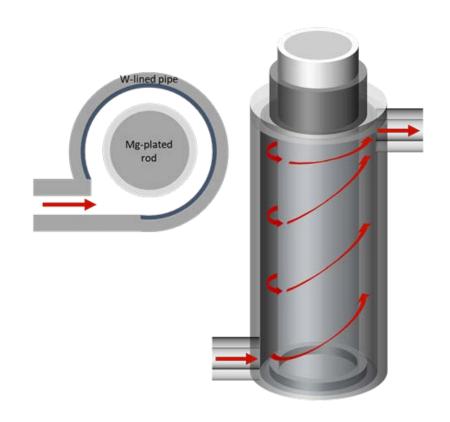
- **Titration**
- H<sub>2</sub> detection

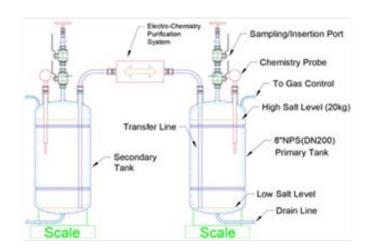
In a lab-scale batch reactor, MgOHCl can be reduced to target 0.1 wt% in about 45 min.

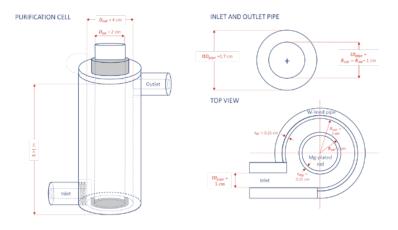
### **Challenges:**

- Mass Transfer
- Passivation of electrode due to MgO buildup

We believe these challenges will be mitigated in a flow cell







### **Outlook and next steps:**

- Building a lab-scale flow system and flow purification cell
- Evaluate and optimize kinetics for *in-situ* control of corrosive impurities during operation of molten-chloride salt based CSP plants
- Maintenence cost, materials costs, and ultimately LCOE of CSP could be reduced

## Thank you for your attention!

#### **Team Members:**

Dr. Judith Vidal, NREL

Dr. Patrick Taylor, CSM

Dr. Mark Anderson, UW

Liam Witteman, CSM

#### NREL/PR-5500-80080

This work was authored [in part] by the National Renewable Energy Laboratory, operated by Alliance for Sustainable Energy, LLC, for the U.S. Department of Energy (DOE) under Contract No. DE-AC36-08GO28308. Funding provided by the U.S. Department of Energy Office of Energy Efficiency and Renewable Energy Solar Energy Technologies Office. The views expressed in the article do not necessarily represent the views of the DOE or the U.S. Government. The U.S. Government retains and the publisher, by accepting the article for publication, acknowledges that the U.S. Government retains a nonexclusive, paid-up, irrevocable, worldwide license to publish or reproduce the published form of this work, or allow others to do so, for U.S. Government purposes.