Advanced Water Purification System for *In Situ* Resource Utilization

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• One of NASA’s goals is to enable long-term human presence in space, without the need for continuous replenishment of consumables from Earth.

• *In situ* resource utilization (ISRU) is the use of extraterrestrial resources to support activities such as human life-support, material fabrication and repair, and radiation shielding.

• Potential sources of ISRU resources include lunar and Martian regolith, and Martian atmosphere.

*Crops Mars ISRU Demonstration Unit*
• Water and byproducts (including hydrochloric and hydrofluoric acids) can be produced from lunar regolith via a high-temperature hydrogen reduction reaction and passing the produced gas through a condenser.

\[ \text{FeO} + \text{H}_2 \xrightarrow{900^\circ C} \text{Fe} + \text{H}_2\text{O} \]

• Due to the high solubility of HCl and HF in water, these byproducts are expected to be present in the product stream (up to 20,000 ppm) and must be removed (less than 10 ppm) prior to water consumption or electrolysis.
Due to their consumable nature, typical water purification methods may not be suitable for HCl and HF removal in extraterrestrial applications. Membranes and adsorbents are often regenerated with large amounts of water and/or basic solutions, which aren’t available in the lunar environment. Naturally-occurring adsorbents may be replaced rather than regenerated.

<table>
<thead>
<tr>
<th>Method</th>
<th>Examples</th>
</tr>
</thead>
<tbody>
<tr>
<td>Membrane (solute retention)</td>
<td>Reverse osmosis</td>
</tr>
<tr>
<td></td>
<td>Nanofiltration</td>
</tr>
<tr>
<td></td>
<td>Proton Exchange</td>
</tr>
<tr>
<td>Membrane (solute transport)</td>
<td>Dialysis</td>
</tr>
<tr>
<td></td>
<td>Electrodialysis</td>
</tr>
<tr>
<td></td>
<td>Anion Exchange</td>
</tr>
<tr>
<td>Adsorption</td>
<td>Alumina-based adsorbents</td>
</tr>
<tr>
<td></td>
<td>Natural adsorbents (mud, ore, clay, soil, chitosan)</td>
</tr>
</tbody>
</table>
Nafion®, a common fuel cell proton exchange membrane, was investigated for its chloride and fluoride rejection capability in a previous study.

- The ability to reject 98-99.9% of chloride and 50-80% of fluoride was demonstrated.
- This rate of rejection was not sufficient to produce electrolysis-grade water.
- Since water had to diffuse across the membrane and be recovered in the vapor phase, relatively small amounts of water were recovered using large amounts of carrier gas.

Other technologies were researched with an emphasis on maximizing water recovery and contaminant removal:

- Higher contaminant removal/rejection rate
- Contaminant removal/water retention instead of contaminant rejection/water transport
- Liquid phase only process

Electrodialysis for Contaminant Removal
High Efficiency Electrodialysis (HEED®) membrane stack purchased from EET Corp.
Stack contains 20 alternating Ralex® anion and cation exchange membranes
0.5% NaSO₄ solution circulated through outer cells to protect electrodes
Baseline Contaminant Removal: 20,000 ppm HCl/HF Solution

- Starting volume
  - 500 mL diluent
  - 500 mL concentrate
- Final volume
  - 295 mL diluent
  - 685 mL concentrate
- Power supply
  - 30 V maximum
  - 3 A maximum
- Pump speed
  - 100% diluent
  - 100% concentrate
- Reached steady state within 30 min

<table>
<thead>
<tr>
<th>Diluent Stream</th>
<th>Fluoride</th>
<th>Chloride</th>
</tr>
</thead>
<tbody>
<tr>
<td>Initial (ppm)</td>
<td>18,800</td>
<td>26,400</td>
</tr>
<tr>
<td>Final (ppm)</td>
<td>71.1</td>
<td>11.4</td>
</tr>
<tr>
<td>Removal (%)</td>
<td>99.44</td>
<td>99.94</td>
</tr>
</tbody>
</table>
Baseline Contaminant Removal: 200 ppm HCl/HF Solution

- Starting volume
  - 500 mL diluent
  - 500 mL concentrate
- Final volume
  - 500 mL diluent
  - 500 mL concentrate
- Power supply
  - 30 V maximum
  - 3 A maximum
- Pump speed
  - 100% diluent
  - 100% concentrate
- Reached steady state in less than 10 min

<table>
<thead>
<tr>
<th>Diluent Stream</th>
<th>Fluoride (ppm)</th>
<th>Chloride (ppm)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Initial</td>
<td>166</td>
<td>179</td>
</tr>
<tr>
<td>Final</td>
<td>0.26</td>
<td>0.07</td>
</tr>
<tr>
<td>Removal (%)</td>
<td>99.85</td>
<td>99.96</td>
</tr>
</tbody>
</table>
Variable Matrix Testing: Concentrate Volume

<table>
<thead>
<tr>
<th>Initial Concentrate Volume (mL)</th>
<th>Final Diluent Volume (mL)</th>
<th>Final Concentrate Volume (mL)</th>
<th>Final Fluoride Concentration (ppm)</th>
<th>Final Chloride Concentration (ppm)</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>170</td>
<td>290</td>
<td>550</td>
<td>23</td>
</tr>
<tr>
<td>130</td>
<td>305</td>
<td>310</td>
<td>290</td>
<td>25</td>
</tr>
<tr>
<td>250</td>
<td>335</td>
<td>400</td>
<td>167</td>
<td>22</td>
</tr>
<tr>
<td>500</td>
<td>295</td>
<td>685</td>
<td>71</td>
<td>11</td>
</tr>
</tbody>
</table>

- Parameters:
  - Initial diluent volume: 500 mL
  - Initial diluent concentration: 20,000 ppm HCl/HF
  - Pump speed: 100% of maximum
  - Voltage: 30 V maximum
  - Current: 3 A maximum
- 130 mL is minimum volume required to fill the concentrate membrane cells and tubing
- Final fluoride concentration decreases substantially with decreasing initial concentrate volume
- Final chloride concentration does not decrease with increasing initial concentrate volume below 250 mL
### Variable Matrix Testing: Pump Speed

<table>
<thead>
<tr>
<th>Pump Speed (% of maximum)</th>
<th>Final Diluent Volume (mL)</th>
<th>Final Concentrate Volume (mL)</th>
<th>Final Fluoride Concentration (ppm)</th>
<th>Final Chloride Concentration (ppm)</th>
</tr>
</thead>
<tbody>
<tr>
<td>&lt; 10</td>
<td>500</td>
<td>510</td>
<td>14,500</td>
<td>13,700</td>
</tr>
<tr>
<td>20</td>
<td>280</td>
<td>650</td>
<td>71</td>
<td>8.6</td>
</tr>
<tr>
<td>50</td>
<td>300</td>
<td>700</td>
<td>158</td>
<td>20</td>
</tr>
<tr>
<td>100</td>
<td>295</td>
<td>685</td>
<td>71</td>
<td>11</td>
</tr>
</tbody>
</table>

- **Parameters:**
  - Initial diluent volume: 500 mL
  - Initial diluent concentration: 20,000 ppm HCl/HF
  - Initial concentrate volume: 500 mL
  - Voltage: 30 V maximum
  - Current: 3 A maximum

- Pump speed does not affect final fluoride and chloride concentrations, or final diluent and concentrate volumes at or above 20% of maximum setting (at least 200 mL/min)
- Minimum pump speed does not appear to allow for any ion exchange
## Variable Matrix Testing: Power Supply

<table>
<thead>
<tr>
<th>Maximum Voltage (V)</th>
<th>Maximum Current (A)</th>
<th>Final Diluent Volume (mL)</th>
<th>Final Concentrate Volume (mL)</th>
<th>Final Fluoride Concentration (ppm)</th>
<th>Final Chloride Concentration (ppm)</th>
</tr>
</thead>
<tbody>
<tr>
<td>30</td>
<td>3.0</td>
<td>296</td>
<td>685</td>
<td>71</td>
<td>11</td>
</tr>
<tr>
<td>15</td>
<td>3.0</td>
<td>325</td>
<td>675</td>
<td>1,170</td>
<td>47</td>
</tr>
<tr>
<td>60</td>
<td>1.5</td>
<td>350</td>
<td>640</td>
<td>26</td>
<td>2.8</td>
</tr>
<tr>
<td>30</td>
<td>6.0</td>
<td>355</td>
<td>610</td>
<td>132</td>
<td>26</td>
</tr>
</tbody>
</table>

- **Parameters:**
  - Initial diluent volume: 500 mL
  - Initial diluent concentration: 20,000 ppm HCl/HF
  - Initial concentrate volume: 500 mL
  - Pump speed: 100% of maximum

- Decreasing voltage or current increases final diluent fluoride and chloride concentrations
- Increasing voltage decreases final diluent chloride and fluoride concentrations
- Increasing current does not affect final diluent chloride and fluoride concentrations
• Initial volumes:
  - 500 mL diluent
  - 100 mL concentrate (replaced every hour, 300 mL total)
• Power supply:
  - 30 V maximum
  - 3.0 A maximum
• Pump speed: 100%
• Final volumes:
  - 250 mL diluent
  - 600 mL concentrate
• Contaminant removal improved versus baseline, but at the expense of longer processing time

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<th>Fluoride (ppm)</th>
<th>Chloride (ppm)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Initial</td>
<td>17,000</td>
<td>17,400</td>
</tr>
<tr>
<td>Final</td>
<td>27</td>
<td>3.9</td>
</tr>
<tr>
<td>Removal (%)</td>
<td>99.92</td>
<td>99.99</td>
</tr>
</tbody>
</table>
• Initial volumes:
  - 250 mL diluent (replaced every hour, 1250 mL total)
  - 250 mL concentrate
• Power supply:
  - 30 V maximum
  - 3.0 A maximum
• Pump speed: 100%
• Final volumes:
  - 360 mL diluent
  - 830 mL concentrate
• Contaminant removal was not significantly improved versus simply processing at a higher diluent to concentrate volume ratio
### ISRU Scenario Testing: Obtaining Electrolysis-Grade Water

**1000 mL HCl/HF Solution**
- 390 mL Clean Water
  - **Diluent “A”**
    - 500 mL
    - 130 mL
  - **Concentrate “A”**
    - 130 mL
    - 330 mL
  - **Diluent “B”**
    - 300 mL
    - 130 mL
  - **Concentrate “B”**
    - 130 mL
    - 330 mL
  - **Diluent “C”**
    - 300 mL
    - 130 mL
    - 330 mL
  - **Concentrate “C”**
    - 130 mL
    - 330 mL
  - **Diluent “D”**
    - 435 mL
    - 565 mL
    - 130 mL
    - 330 mL
  - **Concentrate “D”**
    - 130 mL

**Diluent “A”**
- **Fluoride**
  - Initial (ppm): 14,200
  - Final (ppm): 736
  - Removal (%): 94.8
- **Chloride**
  - Initial (ppm): 15,100
  - Final (ppm): 124
  - Removal (%): 99.2

**Diluent “B”**
- **Fluoride**
  - Initial (ppm): 16,200
  - Final (ppm): 214
  - Removal (%): 98.7
- **Chloride**
  - Initial (ppm): 19,000
  - Final (ppm): 126
  - Removal (%): 99.3

**Diluent “C”**
- **Fluoride**
  - Initial (ppm): 1,140
  - Final (ppm): 12.6
  - Removal (%): 98.9
- **Chloride**
  - Initial (ppm): 222
  - Final (ppm): 10.9
  - Removal (%): 95.1

**Diluent “D”**
- **Fluoride**
  - Initial (ppm): 80.4
  - Final (ppm): 0.41
  - Removal (%): 99.5
- **Chloride**
  - Initial (ppm): 41.4
  - Final (ppm): 0.16
  - Removal (%): 99.6
• Added new and additional components to increase system robustness and minimize processing time:
  – Larger membrane stack that contains two 20 membrane cells which can run in series or parallel
  – Higher voltage/current power supply to achieve rapid equilibrium
  – In-line conductivity meters for real-time measurement of water purity
  – Hastelloy® electrodes which offer superior protection against hydrofluoric acid versus stainless steel

• Other considerations for future work:
  – Different types of anion/cation exchange membranes
  – Alternate membrane stack configurations
  – Integration with additional water processing unit (e.g., Nafion®)
  – Increase system automation

http://www.eetcorp.com/lts/flowconfig.htm
I,

<table>
<thead>
<tr>
<th>Ion</th>
<th>C (wt%)</th>
<th>C_{th} (wt%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Na⁺</td>
<td>0.27</td>
<td>0.10</td>
</tr>
<tr>
<td>Mg^{2+}</td>
<td>0.44</td>
<td>0.60</td>
</tr>
<tr>
<td>K⁺</td>
<td>0.28</td>
<td>0.03</td>
</tr>
<tr>
<td>Ca^{2+}</td>
<td>0.05</td>
<td>0.04</td>
</tr>
<tr>
<td>Cl⁻</td>
<td>0.79</td>
<td>0.04</td>
</tr>
<tr>
<td>ClO₄⁻</td>
<td>0.72</td>
<td>0.60</td>
</tr>
<tr>
<td>HCO₃⁻</td>
<td>0.73</td>
<td>?</td>
</tr>
<tr>
<td>SO₄^{2-}</td>
<td>0.78</td>
<td>?</td>
</tr>
</tbody>
</table>


- Attempted to simulate ionic contaminants in water recovered by Phoenix lander
- In absence of specific ion standards, measured contaminant removal indirectly via conductivity meters in diluent and concentrate streams
- Achieved minimum diluent conductivity after about 10 min
Collaborators

- Dr. Scott Jolley, QinetiQ North America
- Jim Captain, QinetiQ North America

Funding Sources

- KSC Center Innovation Fund
- KSC Core Technical Capabilities Program
- KSC Independent Research & Technology Development Program
Advanced Water Purification System for In Situ Resource Utilization

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