

#### Electrochemical Conversion of Carbon Dioxide to Oxygen in Ionic Liquid Media

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Presented at the Northeast Regional Meeting of the<br>
American Chemical Society<br>
Burlington, VT<br>
July 1, 2008<br>
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- NASA missions need to extract oxygen from carbon dioxide for advanced life support and human exploration missions **Problem Background<br>
MASA missions need to extract oxygen from carbon dioxide for<br>
dyanced life support and human exploration missions<br>
Martian In-Situ Resource Utilization (ISRU) aims to process<br>
arbon dioxide (95% in at**
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	-
	-
- **Problem Background**<br>
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dvanced life support and human exploration missions<br>
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arbon dioxide (95% in atmosphe • Ionic liquid based carbon dioxide reduction can enable low temperature operation less than 100 °C JASA missions need to extract oxygen from carbon dioxid<br>dvanced life support and human exploration missions<br>Aratian In-Situ Resource Utilization (ISRU) aims to proces<br>arbon dioxide (95% in atmosphere) to oxygen<br>– Sabatier
- Goal is to electrochemically reduce  $CO<sub>2</sub>$  in an electrochemical reactor based on an immobilized ionic liquid in the separator advanced life support and human exploration missions<br>
• Martian In-Situ Resource Utilization (ISRU) aims to process<br>
carbon dioxide (95% in atmosphere) to oxygen<br>
– Sabatier: CO<sub>2</sub> + 4H<sub>2</sub> → CH<sub>4</sub> + 2H<sub>2</sub>O and then H<sub>2</sub>O
	-
	-
- an oxide ion and optimizing the electrochemical reactions



## Electrochemical CO<sub>2</sub> Reactor Approach Electrochemical CO<sub>2</sub> Ream<br>
M&D activities focused on<br>
- Membrane separator<br>
- Ion-exchange: Nafion<br>
- Ionic liquid electrolyte

- R&D activities focused on
	- -
		-
	- -
		-
		-
	- -
	- -
		- activity
	- -





#### Methodology for Screening Ionic Liquid Imbibed Membranes

- Developed a list of ionic liquid anion/cation combinations considering
	- aprotic, protic, zwitterion type ILs, ionic association, size
- Utilize a chemical catalyzation technique to electrolessly deposit platinum catalyst onto Nafion 115 ion-exchange membranes
- Imbibe ionic liquids into platinized Nafion films
- Ionic liquid cation groups
	- Emim: 1-ethyl,3-methylimidazolium
	- Bmim: 1-butyl,3-methylimidazolium
	- BFP: butylmethylpyrrolidinium
- Ionic liquid anion groups
	- BF4: tetrafluoroborate
	- PF6: hexafluorophosphate
	- FMS: trifluoromethanesulfonate
	- TFSI: bis-trifluoromethylsulfonylimide
	- CH3CO2: Acetate
- Evaluate samples using cyclic voltammetry and chronoamperometry



#### Representative Cyclic Voltammogram for  $CO<sub>2</sub>$  Reduction -Slight Increase in Reactivity with Increasing Pressure





#### Comparison of  $CO<sub>2</sub>$  Reduction Over N<sub>2</sub> Baseline Emim-BF4 Gives Highest CO<sub>2</sub> Reduction Rate





#### Summary of CO<sub>2</sub> Reduction in IL Imbibed MEAs

- High  $CO<sub>2</sub>$  reduction rates were observed with MEAs imbibed with ionic liquids having the  $\mathsf{BF}_{4}$  anion, TFMS also gives high performance
- Low  $CO<sub>2</sub>$  reduction rates were observed with MEAs imbibed with TFSI
- MEAs having the ionic liquids with the EMIM cation performed better than those with the BMIM cation
- Results indicate decreasing reactivity with increasing anion size
	- $\bullet$   $\,$  CO $_{2}$  reactivity trend: BF $_{4}$ > TFMS> TFSI
	- Consistent with a cluster network model of the Ionic liquid/Nafion Composite **Structure**
	- Primary charge transfer from counter ions: imidazolium cations
	- Large anions hinder access and decrease IL packing density in pores



• Electrode surface area is also playing an important role in this process. High surface area platinum electrolytically deposited onto the catalyzed MEA gave the highest activity



#### CO2 Reactivity in Ionic Liquids/Platinum Electrodes

- Want to understand what governs the reaction behavior of CO<sub>2</sub> in the  $\quad\rm CO_{2}$  (Gas Phase)  $\rm CO$ ionic liquid system CO<sub>2</sub> Reactivity in Ionic Liquids/Platin<br>
Vant to understand what governs<br>
the reaction behavior of CO<sub>2</sub> in the<br>
polic liquid system<br>
Is it the dissolution of CO<sub>2</sub> in the<br>
ionic liquid, followed by its<br>
diffusion to a r CO<sub>2</sub> Reactivity in Ionic Liquids/PI<br>
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ionic liquid, followed by its<br>
diffusion to a rea
	- Is it the dissolution of  $CO<sub>2</sub>$  in the ionic liquid, followed by its diffusion to a reaction site, or
	- the electrode surface, or
	- of CO $_{\rm 2}$  on the electrode?





**-** Construction of the construction of the construction

#### Experimental Measurements Show Significant Direct CO<sub>2</sub> Gas Phase Reduction on Platinum

- Higher CO<sub>2</sub> Reduction 1.6E-04 Currents Measured When More Platinum Electrode is Exposed to <br>
and <br>  $\sum_{1.0\in 04}$ Gaseous  $CO<sub>2</sub>$ <br>Implications that Higher
- Implications that Higher  $\frac{1}{5}$  8.0E-05 Surface Area Platinum 6.0E-05 Electrodes be Developed on the Outer <sub>20E-05</sub> Surface of the Ionic<br>
interaction of the Ionic Liquid-Membrane  $\frac{0.06+0.0}{25}$   $\frac{1}{25}$   $\frac{1}{50}$   $\frac{1}{25}$   $\frac{1}{50}$



Depth (mils)





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## Micro-Porous Membranes Show Higher Equilibrium Transport Currents Micro-Porous Membranes Show<br>
Transport Curre<br>
upplied 3 V across cell<br>
Measured current over time<br>
Cathode: N<sub>2</sub> or O<sub>2</sub><br>
unode: N<sub>2</sub><br>
Separator<br>
- Nafion 115/emim-BF4<br>
- Celgard/emim-BF4<br>
- Celgard/emim-BF4<br>
- Sale ion t

- Applied 3 V across cell
- Measured current over time
- Cathode:  $N_2$  or  $O_2$
- Anode:  $N_2$
- Separator
	-
	-
- Oxide ion transport current  $\int_{0}^{\infty}$ higher than nitrogen background current
- Interaction of the Nafion pores/ion-exchange groups and ionic liquid lowers the MEA ionic conductivity





#### Equilibrium Current Density for  $CO<sub>2</sub>$  Reduction

- Applied 3 V across cell
- Measured current over time
- Cathode:  $N_2$ ,  $O_2$ , or  $CO_2$  ...
- Anode:  $N_2$
- Separator:
	-
- Oxide ion current (from  $CO<sub>2</sub>$ or  $O_2$  reduction) higher than  $10^{10}$ nitrogen background current
- Faster decay rate for  $CO<sub>2</sub>$ 
	-





## Initial Peak Currents for  $CO<sub>2</sub>$  Reduction Favor Pt Catalyst Over Pt/Ru alloys **Initial Peak Currents for CO<sub>2</sub> F<br>
Catalyst Over Pt/Ru<br>
Applied 3 V across cell<br>
Measured initial current<br>
Cathode: CO<sub>2</sub><br>
- Pt(100%), Pt/Ru(25% molar),<br>
Pt/Ru(50%), Pt/Ru(75%)<br>
Anode: N<sub>2</sub><br>
- Pt (100%) Initial Peak Currents for CO<sub>2</sub> F<br>
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Applied 3 V across cell<br>
Measured initial current<br>
Cathode: CO<sub>2</sub><br>
- Pt(100%), Pt/Ru(25% molar),<br>
Pt/Ru(50%), Pt/Ru(75%)<br>
anode: N<sub>2</sub><br>
- Pt (100%)<br>
Separator:<br>
- Na**

- Applied 3 V across cell
- Measured initial current
- Cathode:  $CO<sub>2</sub>$ 
	- Pt/Ru(50%), Pt/Ru(75%)<br>
	de: N<sub>2</sub>  $\frac{1}{3}$
- Anode:  $N_2$ 
	-
- Separator:
	-
- Highest initial  $CO<sub>2</sub>$  reduction for 100% Pt catalyst on cathode
- Lower activity with Ru addition suggests Pt/Ru composition and catalyst loading could be optimized





### Ruthenium Addition to Platinum Cathode Stabilizes Decay **Ruthenium Addition to Plat<br>
Stabilizes Deca<br>
Applied 3 V across cell<br>
Measured current over time<br>
Cathode:<br>
- CO<sub>2</sub><br>
- Pt(100%), Pt/Ru(25% molar), 08<br>
Pt/Ru(50%), Pt/Ru(75%)<br>
anode: Ruthenium Addition to Plat<br>
Stabilizes Deca<br>
Applied 3 V across cell<br>
Measured current over time<br>
Cathode:<br>
- CO<sub>2</sub><br>
Pt(100%), Pt/Ru(25% molar),**  $\leftarrow \begin{bmatrix} 1 \\ 0.8 \\ 0.9 \\ 0.9 \\ 0.9 \\ 0.9 \\ 0.6 \end{bmatrix}$ **<br>
Pt/Ru(50%), Pt/Ru(75%)<br>**

- Applied 3 V across cell
- Measured current over time
- Cathode:
	-
	- Pt/Ru(50%), Pt/Ru(75%)<br>
	Pt/Ru(50%), Pt/Ru(75%)<br>  $\frac{2}{3}$ <br>
	Pt/Ru(50%), Pt/Ru(75%)<br>  $\frac{2}{3}$ <br>
	Pt/Ru(50%), Pt/Ru(75%)
- Anode:
	-
	-
- Separator:
	-
- Pt/Ru cathode catalysts stabilize performance
- Significant lowering in initial activity suggests Pt/Ru optimization





- $\bullet$  Identified platinum as a preferred catalyst choice for  $\mathrm{CO}_2$ reduction
	-
- **Summary Electrode Development**<br>
dentified platinum as a preferred catalyst choice for CO<sub>2</sub><br>
eduction<br>
 Stable with ionic liquids and potential windows (0 to 3 volts)<br>
 Can be deposited on ion-exchange membranes usin **Summary - Electrode Development<br>
dentified platinum as a preferred catalyst choice for**  $CO_2$ **<br>
eduction<br>
- Stable with ionic liquids and potential windows (0 to 3 volts)<br>
- Can be deposited on ion-exchange membranes using** and electrolytic techniques
- **Summary Electrode Development**<br>
dentified platinum as a preferred catalyst choice for  $CO_2$ <br>
eduction<br>
 Stable with ionic liquids and potential windows (0 to 3 volts)<br>
 Can be deposited on ion-exchange membranes usin the performance **Example 12**<br>
dentified platinum as a preferred catalyst choice for  $CO_2$ <br>
eduction<br>
- Stable with ionic liquids and potential windows (0 to 3 volts)<br>
- Can be deposited on ion-exchange membranes using electroless<br>
and el
- $\bullet$  CO<sub>2</sub> reaction process investigated
	- reaction of  $CO<sub>2</sub>$  on platinum outside the ionic liquid phase
- dentified platinum as a preferred catalyst choice for  $CO_2$ <br>eduction<br>- Stable with ionic liquids and potential windows (0 to 3 volts)<br>- Can be deposited on ion-exchange membranes using electroless<br>and electrolytic techniq that occurs if the Pt catalyst is coated or submerged in the ionic liquid/membrane phase – Stable with ionic liquids and potential windows (0 to 3 volts)<br>
– Can be deposited on ion-exchange membranes using electroless<br>
and electrolytic techniques<br>
– Some indications that CO is poisoning the Pt, adding Ru stab
	- the ionic liquid environment to maximize the reduction of  $CO<sub>2</sub>$



# Summary – Electrochemical CO<sub>2</sub> Reduction Reactor

#### • Ionic Liquid

- **Summary Electrochemical CO<br>
 High CO<sub>2</sub> reduction rates observed with ME<br>
 High CO<sub>2</sub> reduction rates observed with ME<br>
 MEAs having the ionic liquids with the EMIN<br>
 MEAs having the ionic liquids with the EMIN**  $-$  High CO<sub>2</sub> reduction rates observed with MEAs imbibed with ionic liquids having  $\mathsf{BF}_{4}$  anion
- **Summary Electrochemical CO<sub>2</sub> Reduction Reactor**<br>
This Liquid<br>
 High CO<sub>2</sub> reduction rates observed with MEAs imbibed with ionic liquids<br>
 MEAs having the ionic liquids with the EMIM cation performed better than<br>
tho those with the BMIM cation Summary – Electrochemical CO<sub>2</sub> Reduction Reactor<br>
— High CO<sub>2</sub> reduction rates observed with MEAs imbibed with ionic liquids<br>
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— MEAs having the Summary – Electrochemical CO<sub>2</sub> Reduction Reactor<br>
– High CO<sub>2</sub> reduction rates observed with MEAs imbibed with ionic liquids<br>
having BF<sub>4</sub> anion<br>
– MEAs having the ionic liquids with the EMIM cation performed better than
- 
- Processing metrics
	- reactor module can reduce 0.08 g/hr of  $CO<sub>2</sub>$  producing 0.03 g/hr of  $O<sub>2</sub>$

