

# **Reactive CO<sub>2</sub> Capture:** Process Integration for the New Carbon Economy

Peter Agbo, Sarah Baker, Todd Deutsch, Doug Kauffman, Josh Schaidle

February 18<sup>th</sup>-19<sup>th</sup>, 2020 Marriott Denver West, Golden CO

# **Workshop Goal and Objectives**

### Goal

Develop a vision for success for reactive  $CO_2$  capture within the context of a circular carbon economy and define a strategy for achieving that vision

### **Objectives**

- Develop relationships and bridge gaps between CO<sub>2</sub> capture and CO<sub>2</sub> utilization
- Identify major technical challenges, knowledge gaps, and barriers to progress
- Define research needs
- Establish metrics for success

### Why Reactive Capture?

### **Challenge:** High Costs of CO<sub>2</sub> Capture, Transport, and Storage



### **Opportunity**

Reactive CO<sub>2</sub> capture could offset costs of storage and enable a circular carbon economy while reducing need for CO<sub>2</sub> transport

National Petroleum Council, *Meeting the Dual Challenge, A Roadmap to At-Scale Deployment of Carbon Capture, Use, and Storage*, 2019

# **Definitions and Workshop Scope**

**Reactive Capture Definition:** The coupled process of CO<sub>2</sub> separation from mixed gas streams and conversion to valuable product(s)

### Can include:

- Integration of CO<sub>2</sub> separation and conversion in one step (e.g., catalyst-coated membrane)
- Integration of separation and conversion in one unit (e.g., regenerating capture media through CO<sub>2</sub> conversion during recycle)
- Process intensification in the pathway from CO<sub>2</sub> capture to products (e.g., reduced unit ops)

### Workshop Scope:

- Must form a valuable product, or mixture of products (e.g., more valuable than CO<sub>2</sub> loaded capture media)
- The product or mixture of products must be in a more reduced state than CO<sub>2</sub> (i.e., conversion of CO<sub>2</sub> must involve transfer of reducing equivalents)
- CO<sub>2</sub> source agnostic (e.g., atmospheric CO<sub>2</sub> and concentrated point sources included)
- Photosynthesis is outside the scope of this workshop

# Faradaic Electro-Swing Reactive CO<sub>2</sub> Adsorption

### Scientific Approach

- Leverages redox-active species to capture and release CO<sub>2</sub>
- CO<sub>2</sub> is captured via the carboxylation of reduced quinones
- CO<sub>2</sub> is released during cell discharge (reversed polarity)
- Cell architecture maximizes the surface area exposed to gas, allowing for ease of stacking of the cells in a parallel passage contactor bed

### Significance and Impact

- CO<sub>2</sub> uptake achieved at concentrations as low as 6000 ppm
- CO<sub>2</sub> capacity is independent of the inlet feed concentration
- >90% faradaic efficiency
- Offers an alternative to temperature-swing or pressureswing adsorption systems
- Opportunity for direct integration with CO<sub>2</sub> conversion



*Energy Environ. Sci.*, 2019, 12, 3530-3547 *International Journal of Greenhouse Gas Control* 2019, *82*, 48–58

# **Electrochemical Upgrading of CO<sub>2</sub> Capture Solution**

#### Scientific Approach

- CO<sub>2</sub> is captured in KOH solution to form (bi)carbonate ions
- Carbonate is fed to the cation conducting side of a bipolar membrane based CO<sub>2</sub> electrolyzer
- Protons supplied by the bipolar membrane generate CO<sub>2</sub> from carbonate
- CO<sub>2</sub> is reduced to CO, at its point of generation from CO<sub>3</sub><sup>2-</sup>, which also regenerates the hydroxide for further capture
- H<sub>2</sub> is also produced so pure syngas is the cathode product

### Significance and Impact

- Combined capture and conversion demonstration at a relevant current density – 150 mA/cm<sup>2</sup>
- Energy efficiency ~35%
- Stable operation over 145 hours
- Near 100 % carbon utilization no need to remove CO<sub>2</sub> from product stream



ACS Energy Lett. 2019, 4, 1427-1431

## **Renewable Methane Production**

### Scientific Approach

- Utilize excess electricity production for the electrolysis of water to  $\rm H_2$  and  $\rm O_2$
- Optimized strain of methanogenic archaea to perform methanation under industrial conditions
- 125 kW PEM electrolyzer feeds 2.5 kg H<sub>2</sub>/h, continuously producing 4.1 scfm CH<sub>4</sub>
- 98% carbon efficiency of CO<sub>2</sub> to CH<sub>4</sub>
- Post-processing for pipeline quality natural gas

#### Significance and Impact

- Potential long-term storage strategy via conversion of electricity and  $\rm CO_2$  to  $\rm CH_4$
- High efficiency CO<sub>2</sub> capture and conversion strategy
- Demonstrated route to renewable methane
- Large market and NG-grid to absorb curtailed electricity



# **CO<sub>2</sub> to Methanol via KOH/Ethylene Glycol**

#### Scientific Approach

- CO<sub>2</sub> is captured in a mixture of KOH and ethylene glycol
- Captured CO<sub>2</sub> is hydrogenated into methanol at mild temperatures using H<sub>2</sub> and a Ru-based catalyst
- Quantitative methanol yields after 20 hours under 70 bar H<sub>2</sub> and 140 °C
- Lower operating temperatures also possible
- Regeneration of capture solvent allows multiple capture/conversion cycles
- Demonstrated potential for direct air reactive capture

#### Significance and Impact

Clear potential for integrating direct air capture with production of a carbon neutral commodity chemical



#### pubs.acs.org/JACS

Communication

### Hydroxide Based Integrated CO<sub>2</sub> Capture from Air and Conversion to Methanol

Raktim Sen, Alain Goeppert, Sayan Kar, and G. K. Surya Prakash\*





Journal of the American Chemical Society, Article ASAP DOI: 10.1021/jacs.9b12711

## **Carbon Nanotubes via Molten Carbonate Electrolyzers**

#### Scientific Approach

- Molten carbonate electrolyzer
- Governing reactions:
  - (1)  $CO_2(g) + Li_2O \rightarrow Li_2CO_3$
  - (2)  $\text{Li}_2\text{CO}_3 \rightarrow \text{C}(\text{s}) + \text{Li}_2\text{O} + \text{O}_2(\text{g})$
  - (Net)  $CO_2 \rightarrow C + O_2$
- Control carbon nanofiber morphology via current density, electrolyte (Li-K-Na), and electrolytic temperature

#### Significance and Impact

- Potential for high coulombic and carbon efficiencies if Li<sub>2</sub>CO<sub>3</sub> is not consumed during the reaction and is continuously regenerated from CO<sub>2</sub>
- High-value product
- Leverages atmospheric CO<sub>2</sub>





#### One-Pot Synthesis of Carbon Nanofibers from CO<sub>2</sub>

Jiawen Ren,<sup>†</sup> Fang-Fang Li,<sup>†</sup> Jason Lau,<sup>†</sup> Luis González-Urbina,<sup>†</sup> and Stuart Licht<sup>\*,†</sup> <sup>†</sup>Department of Chemistry, The George Washington University, Washington, DC 20052, United States



Nano Lett. 2015, 15, 6142-6148; Energy Conversion and Management 2016, 122, 400-410

## Agenda

### What does success look like?

#### February 18<sup>th</sup>, 2020 February 19<sup>th</sup>, 2020 1:00pm Meeting Kick-off 7:00-8:00am Breakfast Bill Tumas (NREL) and Roger Aines (LLNL) David Miller (NETL) 1:15-2:00pm 8:00-8:30am 2:05-2:50pm Sean Simpson (Lanzatech) 8:35-9:05am David Heldebrant (PNNL) 2:50-3:20pm Break 9:05-9:20am Break 3:20-5:00pm Panel Discussion 9:20-9:50am Etosha Cave (Opus-12) Matthew Kanan (Stanford University) • Ian Rowe (Bioenergy Technologies Office 9:55-10:25am • Lynn Brickett (Office of Fossil Energy) **Organize into Breakout Sessions** 10:25-10:45am • Paul Kenis (University of Illinois) 10:45-12:00pm Breakout Session (5 rooms) Todd Wilke (Carbon Engineering) 12:00-1:00pm Lunch • Bill Tumas (NREL) 1:00-3:00pm **Breakout Sessions** • Roger Aines (LLNL) 3:00-3:15pm Break 5:00-5:30pm\* Break (Poster presenters set up posters) 3:15-4:00pm **Breakout Session Readouts** 5:30-8:00pm **Reception and Poster Session** 4:00-4:15pm Closing Remarks

How do we achieve success?

\*All moderators please meet in the Golden Ballroom at 5pm today to discuss roles and responsibilities

## **Ground Rules**

- All ideas/thoughts are welcome give everyone a chance to contribute
- Keep an open mind
- Please step out if you need to take a call
- Think big, check your baggage at the door, and have fun

# **Thank You!**

- Speakers and Panelists
- All attendees
- Linda Stolmack
- Co-Organizers
- Department of Energy















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