

Reactive Carbon Capture Observations from ARPA-E 2022 Workshop

Jack Lewnard
ARPA-E Program Director

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2022 ARPA-E CO₂ Utilization/Reactive Carbon Capture Workshop

- ▶ Re-imagine the problem
 - Dramatically reduce cost and environmental footprint of C₁ and/or C₂ products
 - Leverage commercial downstream processes
 - Skip steps, and their related hardware and energy inputs
 - Leverage new materials/chemistry (ionic liquids, MOFs, homogeneous catalysts)
 - Maximize process intensification
 - <https://arpa-e.energy.gov/events/reactive-carbon-capture-workshop>
- ▶ Reactive Carbon Capture
 - Capture CO₂ and react it while in adsorbed/absorbed state
 - No intermediate CO₂ production, purification, compression
 - https://netl.doe.gov/projects/files/SummaryReportoftheReactiveCO2CaptureProcessIntegrationfortheNewCarbonEconomyWorkshop_08242021.pdf
- ▶ React CO₂ and separate the product(s), esp where CO₂ is hot and in reducing environment
 - Replace CO₂ capture with easier product separation (ie MeOH in water wash)

Options for RCC

► Inorganic RCC

- Capture CO₂ with weak or strong base
 - Cement, pozzolanic ash, minerals
 - “Sequesters” the CO₂ as a mineral
 - Can operate over a wide range of temperatures (although rates vary)
 - Minimal concerns about impurities

– Organic RCC

- React CO and/or CO₂ to make useful products
- May be easier to separate products than CO₂ (ie MeOH)

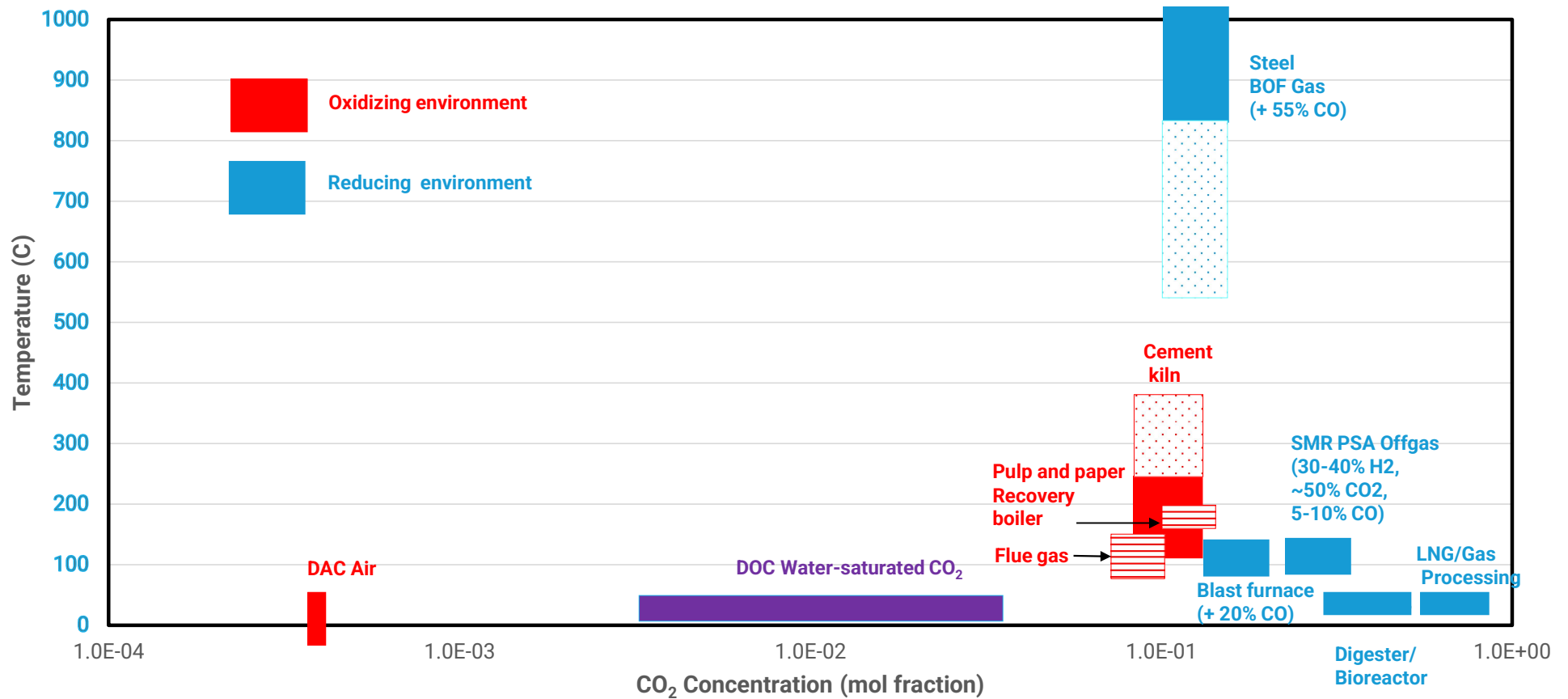
– Biological

- How Nature does it
- Lanza Tech
- ARPA-E ECOSYNBIO

Organic and Biological RCC Challenges

- ▶ Carbon Capture (first step)
 - Prefer low temperature, high pressure, no competing species
 - “Goldilocks” affinity – less than a chemical bond, more than van der Waals forces
 - High selectivity for CO or CO₂ – esp. avoid O₂
- ▶ Reaction (second step)
 - Catalytic processes prefer high temperature, high pressure
 - Plasma processes prefer low pressure
 - No free oxygen to compete for reducing equivalents
- ▶ Biology already does this – what can we learn?

CO and CO₂ Sources/Attributes



System-Level Considerations

- ▶ “Contaminants”
 - O₂, possibly SO_x and NO_x from oxidizing environment (air and combustion point sources)
 - H₂S, NH₃, particulates from reducing environments
 - Possibly many from water-borne CO₂

- ▶ Temperature
 - Adsorption/absorption favors low temperature
 - Reaction favors high temperature
 - Not clear if there is a good middle ground
 - Cooling below 40 C is not easy or cheap

- ▶ Pressure drop to contact CO₂ from air or flue gas can be energy-intensive

- ▶ Liquids can be pumped, easy to change temperature

- ▶ Solids are hard to move, hard to change temperature. Adsorption system capturing CO₂ from oxidizing environments usually require multiple beds with interim purge steps

- ▶ Operating intermittently (to access off-peak electricity or accommodate variable flow CO₂ sources) can be difficult, lowers capital utilization, and increases costs