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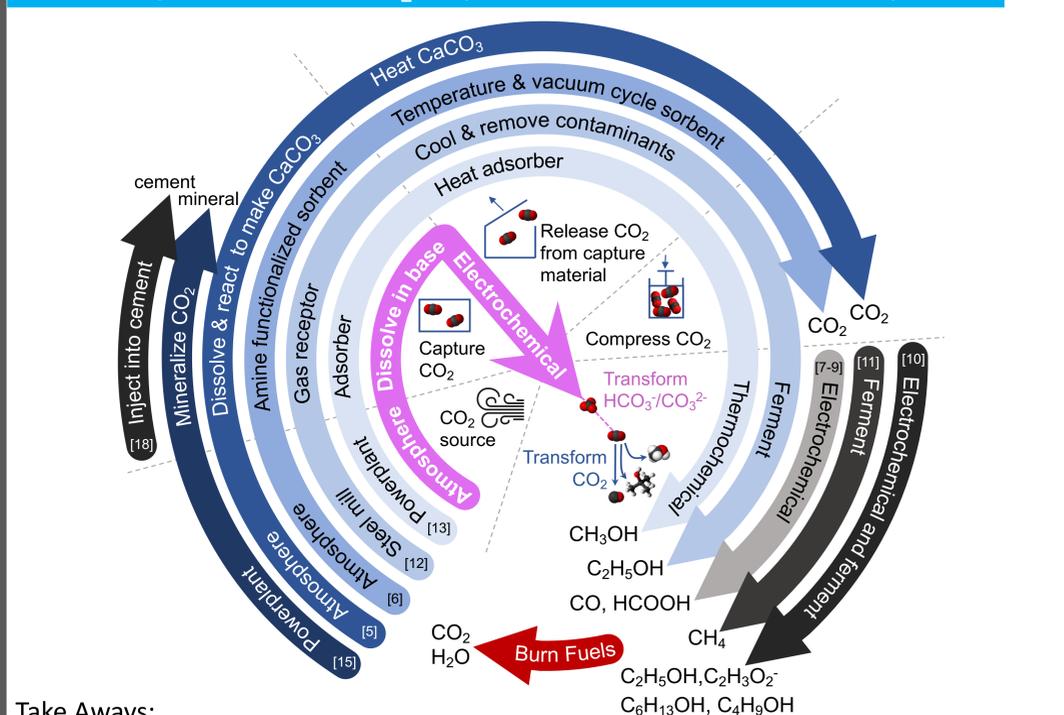
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**Motivation and Overview** Designing a complete system to capture CO<sub>2</sub> from the air and transform it into valuable chemicals is of utmost importance for mitigating climate change. The processes of CO<sub>2</sub> sequestration, CO<sub>2</sub> transformation, and product separation all require significant energy inputs - therefore devising a system that simultaneously minimizes all of these steps is challenging. To date, a variety of CO<sub>2</sub> sequestration and/or conversion systems have been built targeting these individual aspects. Here we propose a new paradigm for designing CO<sub>2</sub> capture and conversion systems: (i) formation of bicarbonates/carbonates through dissolution of CO<sub>2</sub> into basic solutions; followed by (ii) electrochemical reduction; and (iii) transformation into valuable chemicals via industrial processes. Unlike traditional systems in which gaseous CO<sub>2</sub> reacts with a catalyst, our design focuses on the transformation of bicarbonate or carbonate ions from solution which offers several advantages. First, the CO<sub>2</sub> sequestration from the atmosphere does not require an energy intensive heating step to recover gaseous CO<sub>2</sub> for later transformation, and 75% of the CO<sub>2</sub> is removed on the first pass. Second, by transforming bicarbonate/carbonate ions, the process avoids any energy intensive CO<sub>2</sub> compression and allows for significantly higher conversion percentages. Taken together, we anticipate significant reductions in overall energy consumption can be achieved by focusing attention on the conversion of carbonate/bicarbonate ions instead of gaseous CO<sub>2</sub>.

## Summary of Current CO<sub>2</sub> Capture and Conversion Methods

CO <sub>2</sub> Source	Capture CO <sub>2</sub>	Release CO <sub>2</sub> from capture material	Compress CO <sub>2</sub>	Transform CO <sub>2</sub>	Products
Environment - Air - Seawater	Absorption - e.g. amines	Heat	~15 PSIA for fermentation	Electrochemical	C <sub>2</sub> H <sub>4</sub> O <sub>2</sub> C <sub>2</sub> H <sub>2x</sub> C <sub>x</sub> H <sub>2x+1</sub> C <sub>x</sub> H <sub>2x+2</sub>
Point Source - Petrochemical Industry - e.g. ethylene - Iron and Steel - Cement production - Natural gas sweetening - Fossil fuels or Minerals - e.g. coal	Adsorption - e.g. zeolites  Membranes - e.g. fibers  Biological - e.g. algae  Other - e.g. mineralization	Vacuum cycle	~15-50 PSIA for electrochemical processes  ~250-1450 PSIA for thermochemical processes	Fermentation  Photo-electrochemical  Photocatalysis  Thermocatalytic  Thermolysis  Industrial Processes	Diesel Syngas Ethylene Butanol Hexanol Propanol Ethanol Plastics

## Summary of Current CO<sub>2</sub> Capture and Conversion Companies



### Take Aways:

- Releasing CO<sub>2</sub> from the capture media and compressing CO<sub>2</sub> for storage or for use by conversion processes is energy intensive
- Direct air capture is more difficult than point-source capture, and CO<sub>2</sub> transformation is more energy intensive than CO<sub>2</sub> capture
- Significant energy savings are possible when you do not have to compress the CO<sub>2</sub> or release it from the capture material

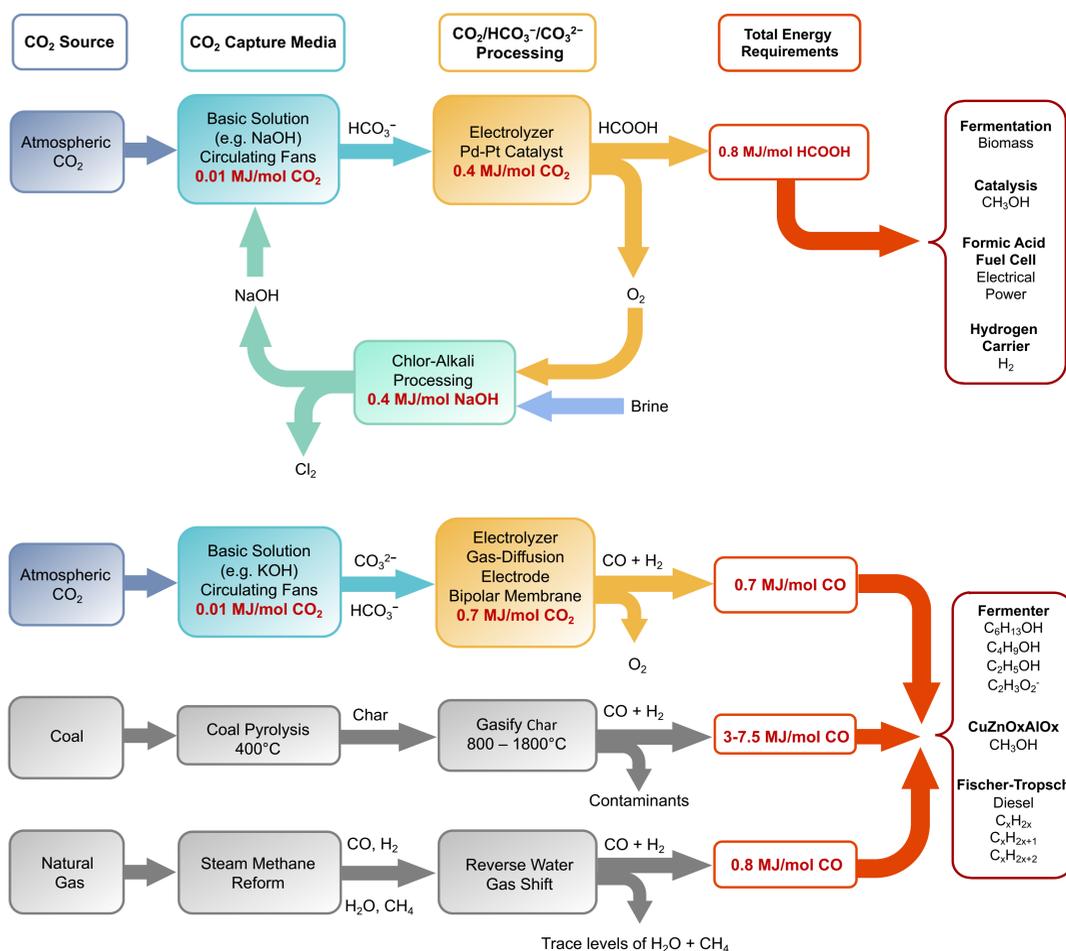
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## Results



### Take Aways:

- The proposed process generates syngas for 0.7 kJ/mol CO<sub>2</sub>, which could then be used in a variety of industrial processes to generate valuable chemicals
- Generating syngas in our proposed method leads to an extremely pure syngas feed unlike industrial processes
- Generating syngas from coal and natural gas is more energy expensive than our proposed process, and the reported energies do not include the energy required to mine the coal or drill for the natural gas
- Our proposed processes would allow for 100% utilization of captured CO<sub>2</sub>
- Energy required by other pilot CO<sub>2</sub> capture and conversion companies is higher than what we have proposed for a system that does both direct air capture and syngas synthesis

### Comments on Calculations:

- All electrolyzers are assumed to perform the oxygen evolution reaction at the anode
- The total energies calculated would most likely be higher in a realistic plant due to the need for pumps, possible catalyst regeneration, etc.

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