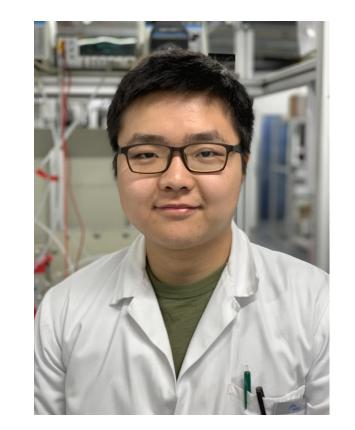
### Sustainable Carbon Cycle



european centre of **excellence** for sustainable **water technology** 

# Novel methods for electrochemical capture and conversion of CO<sub>2</sub>



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#### Motivation

Global warming is one of the most critical global challenges.

	Pure CO <sub>2</sub>	
		<b>^</b>
HRES system		

Increasing atmospheric  $CO_2$  concentration brought by anthropogenic emission is the major reason for this climate change problem. Capturing  $CO_2$  from emission points and even directly from air provides a feasible solution to mitigate the amount of  $CO_2$  emission and reduce the atmospheric  $CO_2$  concentration. At Wetsus, under the theme Sustainable Carbon Cycle, we aim to develop novel  $CO_2$  capture technologies that could be potentially energy efficient and environmentally benign <sup>1</sup>.

# **Technological challenge**

 $CO_2$  can be captured by alkaline aqueous sorbent due to the high solubility of  $CO_2$  under high pH. However, the conventional regeneration of the sorbent by calcination consumes a large amount of energy. Therefore, we aim to develop an alternative process for regenerating the alkaline solution in wet scrubbing process for direct air capture (DAC) application.

This alternative process is designed to create a pH-swing of the

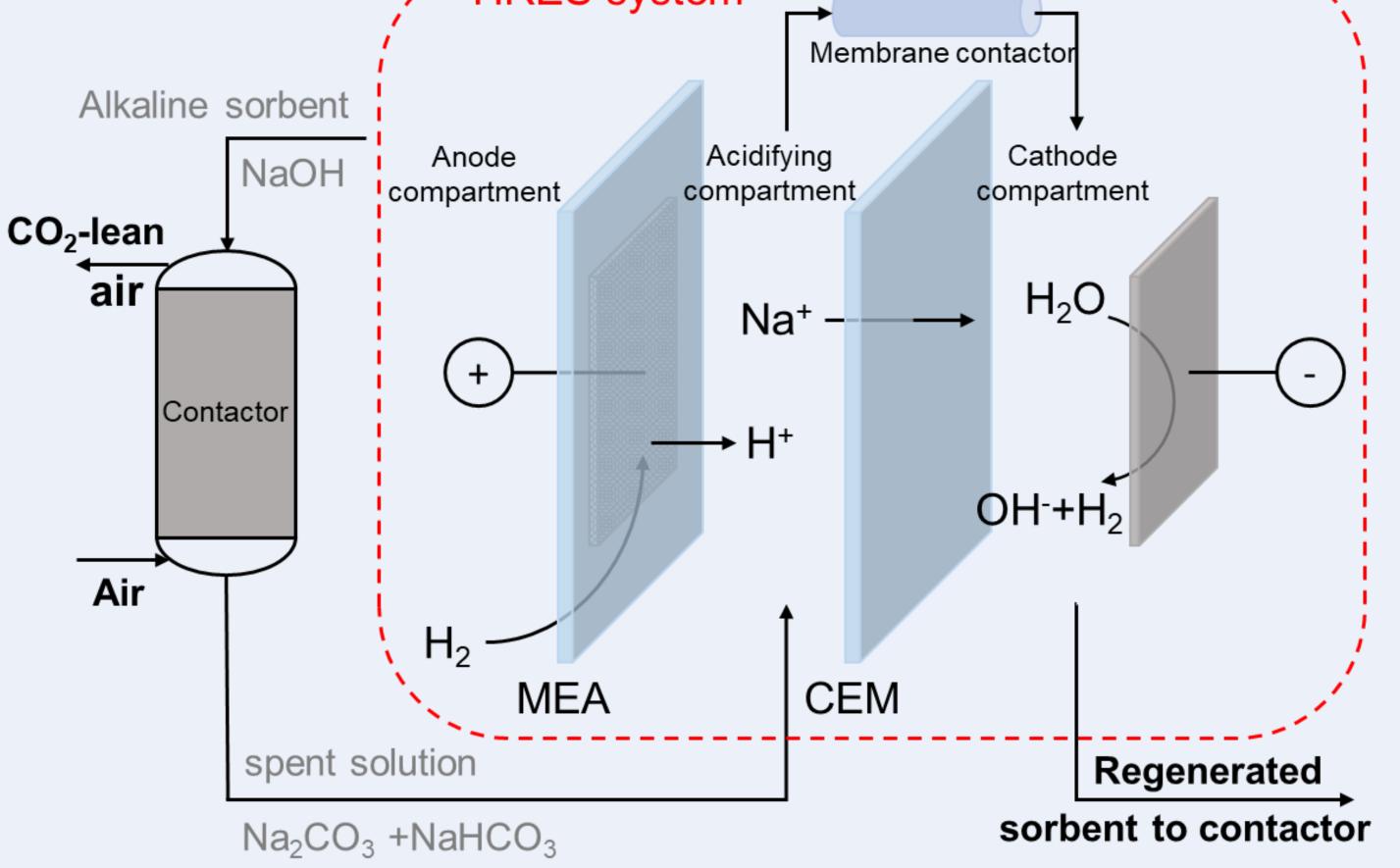
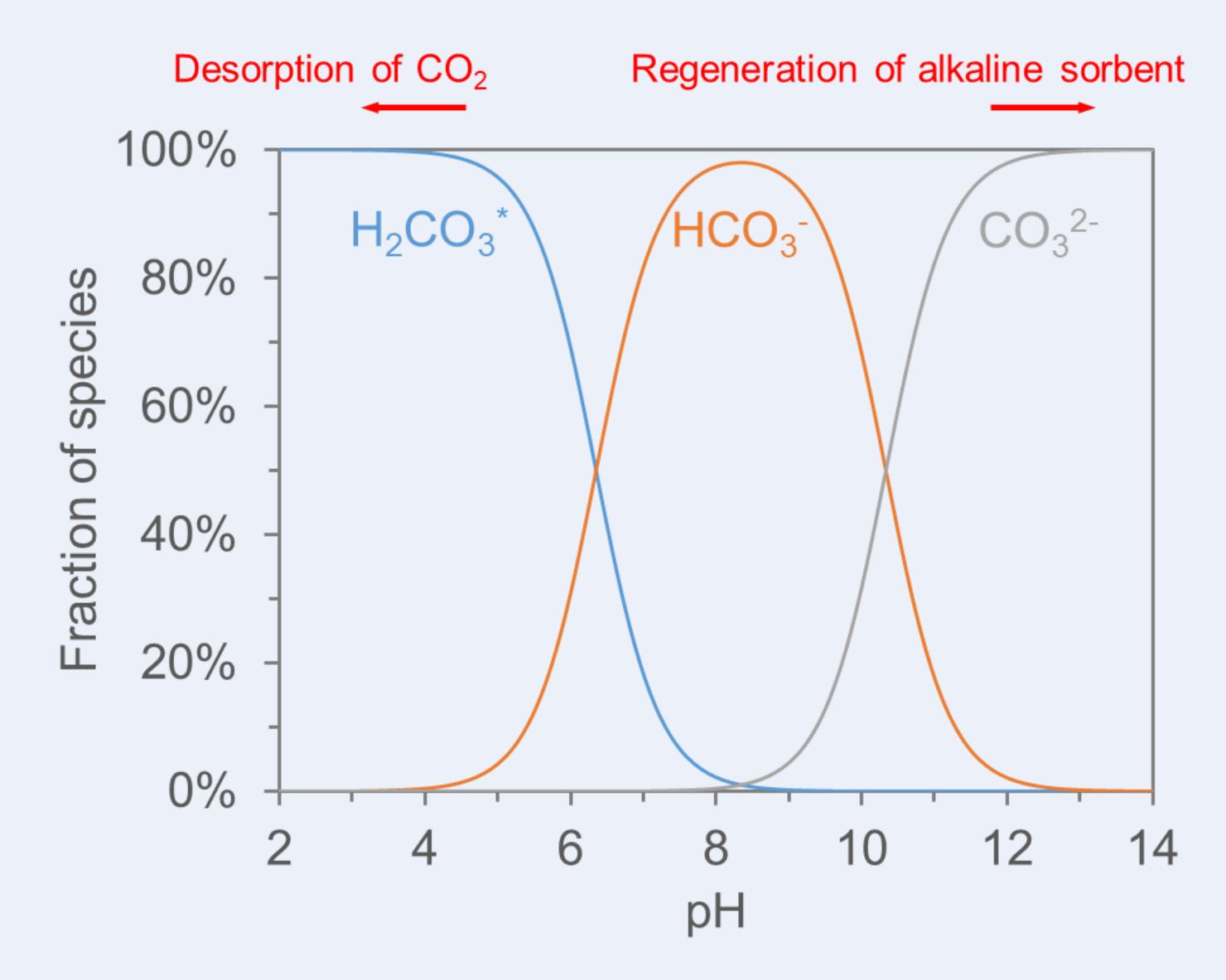


Fig 2. Principle of DAC process using a  $H_2$ -recycling electrochemical system (HRES) for  $CO_2$  desorption and regeneration of alkaline sorbent.

The process is based on a  $H_2$ -recycling electrochemical system (HRES) which was originally developed for nitrogen recovery from wastewater <sup>2</sup>. Alkaline and acidic conditions are created in two adjacent compartments respectively in HRES, so simultaneous desorption of CO<sub>2</sub> and regeneration of alkaline sorbent could be achieved after the spent solution flows through the electrochemical cell <sup>3</sup>.

solution. We have noticed that the solubility of  $CO_2$  in aqueous solvent is dependent on the pH of the solution (Figure 1). Thus,  $CO_2$  can be desorbed at low pH and the sorbent can be regenerated at high pH. The configuration of the system is shown in Figure 2.



## **Research goals**

- Developing a novel electrochemical system for CO<sub>2</sub> capture based on the scheme shown in Figure 2
- Investigating the CO<sub>2</sub> capture performance and energy consumption of the system under different conditions
- Studying the performance of the system with different sorbents
- Developing a mathematical model of the system describing the kinetics and transport of different components
- Scaling-up study by integrating pairs of bipolar membranes in the electrochemical cell

Fig 1. The fractions of different aqueous carbon species at different pH.

[1] Legrand et al., Solvent-Free CO2 Capture Using Membrane Capacitive Deionization. *Environmental science & technology* 2018, *52*, (16), 9478-9485.
[2] Kuntke et al., Hydrogen Gas Recycling for Energy Efficient Ammonia Recovery in Electrochemical Systems. *Environmental Science & Technology* 2017, *51* (5), 3110-3116.
[3] Hamelers et al, Electrochemical device, system and method for electrochemical recovery and/or regeneration of carbon dioxide from a stream, NL2025044

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