

# Development of a CO<sub>2</sub> Direct Ocean Capture System Using Bipolar Membrane Electrodesialysis

This work investigates the acid and base generation mechanism, efficiency and durability of bipolar membrane electrodesialysis and its application in CO<sub>2</sub> direct ocean capture using pH swing. The system provides an efficient and cost-effective solution to reduce CO<sub>2</sub> level and ocean acidification.

Kaiwen Wang<sup>1</sup>, Chengxiang Xiang<sup>1,2</sup>

1. California Institute of Technology, Pasadena, CA, USA

2. Captura, Pasadena, CA, USA

## Introduction & Goals

Based on the pH-swing of the dissolved inorganic carbon species, a CO<sub>2</sub> direct ocean capture system exploiting bipolar membrane (BPM) electrodesialysis was designed. Models using CFD and tertiary current distribution are developed focusing on optimizing the cell structure to improve faradaic efficiency for acid and base generation and prevent over-heating for better durability even when operating at high current density. The environmental impact and CO<sub>2</sub> drawdown ability is also investigated with a mesoscale CFD and mass transport model on an ocean level.

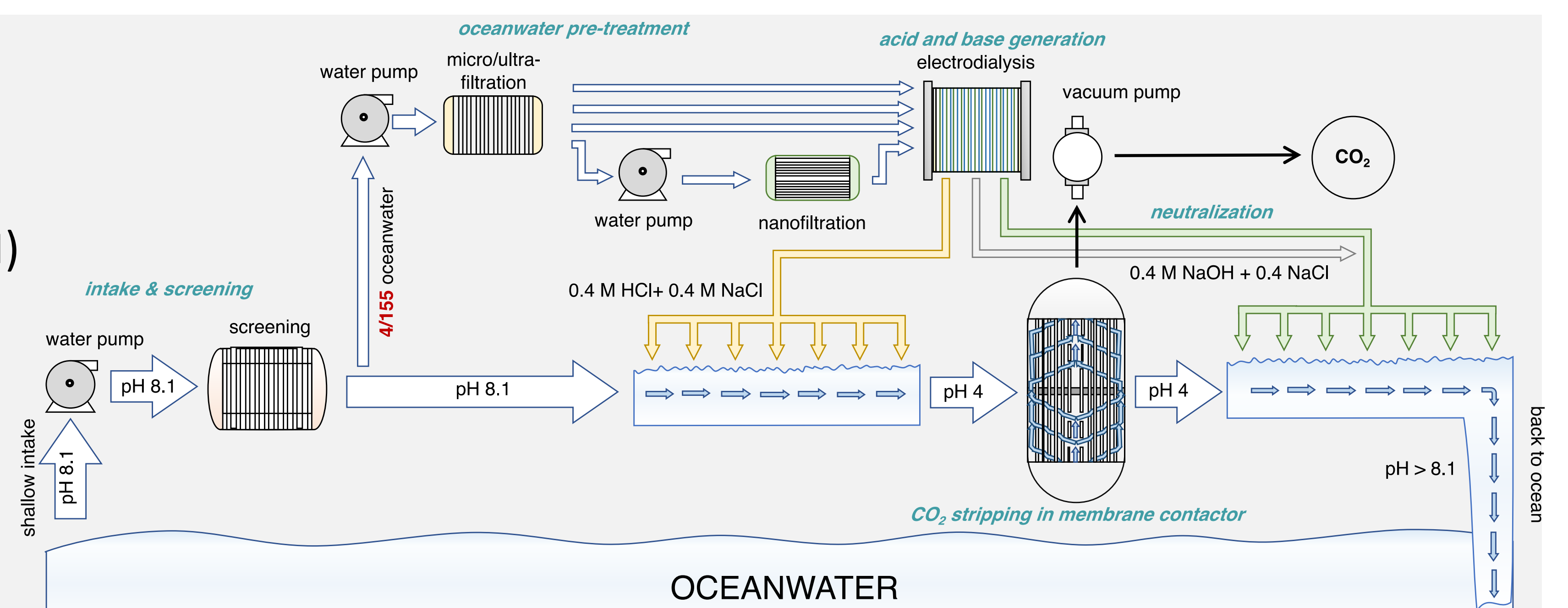
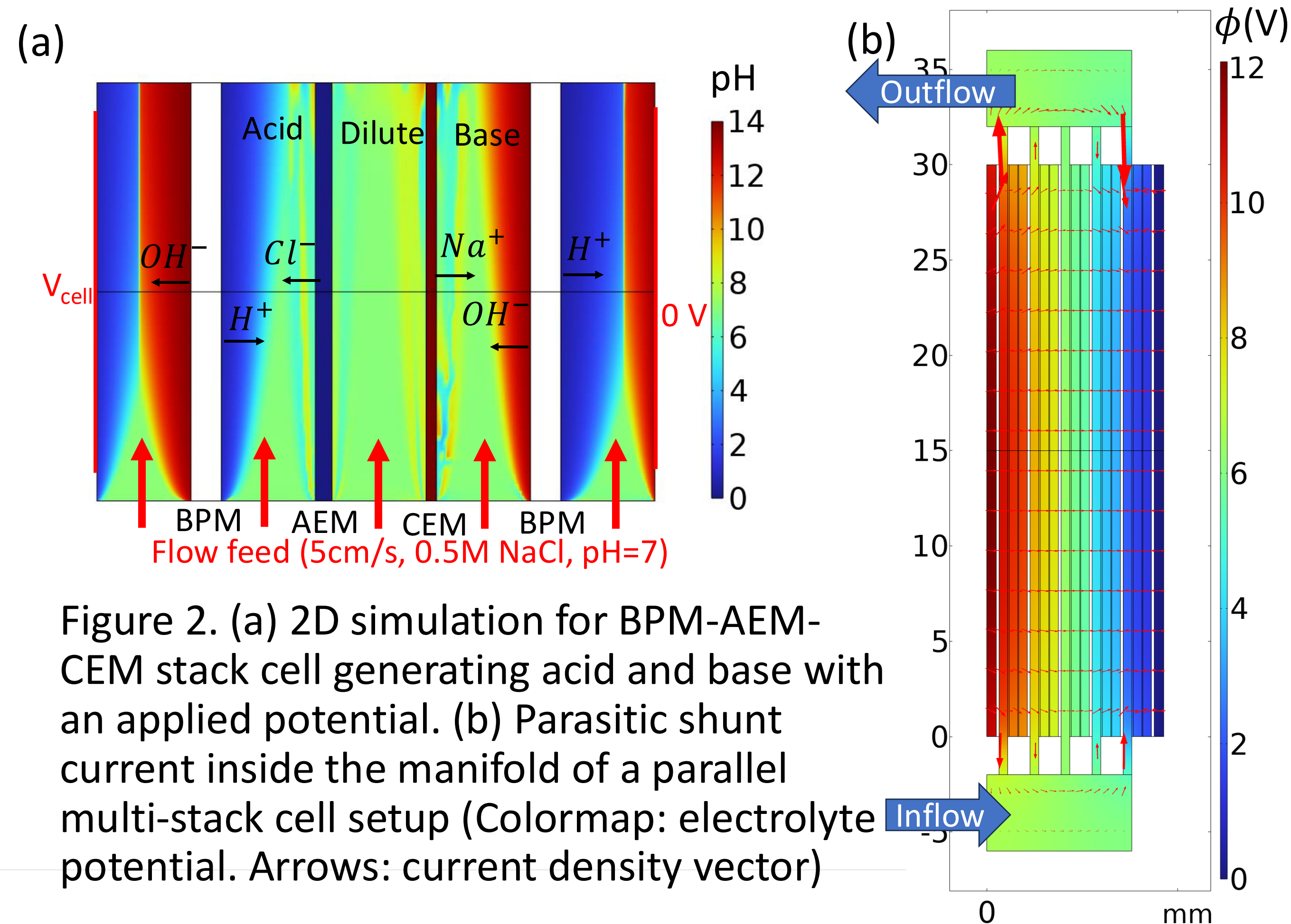


Figure 1. General schematic of the direct ocean capture CO<sub>2</sub> removal system



## Bipolar Membrane Electrodesialysis

A BPM electrodesialysis stack cell is modeled in 2D using the CFD and tertiary current distribution module. The selectivity of the anion exchange membrane (AEM) and cation exchange membrane (CEM) are simulated through a Donnan boundary condition at the membrane surface:

$$\phi_l - \phi_m = -\frac{RT}{zF} \ln\left(\frac{c_l}{c_m}\right)$$

BPM is a combination of the AEM and CEM with a very thin catalyst layer in between them, which promotes water dissociation ( $H_2O \rightleftharpoons H^+ + OH^-$ ). Through applied potential and ion selectivity of the membranes, the electrodesialysis cell can generate acid and base.

The model was used to improve the faradaic efficiency of the cell by optimizing geometry to reduce shunt current, which is the parasitic current flowing through the manifold when multiple cell are setup in parallel.

## CO<sub>2</sub> Ocean Drawdown

To ensure efficient CO<sub>2</sub> capture while not causing too aggressive disturbance to the marine ecosystem, an area with slightly higher pH needs to be established as an enhanced CO<sub>2</sub> drawdown site. This ocean mesoscale model studies the turbulent flow and its influence on the mixing and drawdown of CO<sub>2</sub>.

A flux boundary condition was assigned to the top of the seawater domain depending on the gas transfer velocity [2]:

$$J_{CO_2} = k(Hp_{CO_2,air} - c_{CO_2,water})$$

Inside the domain, an enhanced diffusivity ( $D_T = \nu_T / Sc_T$ ) was used to address the turbulent mixing. And reactions between dissolved inorganic carbon species were

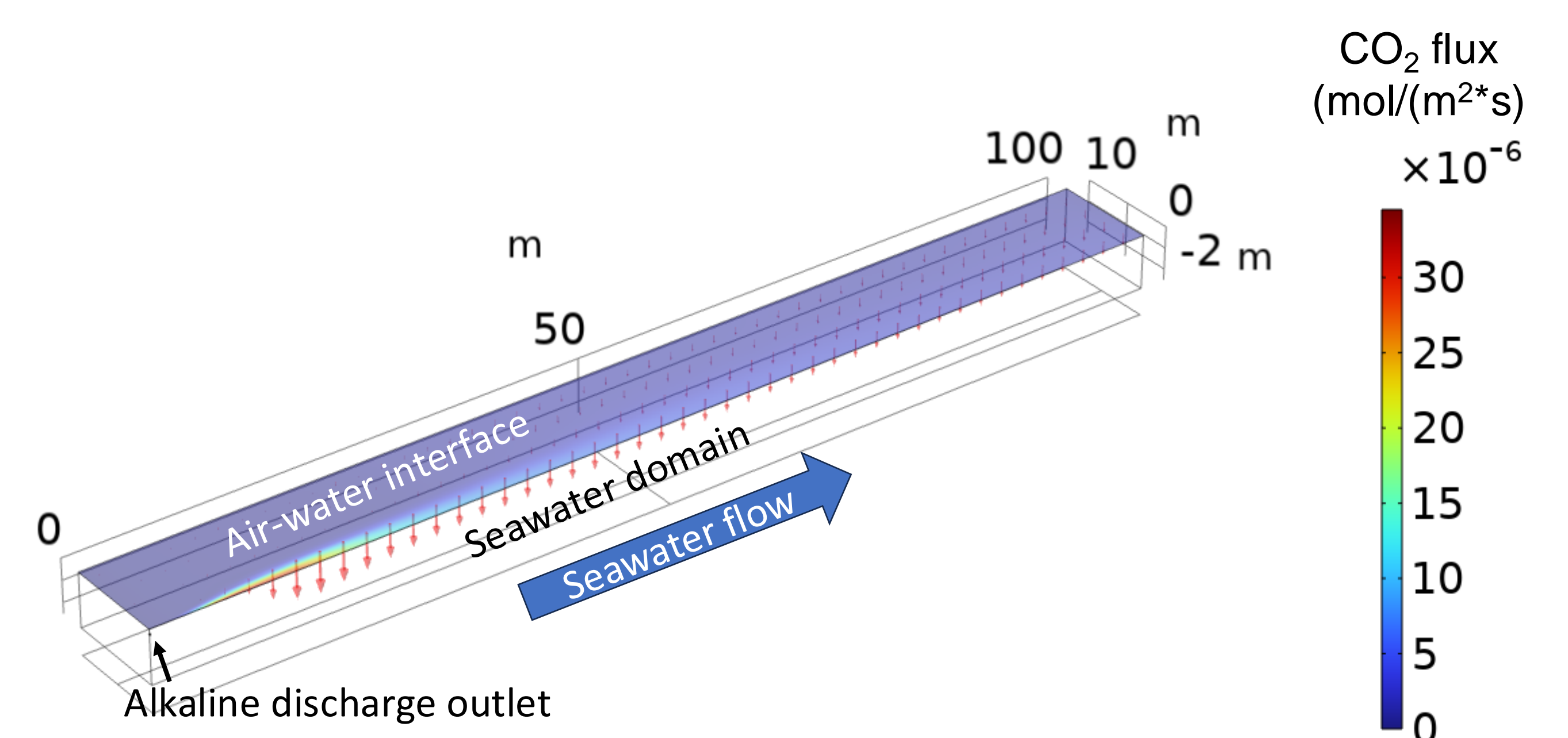
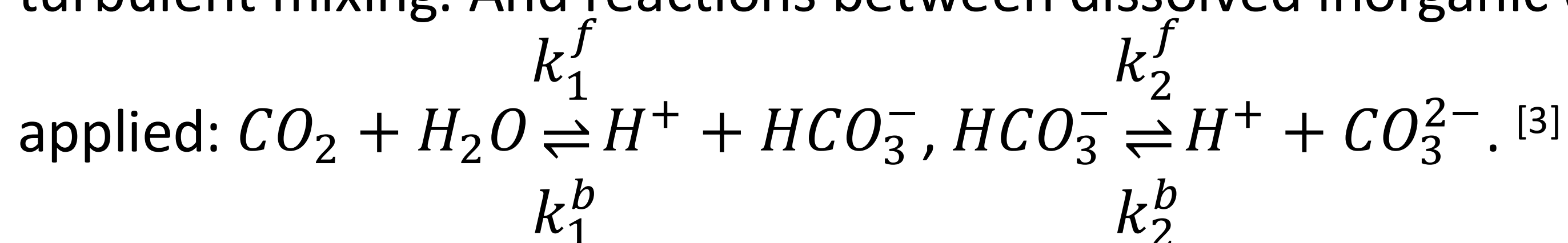


Figure 3. CO<sub>2</sub> drawdown rate at the alkaline discharge.

## REFERENCES

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