Mixing-controlled reactive transport at the pore scale and its impact on flow field and upscaling of reactive transport

H. YOON^{1*} AND T. DEWERS¹

¹Sandia National Laboratories, Albuquerque, NM 87123, USA (*correspondence: hyoon@sandia.gov)

Mixing-controlled pore-scale reactive transport

Dissolved CO2 during geological CO2 storage may react with minerals in fractured rocks or confined aquifers resulting in mineral precipitation and dissolution. Complex pore structures in reservoirs also influence flow complexities that can have a significant impact on coupled pore-scale phenomena including fluid dynamics in capillary systems, multiphase flow, and heterogeneous geochemical reactions. In this work, pore-scale experiments on transverse-mixing induced geochemical reactions in microfluidic pore networks (micromodel) are used as a basis for understanding coupled processes among hydrodynamics, transport, and reactions at the (sub) pore-scale. In particular, precipitation and dissolution dynamics affect the flow field, resulting in transient behavior of reactions. We will apply micro particle image velocimetry (µPIV) using laser scanning confocal microscopy for characterizing 3-D fluid velocities in dynamic precipitation and dissolution regimes. Pore-scale models of coupled fluid flow, reactive transport, and heterogeneous reactions are applied to account for transient 3-D experimental results of CaCO₃ precipitation and dissolution.

Upscaling of reaction rates

Response function of reaction rates will be constructed from pore-scale simulations which account for a range of reaction regimes characterized by the Damkohler and Peclet numbers under different pore configurations. Newly developed response functions will be used in a continuum scale model that may account for large-scale phenomena. In particular, this work is motivated by the observed CO₂ seeps from a natural analog to geologic CO2 sequestration at Crystal Geyser, Utah where the lateral migration of CO2 pathways was observed at a scale of ~ 100 meters over time. Implications for evaluating mineral precipitation patterns observed in natural analogues for CO2 storage and leakage are discussed to link pore-scale processes to the field scale observations. Comparison of field observations and continuum-scale simulation results will provide mechanistic explanations of the lateral migration of CO₂ pathways.

Sandia National Laboratories is a multi-program laboratory managed and operated by Sandia Corporation, a wholly owned subsidiary of Lockheed Martin Corporation, for the U.S. Department of Energy's National Nuclear Security Administration under contract DE-AC04-94AL85000.