

MODELING CO₂-WATER-ROCK REACTIONS IN A BASALT FRACTURE NETWORK

CO₂ mineral trapping
Permeability alteration
Fracture network
Carbon sequestration

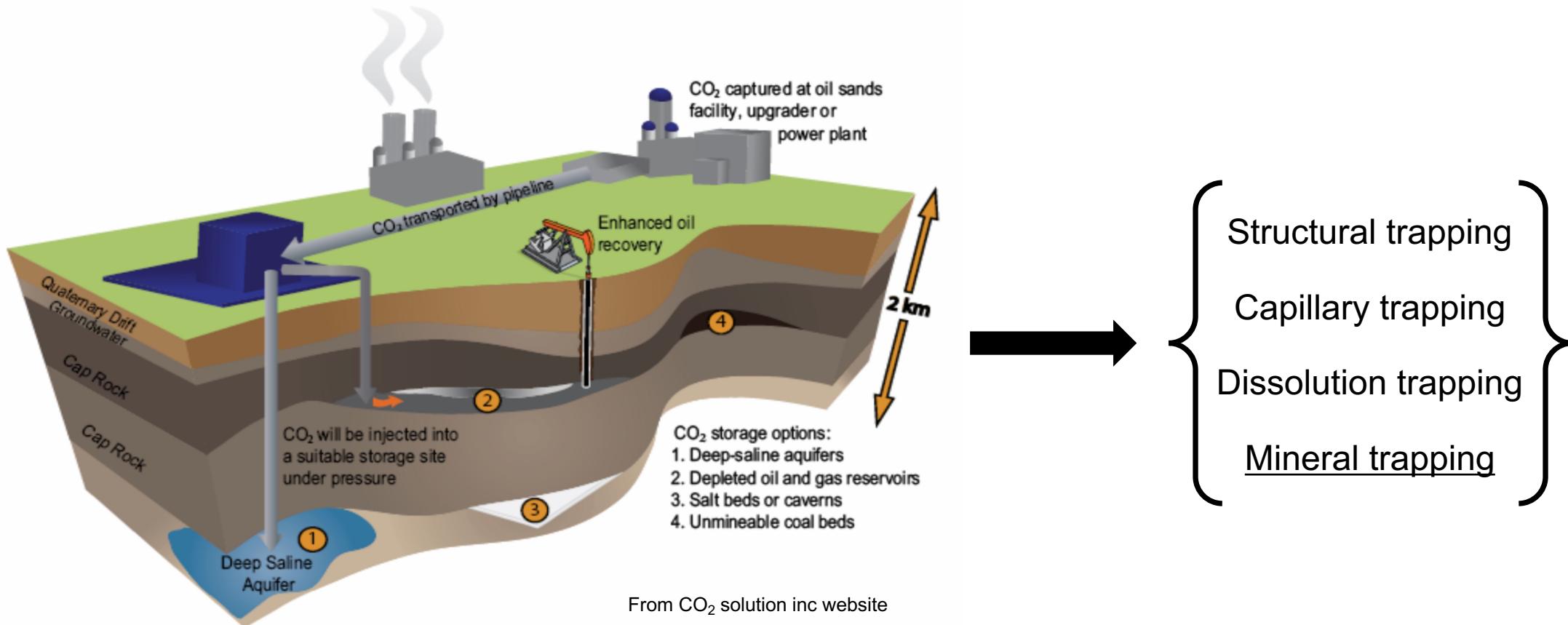


WU Hao (LANL)
Richard S. Jayne (Sandia)
Robert Bodnar (VT)
Ryan M. Pollyea (VT)



CARBON SEQUESTRATION

Carbon Capture and Storage (CCS)



MINERAL TRAPPING IN BASALT

Basalt reservoirs

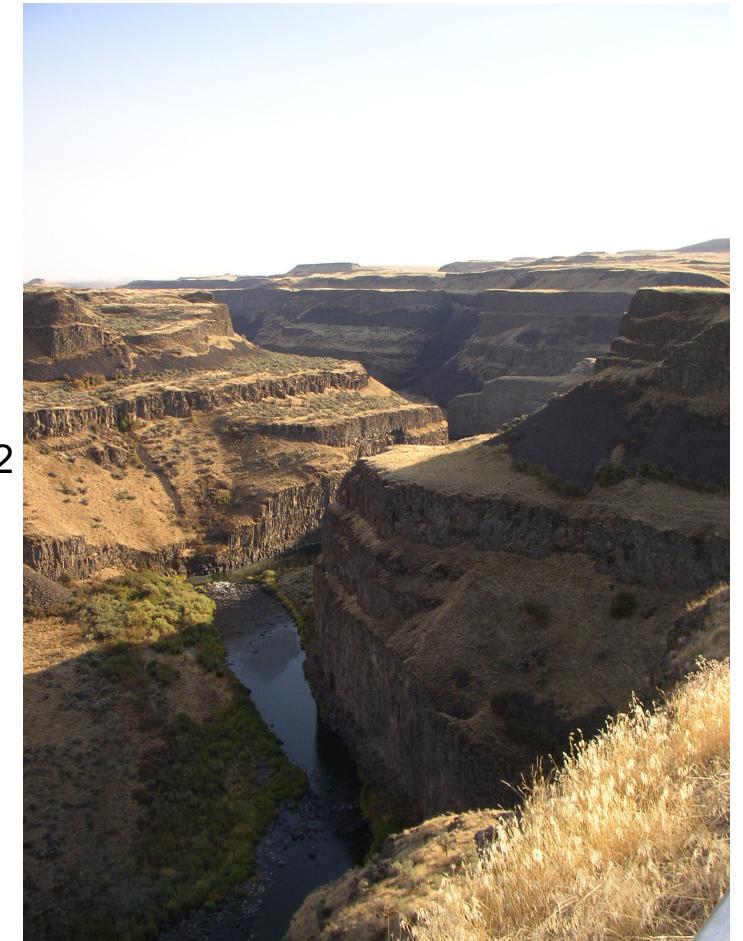
- High concentration of cations (Ca, Mg, Fe)
- Worldwide distribution & immense volumes

Columbia River Basalt Group: 164,000 km²

India's Deccan Volcanic Province (DVP): ~500,000 km²

Pilot-scale CO₂ injection tests

- Iceland CarbFix Project
95% of injected CO₂ mineralized in two years
- Wallula Basalt Sequestration Pilot Project
carbon mineralization widespread in injection zone

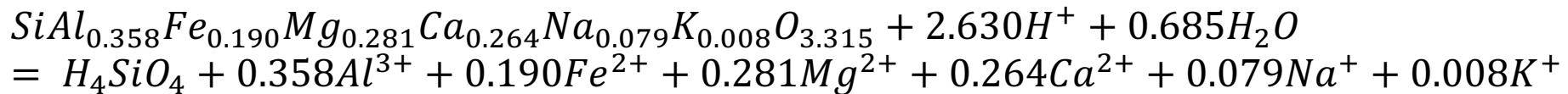


CHEMICAL REACTIONS

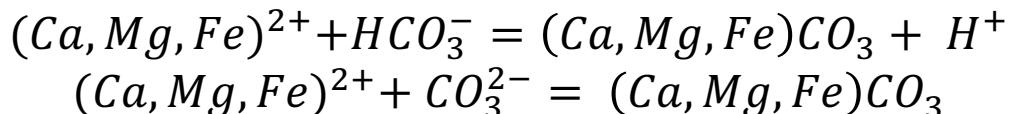
(1) CO₂ Dissolution



(2) Basalt Dissolution (Icelandic basalt)

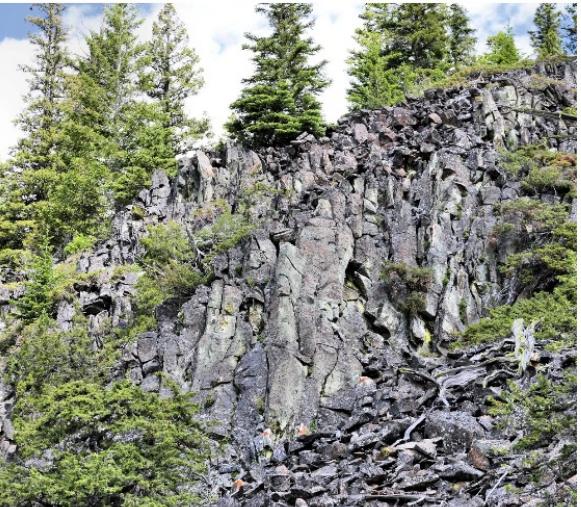


(3) Carbonate Precipitation



From Matter & Kelemen (2009)
Pollyea & Rimstidt (2017)

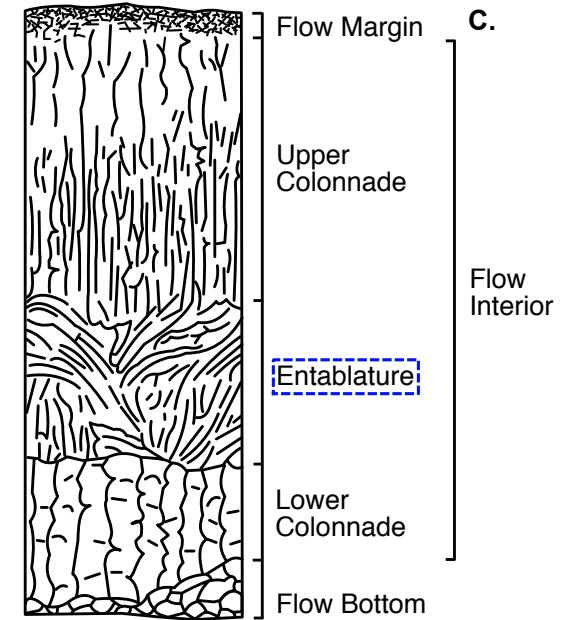
BASALT FORMATIONS ARE HIGHLY FRACTURED



Basalt bluffs of
McConnell Hill,
Canada

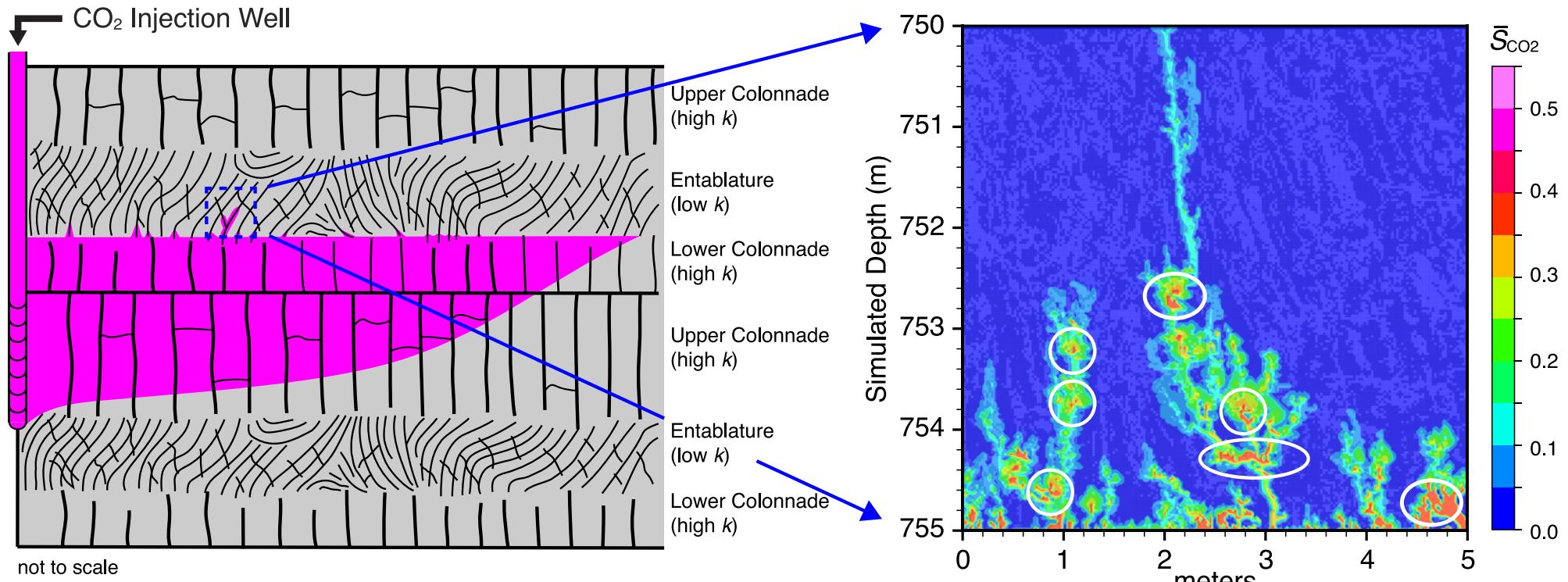


Columbia River
Basalt Group,
Washington



Morphology of Columbia
River Basalt Group
(CRBG)

RESEARCH SHOWS THAT FREE-PHASE CO₂ TENDS TO ACCUMULATE AT FRACTURE INTERSECTIONS



Gierzynski & Polyea (2017)

Stochastic simulations

50 permeability distributions

Conclusion

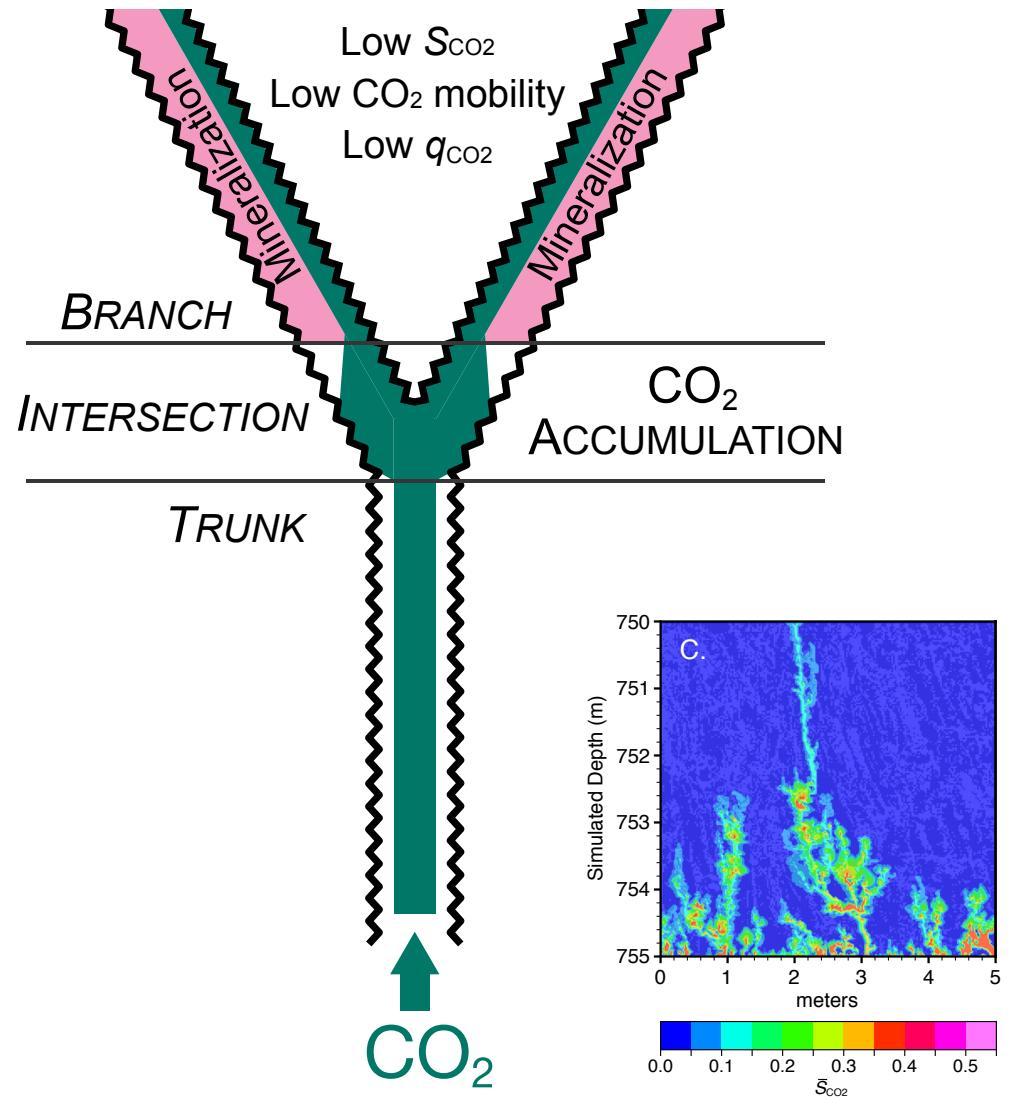
CO₂ accumulates at fracture intersections
due to multiphase flow effects

HYPOTHESIS

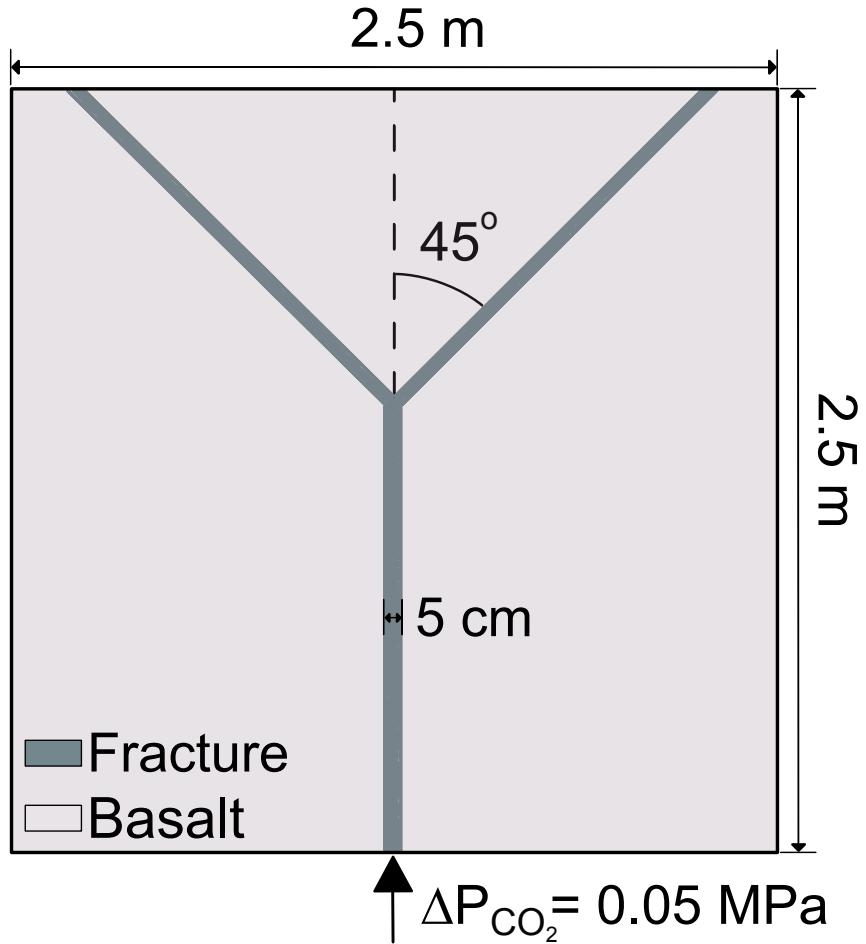
- Carbon mineralization is likely to focus at branching fractures.
- CO₂ vertical migration will be reduced due to carbon mineralization.

Hypothesis Test

Use numerical simulation to model how different permeability alteration scales effect this carbon mineralization process.



MODEL DOMAIN



Boundary Condition

Top: open system

Bottom: 0.05 MPa over pressure

Initial Condition

33 °C

7.5 MPa

Permeability

Fracture: $5.78 \times 10^{-16} \text{ m}^2$

Basalt: 10^{-20} m^2

Simulation

TOUGHREACT

Ten years

PERMEABILITY ALTERATION

$$k = k_o \left(\frac{\phi}{\phi_o} \right)^n$$

Verma & Pruess (1988)

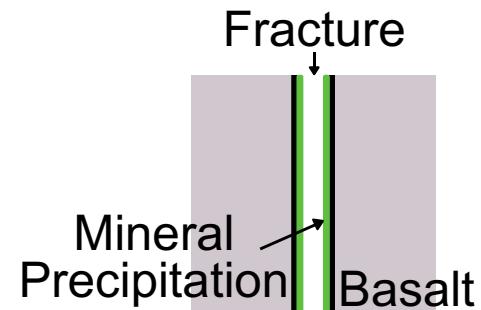
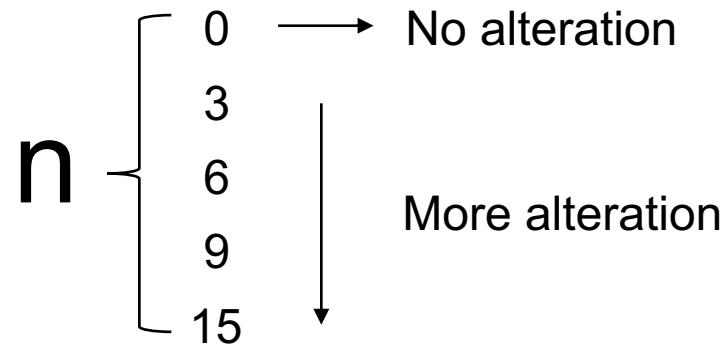
k permeability

k_o initial permeability

ϕ porosity

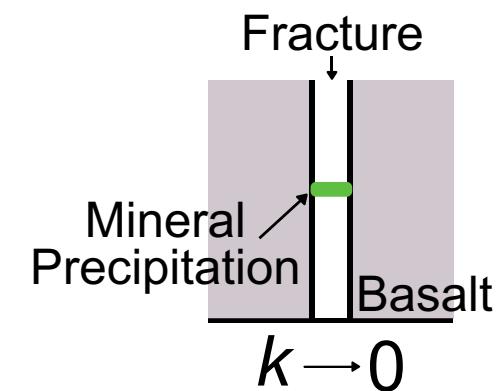
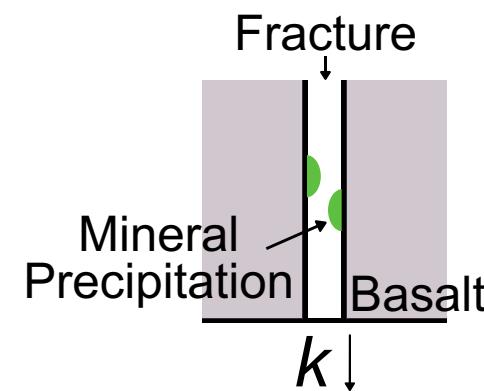
ϕ_o initial porosity

n parameter control the different scales of permeability alteration

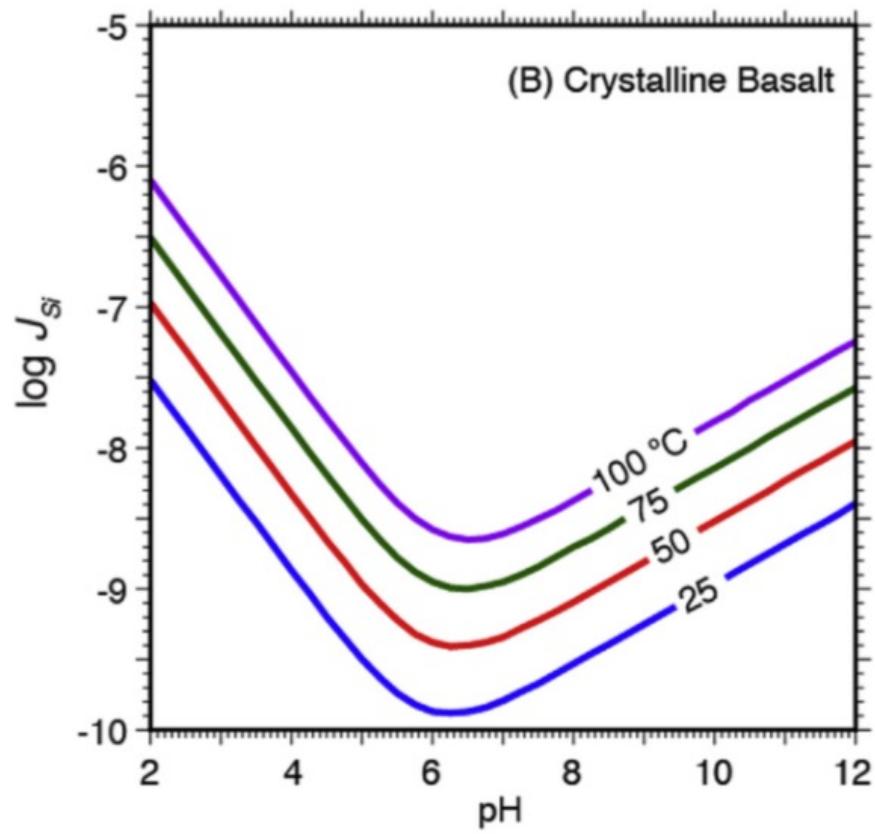


$$k = k_o \left(\frac{\phi}{\phi_o} \right)^3$$

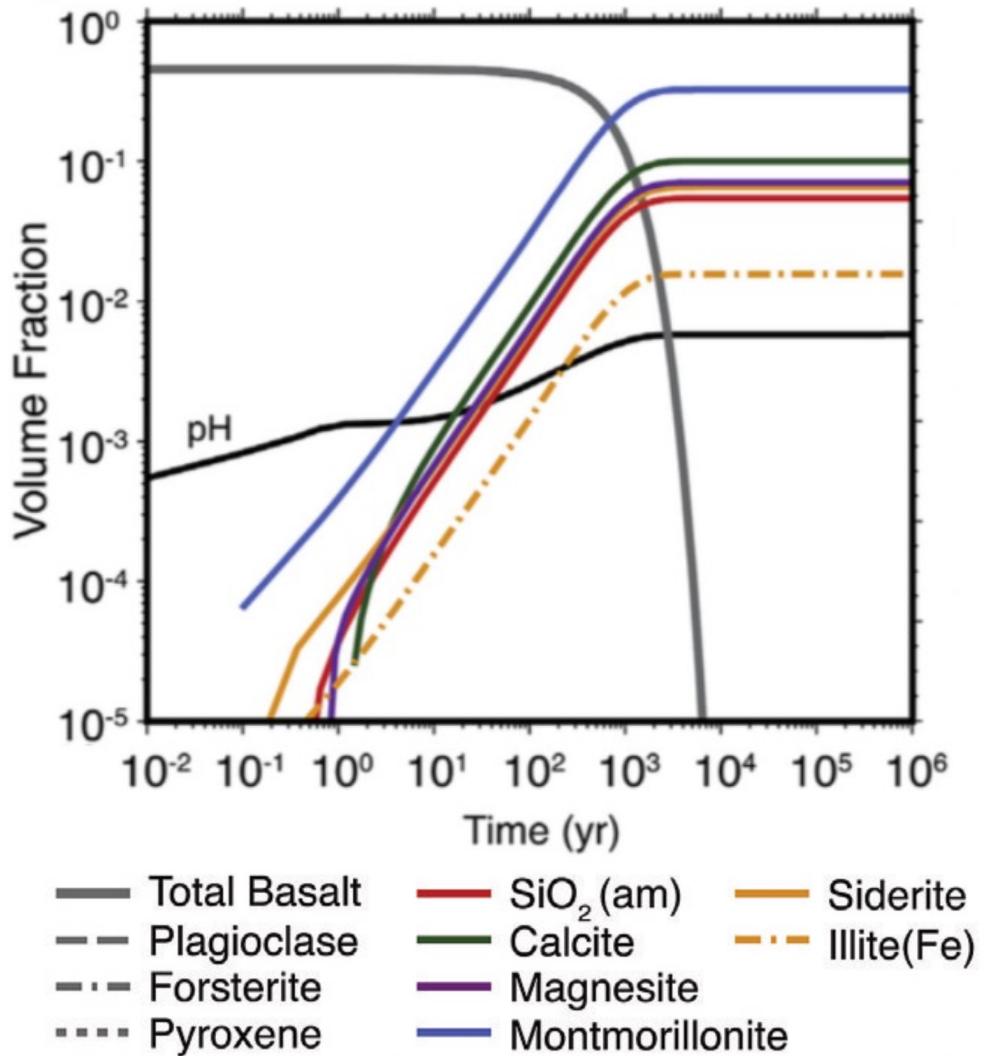
Cubic Law



BASALT DISSOLUTION MODEL



Pollyea & Rimstidt (2017)



MINERAL PRECIPITATION ACCUMULATES AT BRANCHING FRACTURE

Carbonate minerals:

Calcite

Magnesite

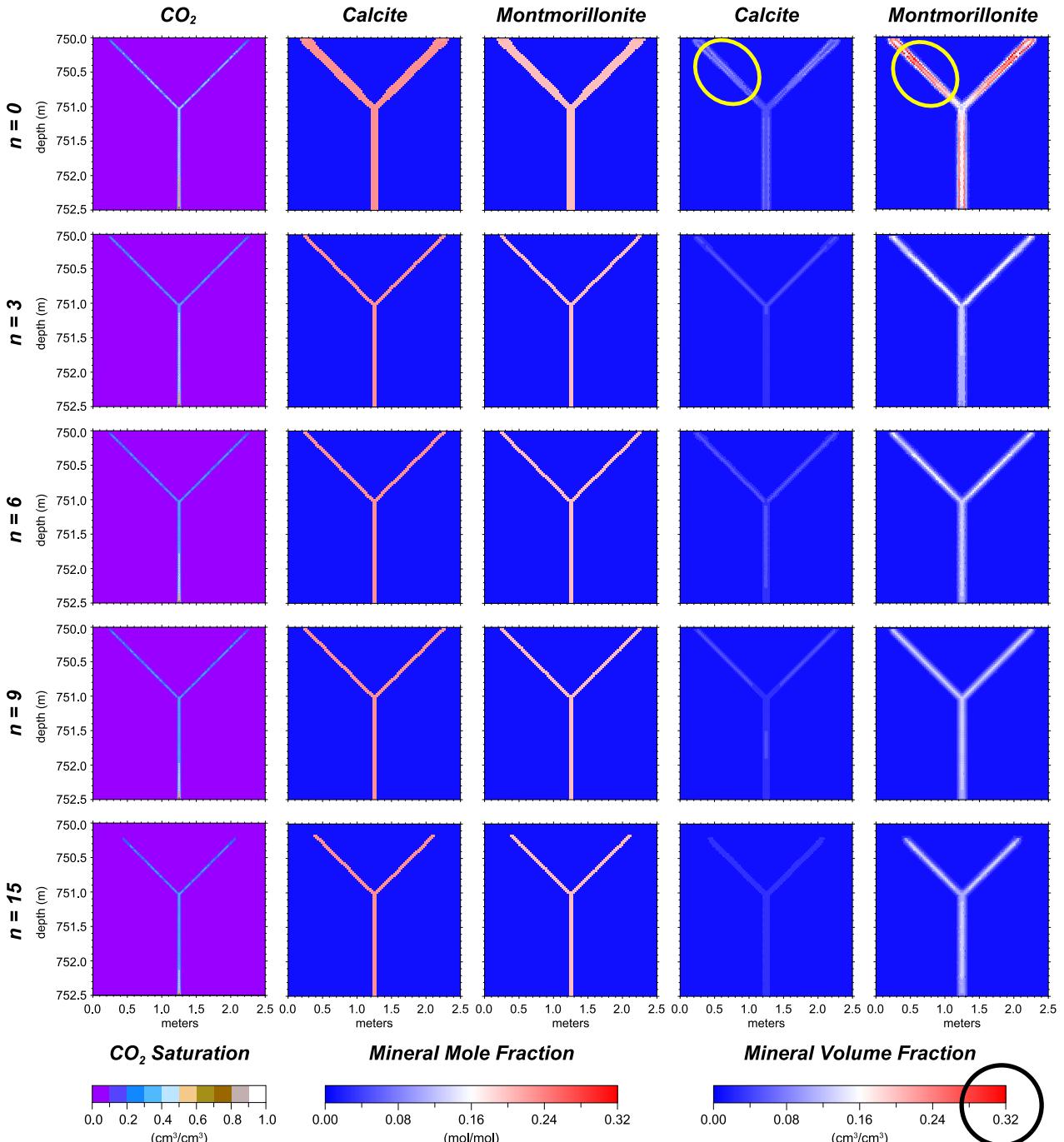
Siderite

Clay minerals:

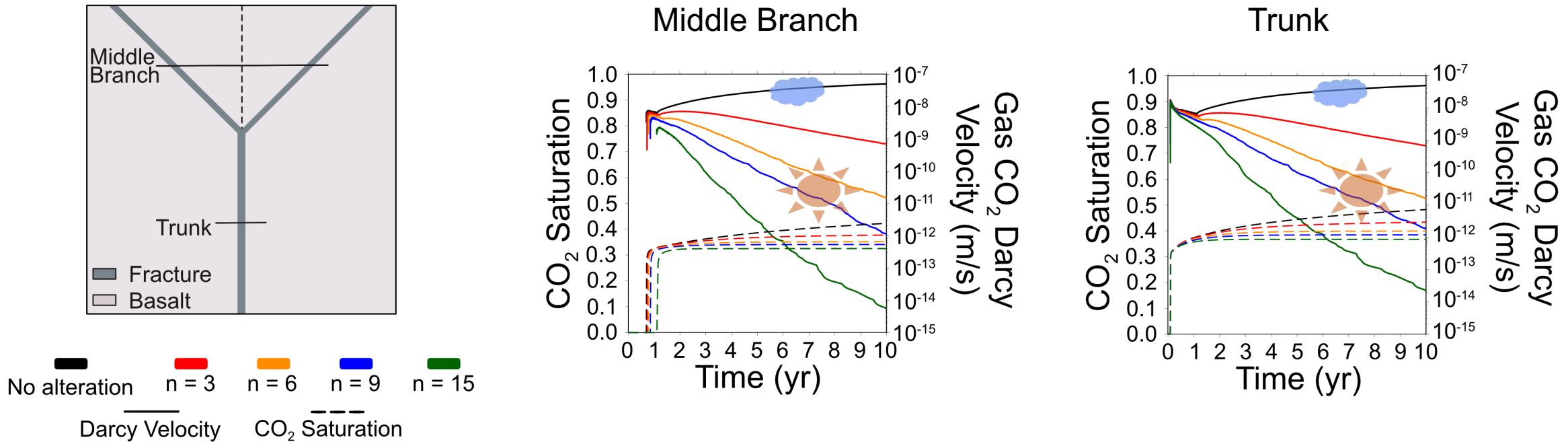
Montmorillonite (MgCa)

Illite (Fell)

Amorphous_silica



DARCY VELOCITY OF FREE-PHASE CO₂ SHOWS MAXIMUM 6 ORDERS OF MAGNITUDES DECREASE



No alteration: Darcy velocity increase

With alteration: Darcy velocity decrease

MASS BALANCE

| (mole)\n n | 0 | 3 | 6 | 9 | 15 |
|--------------|------------|------------|------------|------------|------------|
| Mineral | 62.5 | 30.5 | 31.6 | 31.8 | 27.2 |
| Aqueous | 4.5 | 2.0 | 2.2 | 2.2 | 2.1 |
| Gas | 1.5 | 0.5 | 0.4 | 0.4 | 0.4 |
| Boundary | 40.9 | 4.6 | 1.2 | 0.4 | - |
| Total | 109.4 | 37.6 | 35.4 | 34.8 | 29.7 |
| CMR (-) | 57% | 81% | 89% | 91% | 92% |

$$\text{Carbon Mineralization Ratio (CMR)} = \frac{\text{Mineralized carbon}}{\text{Total carbon in the system}}$$

CMR is >80% over ten years when considering permeability alteration

CONCLUSION

- Carbon mineralization focuses on branching fracture
- Fracture permeability decreases and slows the vertical migration of CO₂
- Different permeability alteration scales show important effects in this carbon mineralization process
- Fracture system may be self-sealing due to secondary mineral precipitation

THANK YOU