



# Subcritical Fracturing of Calcite Single Crystals and Grain Packs

**Anastasia G. Ilgen,<sup>1</sup> R. Charles Choens,<sup>2</sup> and Jennifer Wilson<sup>2</sup>**

<sup>1</sup>Geochemistry Department; <sup>2</sup>Geomechanics Department  
Sandia National Laboratories



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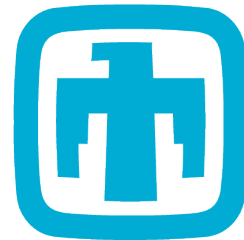
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# Overview: theory of chemically-assisted fracturing



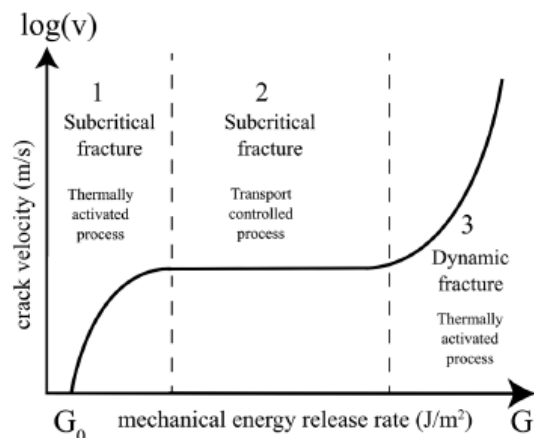
## Constitutive modeling of subcritical crack growth:

Reaction rate theory

$$\ln \left( \frac{r}{r_o} \right) = a - \left( \frac{E^* - v^* \sigma_{rs}}{RT} \right)$$

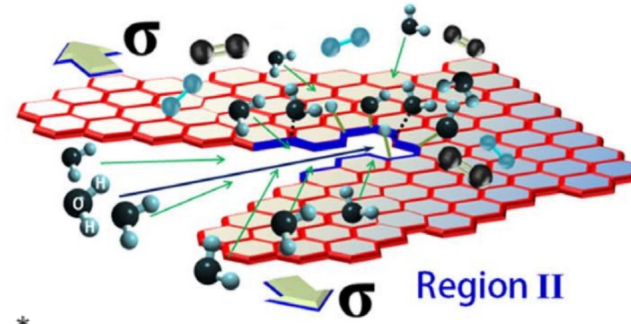
$r$  is reaction rate,  $r_o$  and  $a$  – empirical constants,  $R$  – gas constant,  $T$  – absolute temperature,  $\sigma_{rs}$  – reaction site stress,  $E^*$  and  $v^*$  – apparent activation energy and activation volume.

$$v = 2 \frac{kT}{h} a_0 \exp \left( \frac{-\Delta F}{kT} \right) \sinh \left( \alpha \frac{G - G_0}{kT} \right)$$

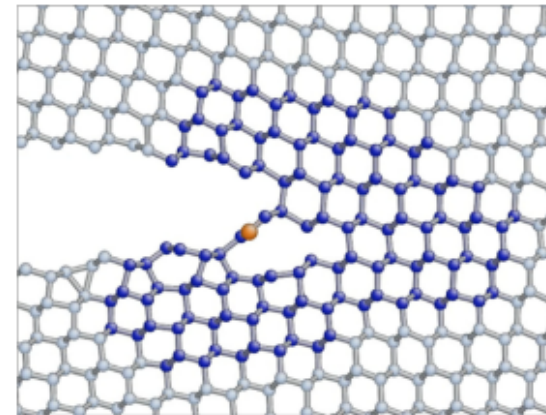


$v$  is crack velocity,  $k$  is Boltzmann's constant,  $h$  is Plank's constant,  $G$  is mechanical energy release ( $G_0$  is theoretical limit),  $a_0$  is characteristic atomic spacing,  $\alpha$  is activation area,  $\Delta F$  is apparent activation barrier.

Bergsaker et al., 2016

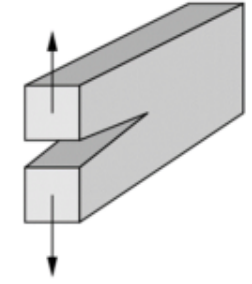


Hwangbo et al., 2014

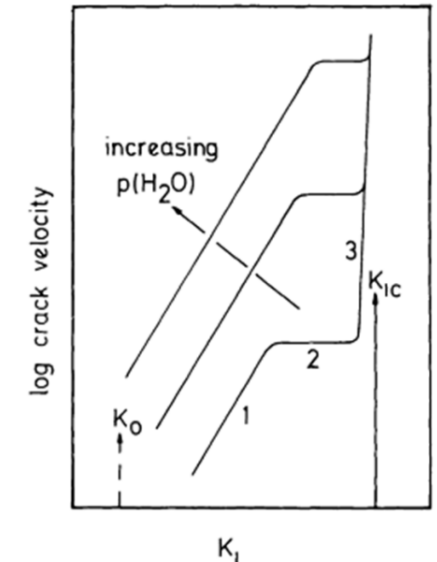


Bitzek et al., 2005

Mode I



Atkinson and Meredith, 1987



Schematic stress intensity factor ( $K_i$ ) and crack velocity diagram for tensile crack growth by stress corrosion.  $K_{ic}$  is the fracture toughness and  $K_0$  is the stress corrosion limit.

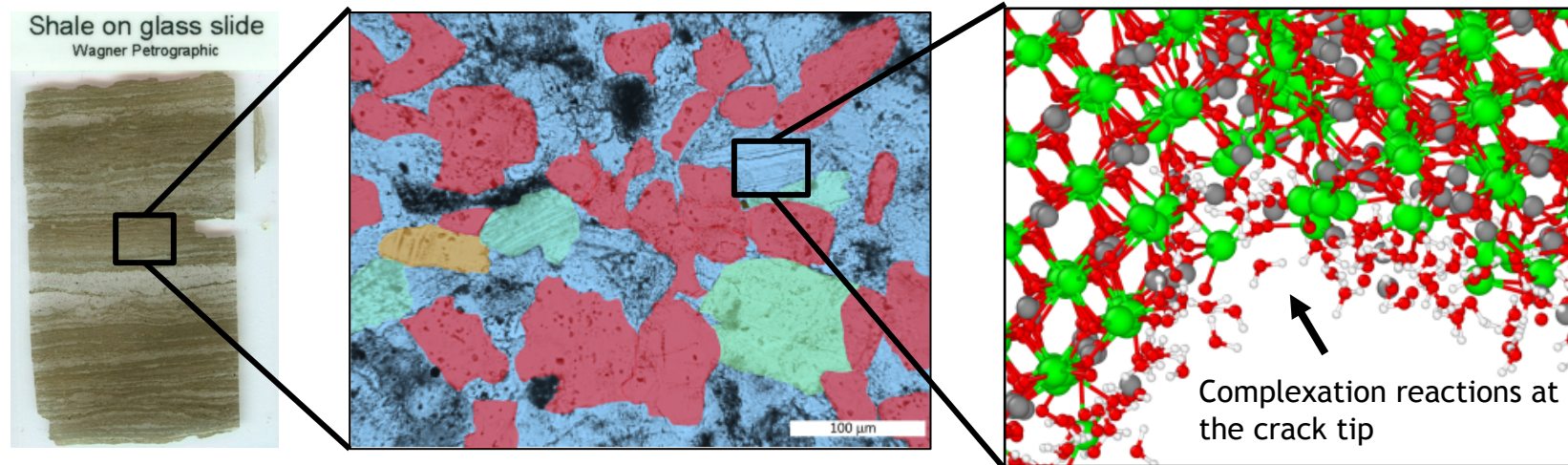


# Chemically-assisted fracturing in calcite: hypothesis



**Science Question:** How and why do chemical complexation reactions at a single crack tip change *in situ* fracture behavior?

**Hypothesis:** With increasing favorability of the cation-ligand complex, the velocity of subcritical crack growth decreases, and the effective fracture toughness increases.



- Fracturing in rocks can occur through intergranular cement, or through mineral grains. Calcite ( $\text{CaCO}_3$ ) and quartz ( $\text{SiO}_2$ ) cements are common intergranular phases in sedimentary rocks;
- Previous studies on subcritical fracture show that:
  - Activity of  $\text{H}_2\text{O}$  controls weakening of chalk <sup>[1]</sup>
  - Dissolution at fracture tip controls fracture growth <sup>[2,3]</sup>
  - Changes in surface energy control fracture propagation <sup>[4-7]</sup>

[1] Risnes et al., 2005

[2] Atkinson, 1984

[3] Royne et al., 2011

[4] Rostom et al., 2012

[5] Griffith, 1921

[6] Kermode et al., 2013

[7] Bergsaker et al., 2016

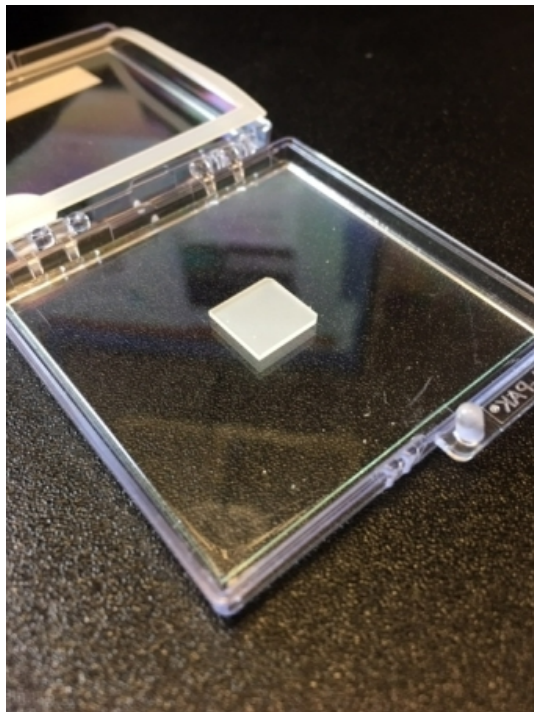
# SCIENTIFIC REPORTS

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## Chemical controls on the propagation rate of fracture in calcite

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A. G. Ilgen<sup>1</sup>, W. M. Mook<sup>2</sup>, A. B. Tigges<sup>1</sup>, R. C. Choens<sup>1</sup>, K. Artyushkova<sup>1</sup> & K. L. Jungjohann<sup>2</sup>



micron → centimeter

minutes → days

single crystal → grain pack

dilute aqueous solutions

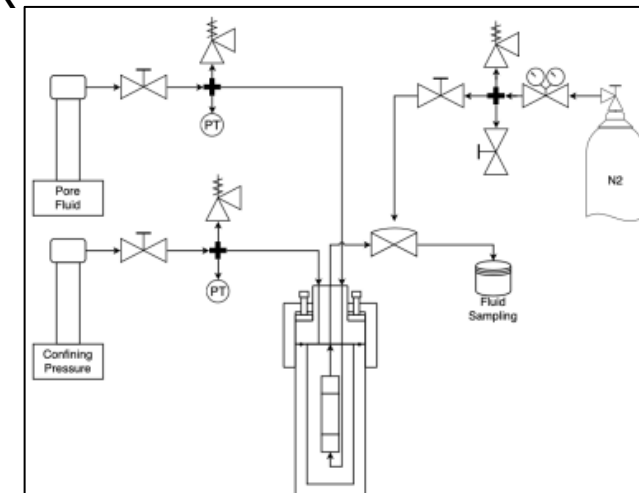
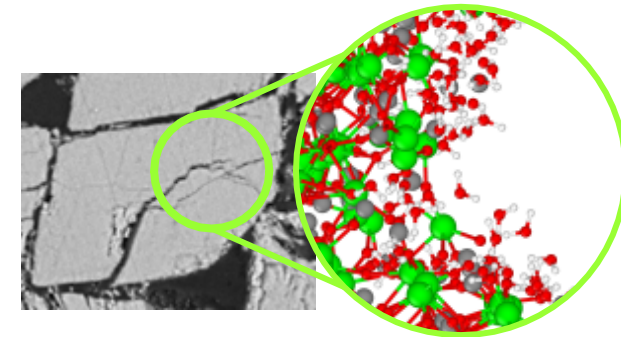
→ 0.5M

brines

Submitted paper:

## Strengthening of Calcite Assemblages through Chemical Complexation Reactions

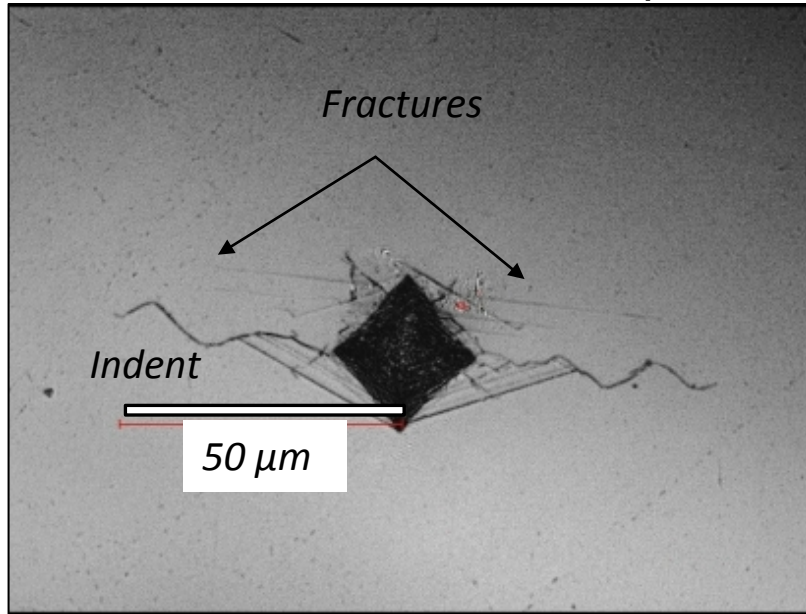
R. C. Choens, J. Wilson, and A. G. Ilgen



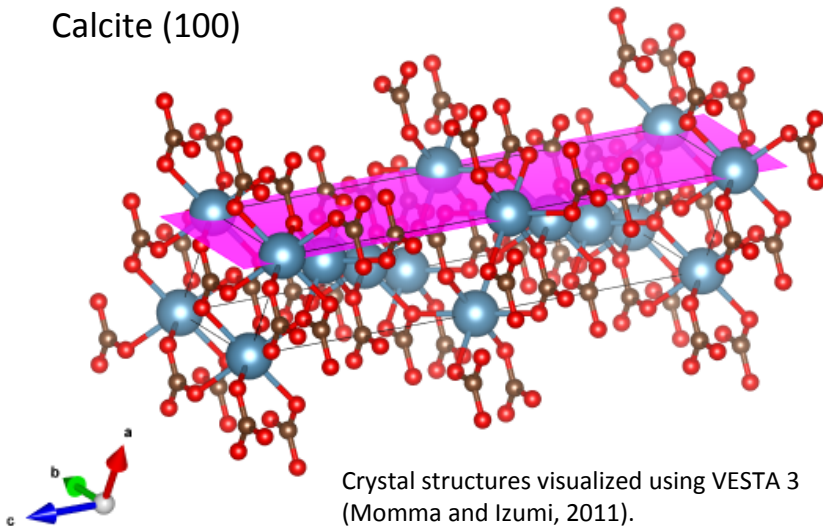
# Methods: nanoindentation and in situ crack growth



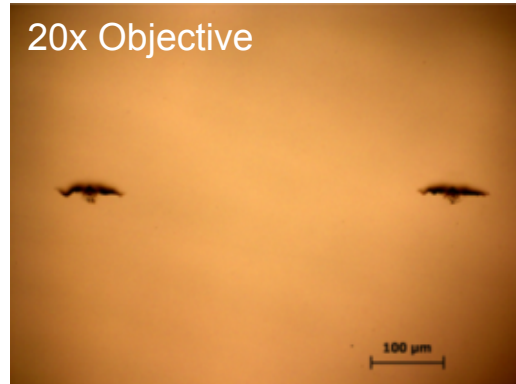
Calcite Indentation, Vickers tip, 400 mN



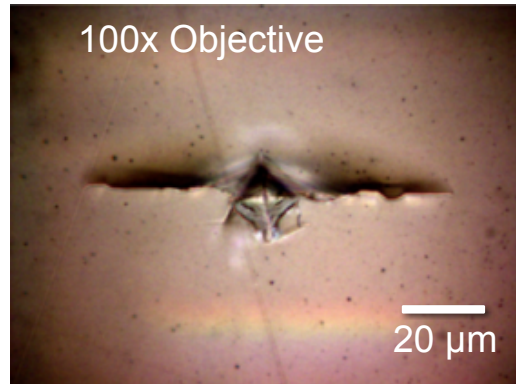
Calcite (100)



20x Objective



100x Objective



Fracture toughness <sup>[1]</sup>:

$$\frac{P}{c^{3/2}} = \frac{1}{\xi} \times \left(\frac{H}{E}\right)^{1/2} \times T$$

[1] Lawn and Cook, 2012

Vickers tip

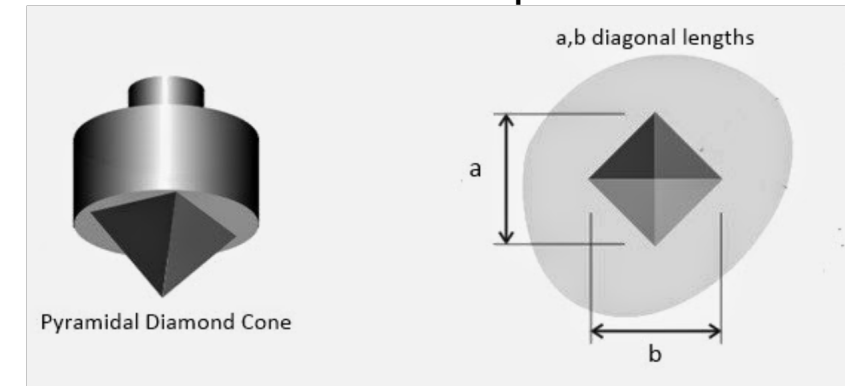


Image from: <http://www.weldpedia.com/2014/10/macrosopic-and-microscopic-examination.html>

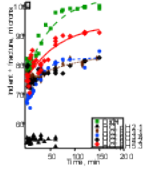
- Laboratory experiments to measure fracture propagation rate *in situ* as a function of chemical composition of the fluid;
- Single crystal calcite (100) indented using Vickers indenter tip at 400 mN force to induce cracking;
- Fractures are imaged *in situ* using optical microscope Nikon Eclipse 80i and SPOT 7.2 camera **Hagen, et al., 2018**



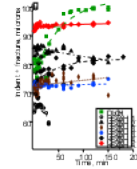
# Results: fracture growth rate



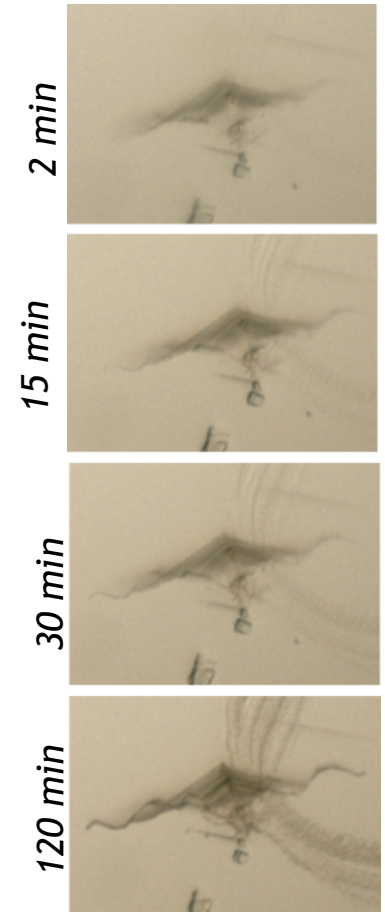
HCl



H<sub>2</sub>SO<sub>4</sub>

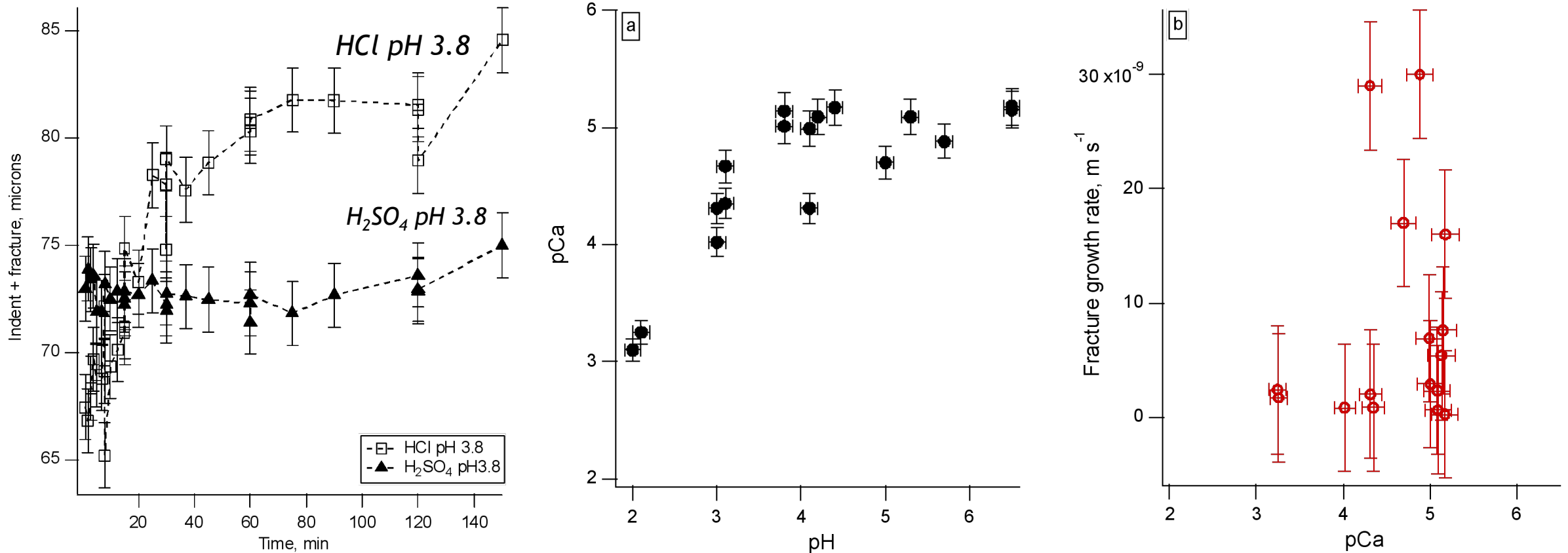


pH 5



- The propagation rate of subcritical fracture measured *in situ* varied from  $1.6 \times 10^{-8} \text{ m s}^{-1}$  to  $2.4 \times 10^{-10} \text{ m s}^{-1}$ .

# Results: what controls crack growth?



- Propagation rate of fracture in calcite is dependent on the anion.
- No correlation between the dissolution rate of calcite and subcritical fracture growth.
- Positive correlation between pCa and pH for all examined reactors;  $pCa = -\log_{10}[Ca^{2+}]$ ;
- No correlation observed between pCa (proxy for the  $\xi$ -potential) and fracture propagation rates



# Results: what controls fracture growth?



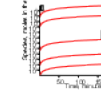
H<sub>2</sub>O



HCl

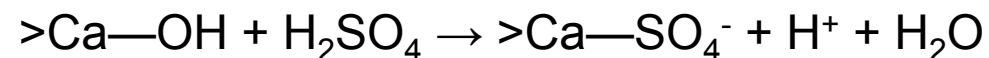
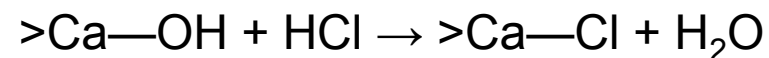
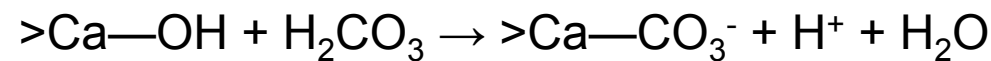
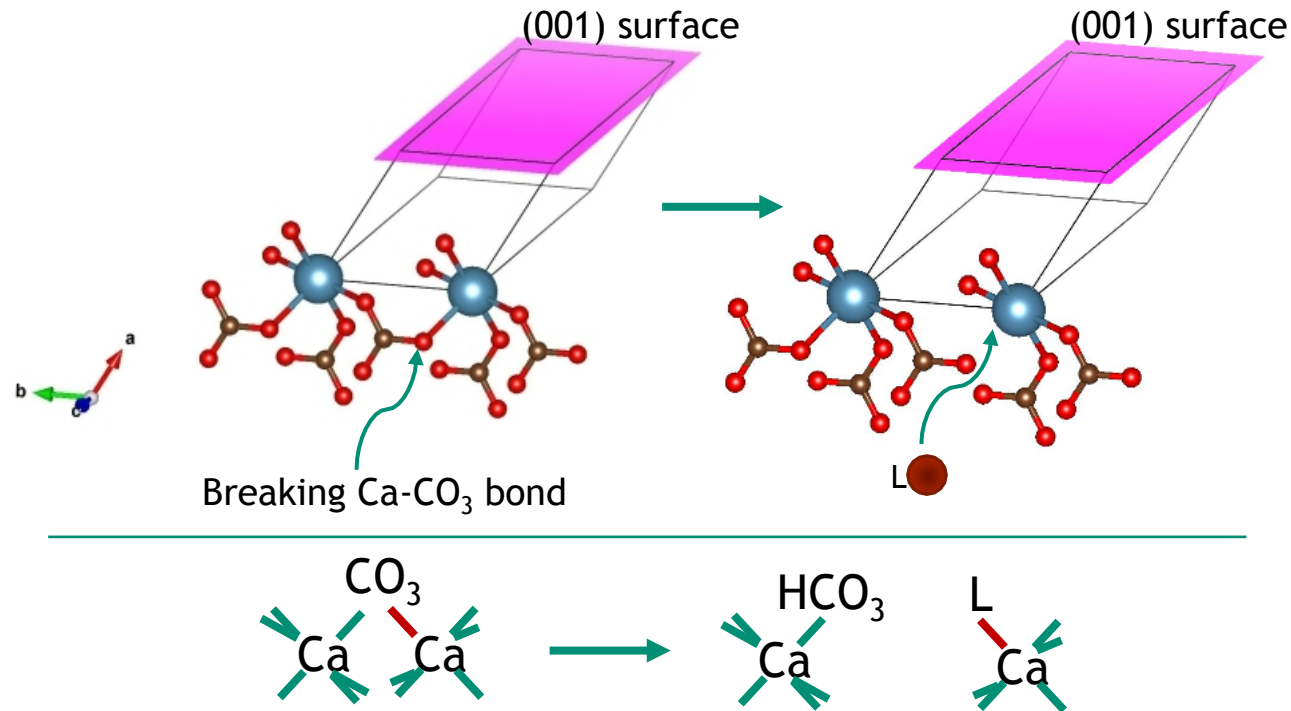


H<sub>2</sub>SO<sub>4</sub>



$K_{\beta}$  for  $\text{CaCO}_3$  is  $10^{-7.128}$ ;  $K_{\beta}$  for  $\text{CaCl}^+$  is  $10^{0.7}$ ; and  $K_{\beta}$  for  $\text{CaSO}_4$  is  $10^{2.32}$

# Results: Conceptual model

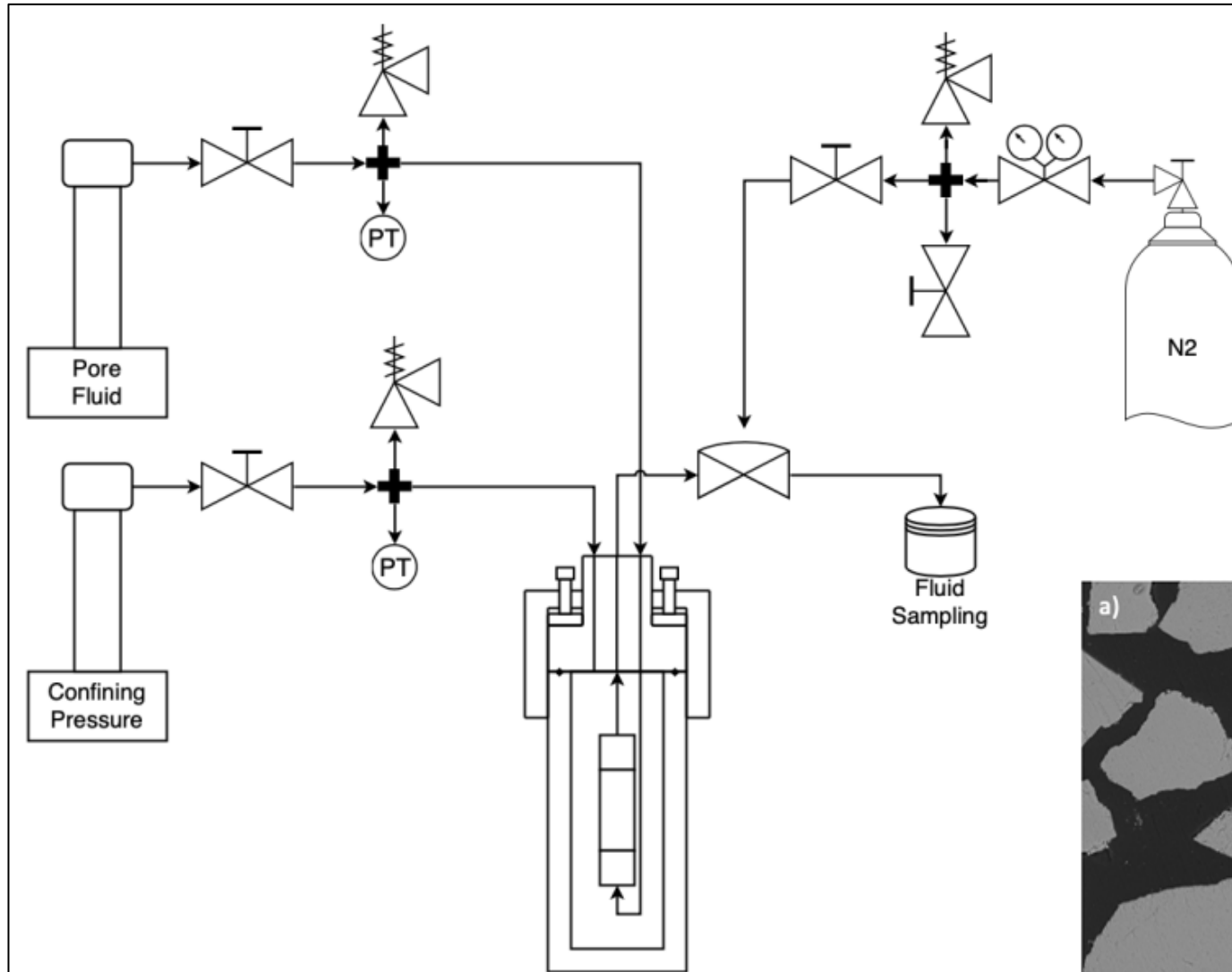


- The estimated fracture toughness prior to *in situ* fracture growth experiment was 0.10 – 0.16 MPa m<sup>1/2</sup>
- Fracture toughness at the end of the fracture growth experiment decreased by 0.01-0.05 units.

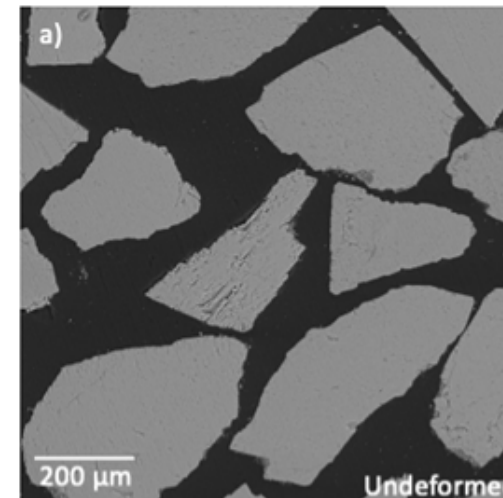
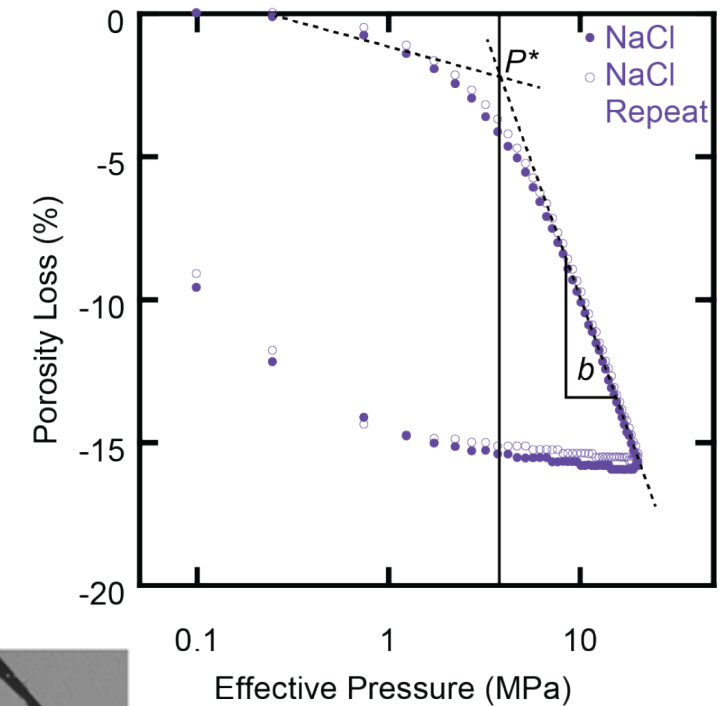
# Experimental apparatus for consolidation tests



Schematic of consolidation apparatus

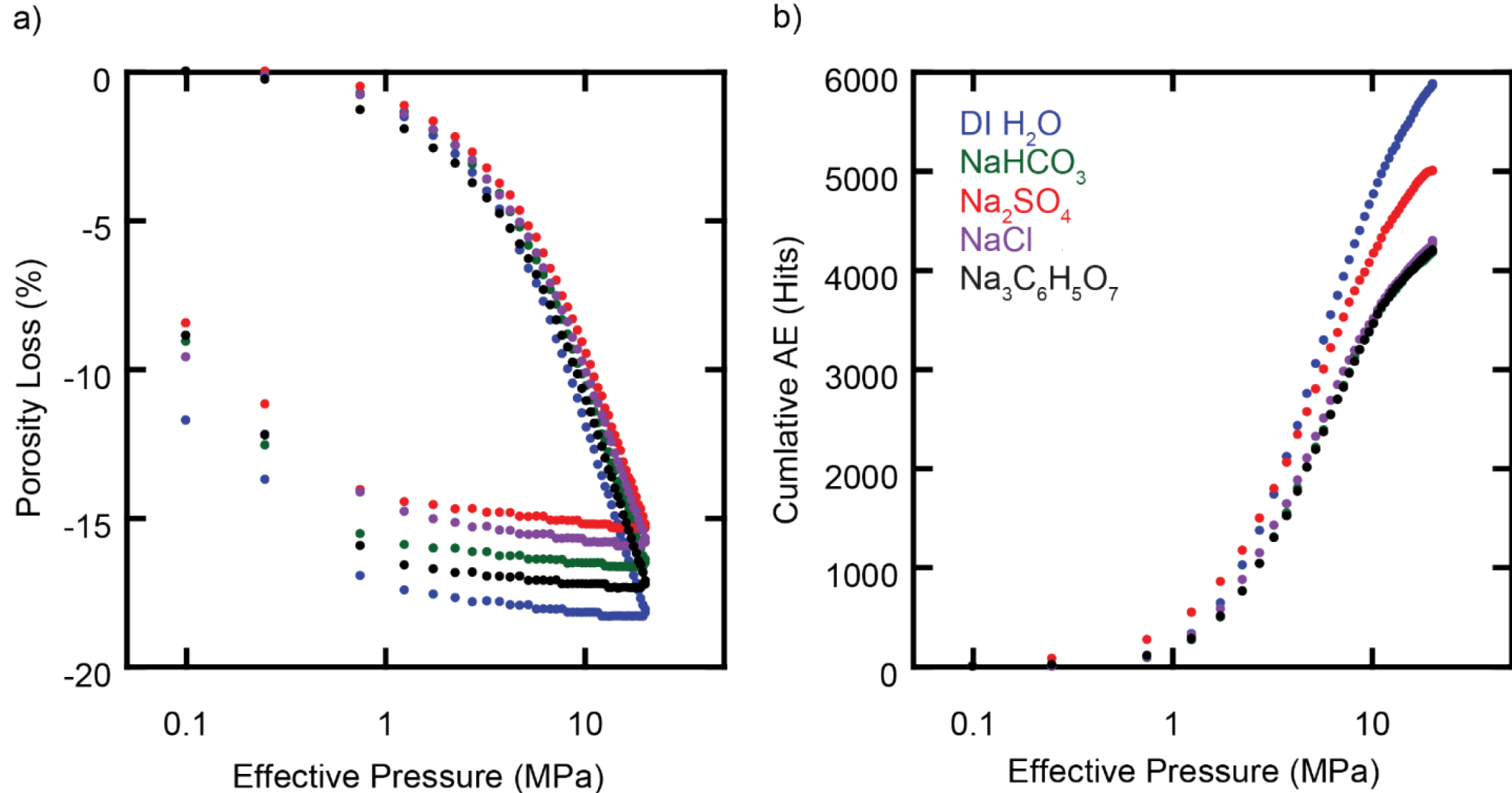


Consolidation curves



Starting calcite material, sieved to a grain size of 300-355  $\mu\text{m}$ .

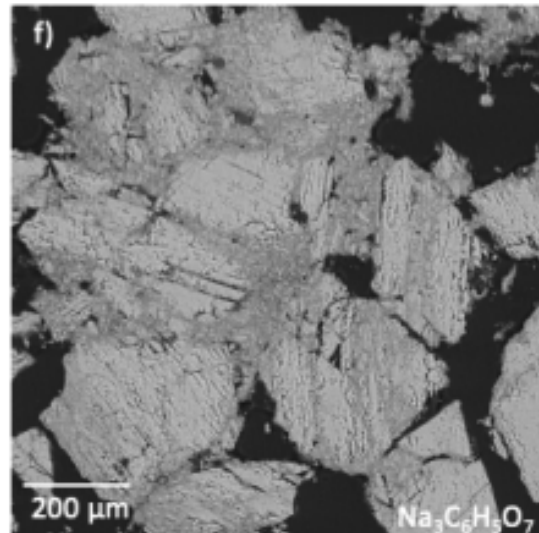
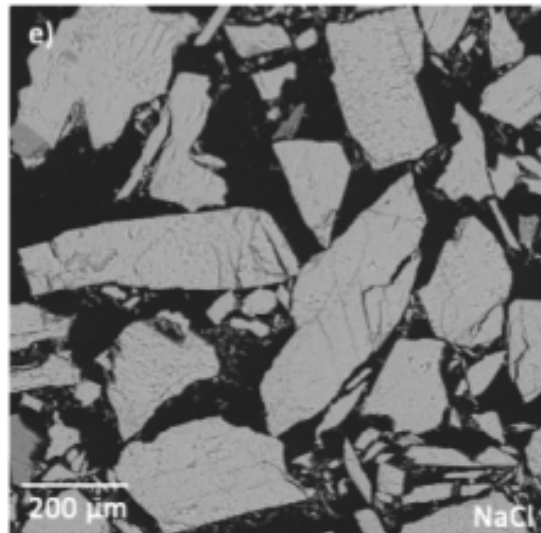
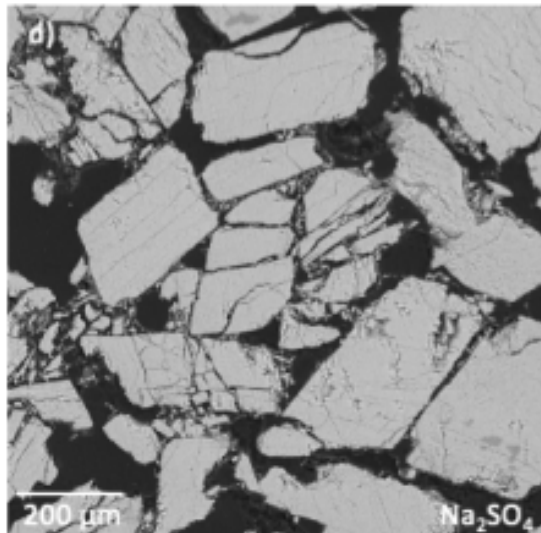
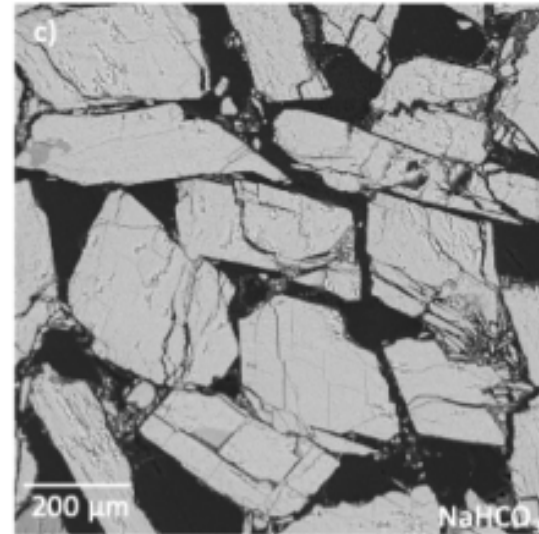
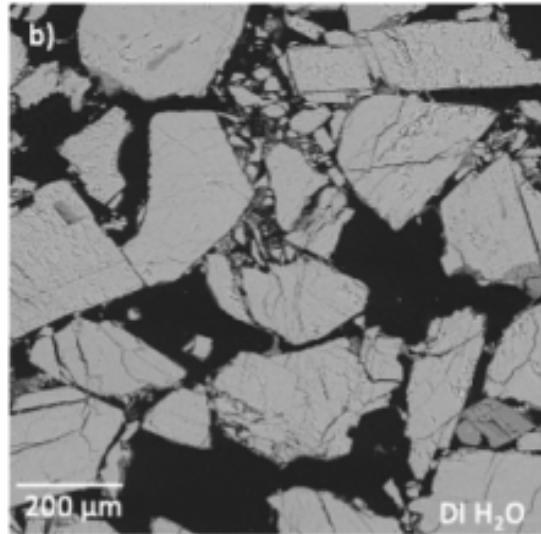
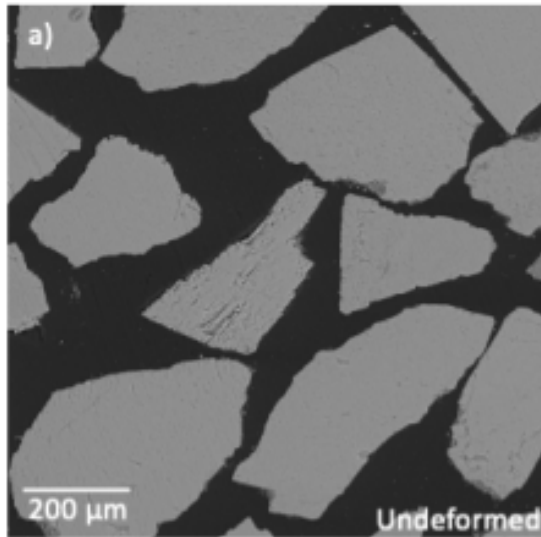
# Results: porosity loss and acoustic emissions



a) Consolidation curves for granular calcite deformed with different interstitial pore fluids, showing porosity loss versus log effective pressure. b) Cumulative Acoustic Emissions (AE) during consolidation versus log effective pressure.



# Results: microfracturing



- Scanning electron microscopy (SEM): a) Starting calcite material, sieved to a grain size of 300-355  $\mu\text{m}$ .
- Samples consolidated in the presence of b) DI  $\text{H}_2\text{O}$ , c)  $\text{NaHCO}_3$ , d)  $\text{Na}_2\text{SO}_4$ , e)  $\text{NaCl}$ , and f)  $\text{Na}_3\text{C}_6\text{H}_5\text{O}_7$ .
- Fragmentation of grains and incorporation of crushed grains into interstitial pore spaces in all consolidated samples.

# Results: microfracturing



**Table 1.** Consolidation results for granular calcite.

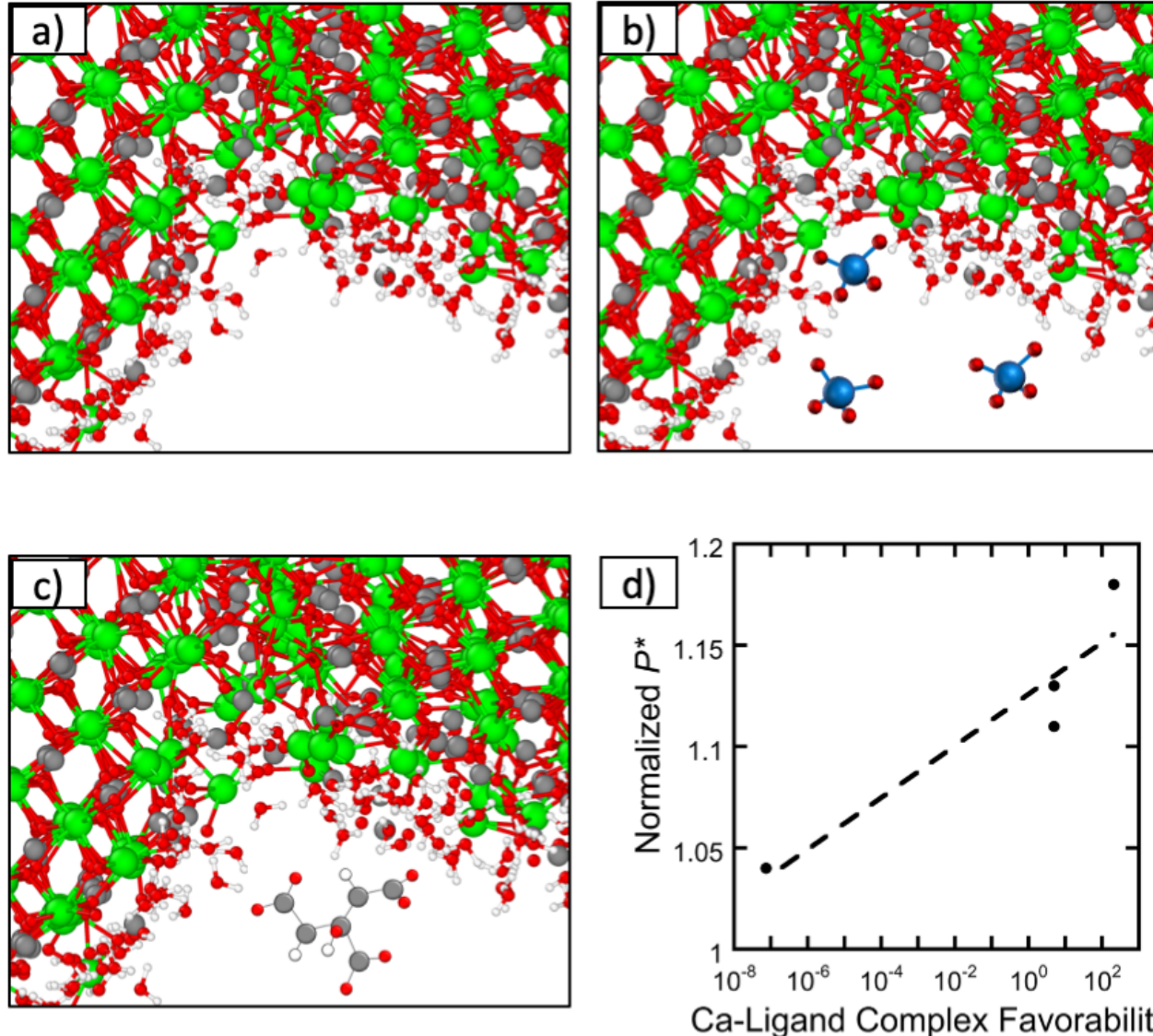
Sample	P* (MPa)	Normalized P* by DI H <sub>2</sub> O	Porosity Loss (%)	Consolidation Slope (%/MPa)	Total Microfracture Density (mm/mm <sup>2</sup> )	% Grains affected by compaction	Modal Grain Size (grain diameter in
Starting CaCO <sub>3</sub>	--	--	--	--	3.5	0	325.0
DI H <sub>2</sub> O	3.5	1	-18	20.5	21.1	71	29.0
NaHCO <sub>3</sub>	3.65	1.04	19.3	19.3	19.9	80	28.0
Na <sub>2</sub> SO <sub>4</sub>	4.12	1.18	18.7	18.7	15.4	78	32.6
NaCl	3.90	1.11	18.6	18.6	13.5	75	28.2
NaCl Repeat	3.94	1.13	18.6	18.6	--	--	--
Na <sub>3</sub> C <sub>6</sub> H <sub>5</sub> O <sub>7</sub>	4.07	1.16	19.8	19.8	20.2	90	43.0

Microfracture density depends on the fluid type and follows the sequence:  
 DI H<sub>2</sub>O > Na<sub>3</sub>C<sub>6</sub>H<sub>5</sub>O<sub>7</sub> > NaHCO<sub>3</sub> > Na<sub>2</sub>SO<sub>4</sub> > NaCl

**Choens, et al., 2018**

Submitted

# Results: Conceptual model



- Molecular-scale schematic of crack tip in calcite: red – oxygen, green – calcium, grey – carbon, blue – sulfur, white – hydrogen.
- (a) Calcite consolidation in de-ionized  $H_2O$ , with water hydrolyzing  $Ca-CO_3$  bonds and promoting crack growth;
- (b) consolidation in  $0.5M Na_2SO_4$  with sulphate forming an  $Ca-SO_4$  complex at the crack tip preventing hydrolysis reaction;
- (c) consolidation in  $0.5M Na_3C_6H_5O_7$  with citrate anion not reaching the crack tip before water does due to slower diffusion, compared to sulphate;
- (d) normalized  $P^*$  versus Ca-anion complex favorability ( $K_\beta$  constant).

Choens, et al., 2018

Submitted



**Thank you.**