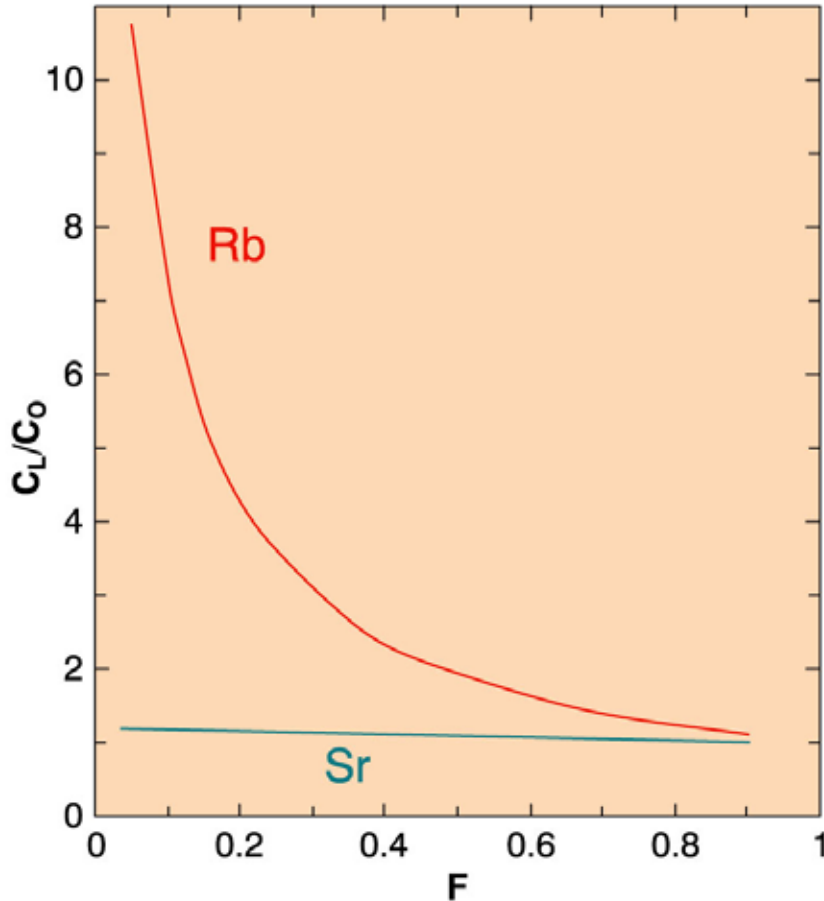


# Announcements

Be on time for Field Trip: 9:45 AM  
on Friday



# Rayleigh Fractionation



$$\frac{C_L}{C_0} = F^{(D-1)}$$

$C_L$  = concentration in melt

$C_0$  = concentration in total original assemblage

$F$  = fraction of melt remaining

$D$  = bulk distribution coefficient =  $X_1K_1 + X_2K_2 + \dots$

**Relevant to... perfect fractional crystallization!!!!!!**

# Rayleigh fractionation in petrology I

- Application: fractional crystallization of a melt
- Trace elements will partition preferentially into a melt or a mineral during this process (chemical fractionation)

Example: Eu

Initial concentration: 2 ppm

$$K_{D_{\text{plag}}} = 4$$

When have 20 wt% fractional crystallization

What is concentration of Eu in melt?

$$C_L = 1 \text{ ppm}$$

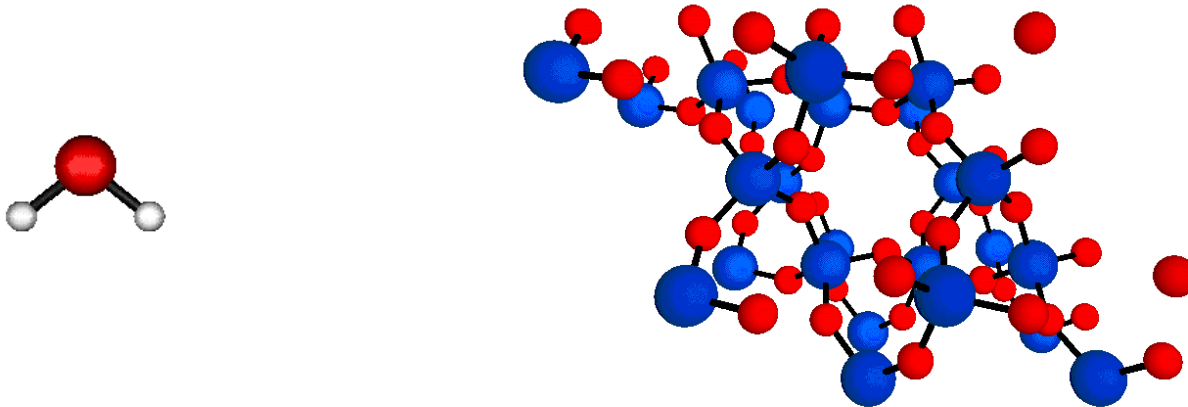
# Stable Isotopes

- Stable: last ~ forever
- Chemical fractionation is (impossible)
- Mass fractionation is the only type possible

# Oxygen isotopes

- Terrestrial abundances:  $^{16}\text{O}=99.76/^{17}\text{O}=0.04/^{18}\text{O}=0.20$
- $$\delta^{18}\text{O} = 1000 * \frac{(^{18}\text{O}/^{16}\text{O}_{\text{sample}} - ^{18}\text{O}/^{16}\text{O}_{\text{std}})}{^{18}\text{O}/^{16}\text{O}_{\text{std}}}$$

Std = Standard Mean Ocean Water (SMOW)

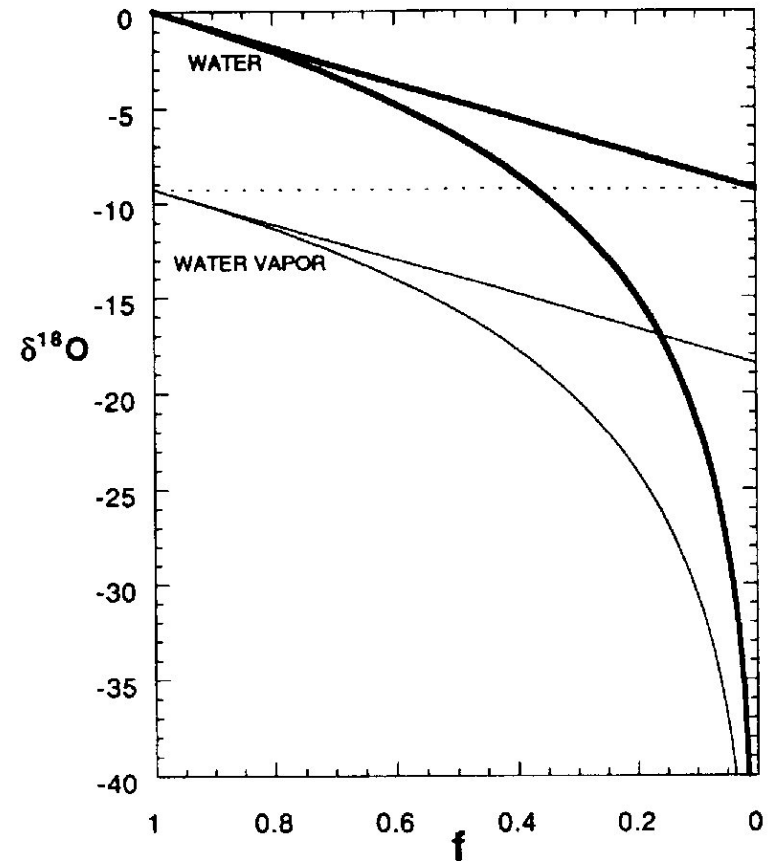
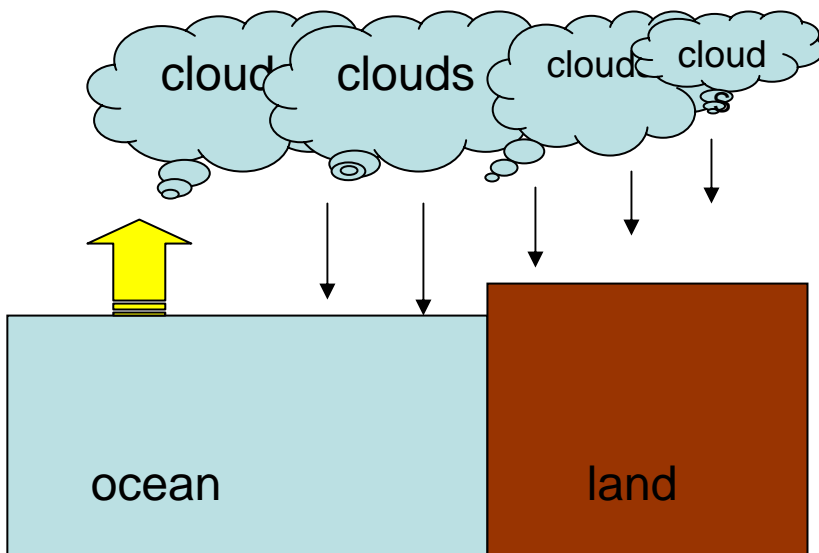


**Mass-dependent fractionation**

# Rayleigh fractionation in petrology

## II

- Oxygen isotopes are fractionated during evaporation and rain cycles



Fraction of water vapor remaining

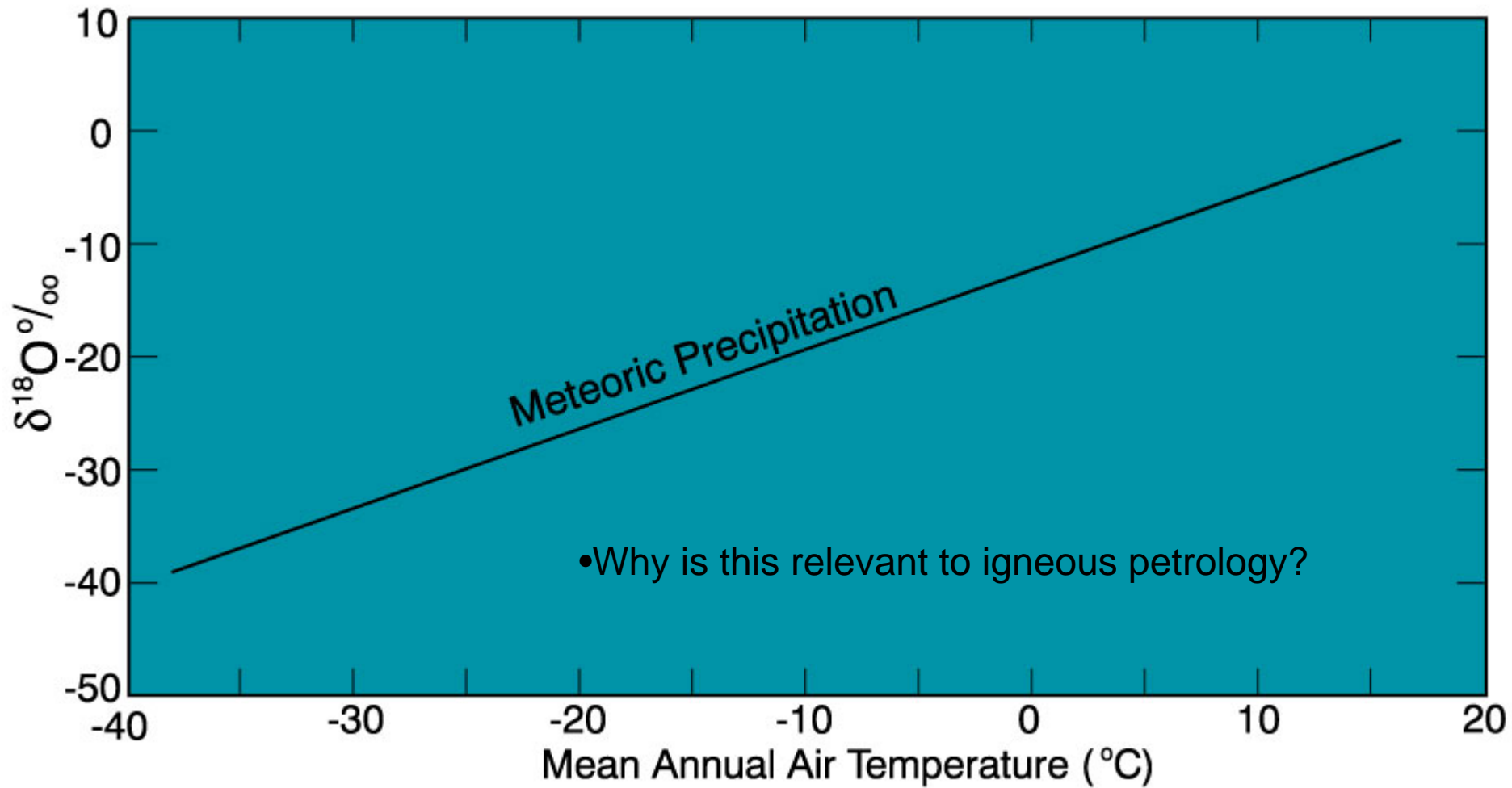
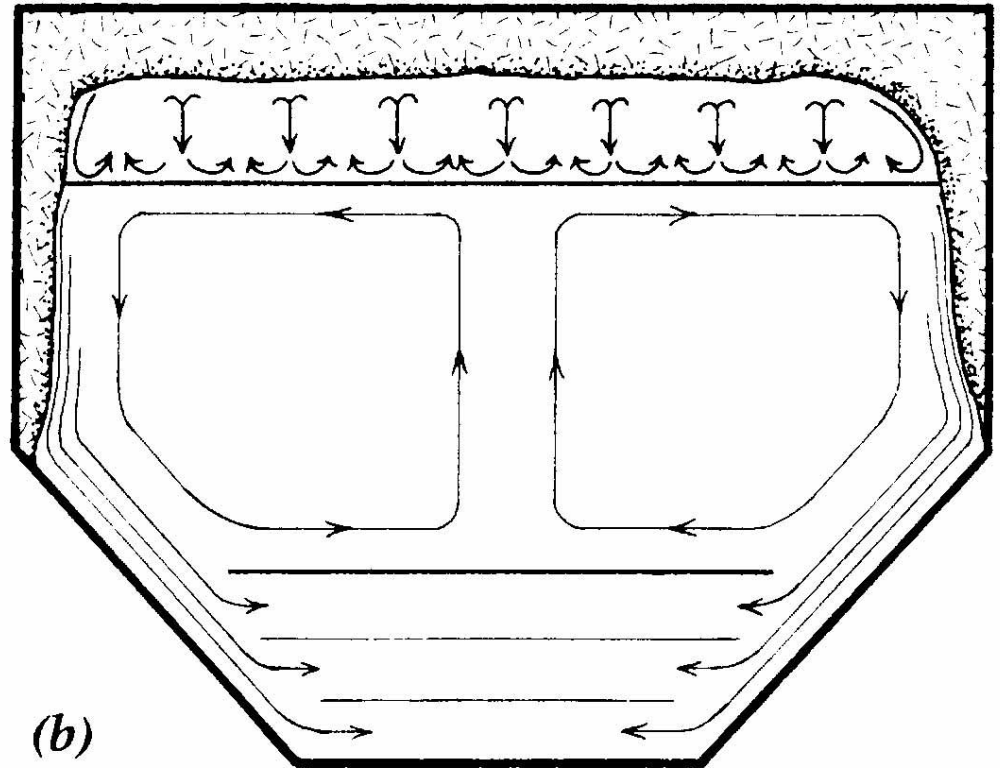


Figure 9-9. Relationship between  $d(^{18}\text{O}/^{16}\text{O})$  and mean annual temperature for meteoric precipitation, after Dansgaard (1964). *Tellus*, **16**, 436-468.

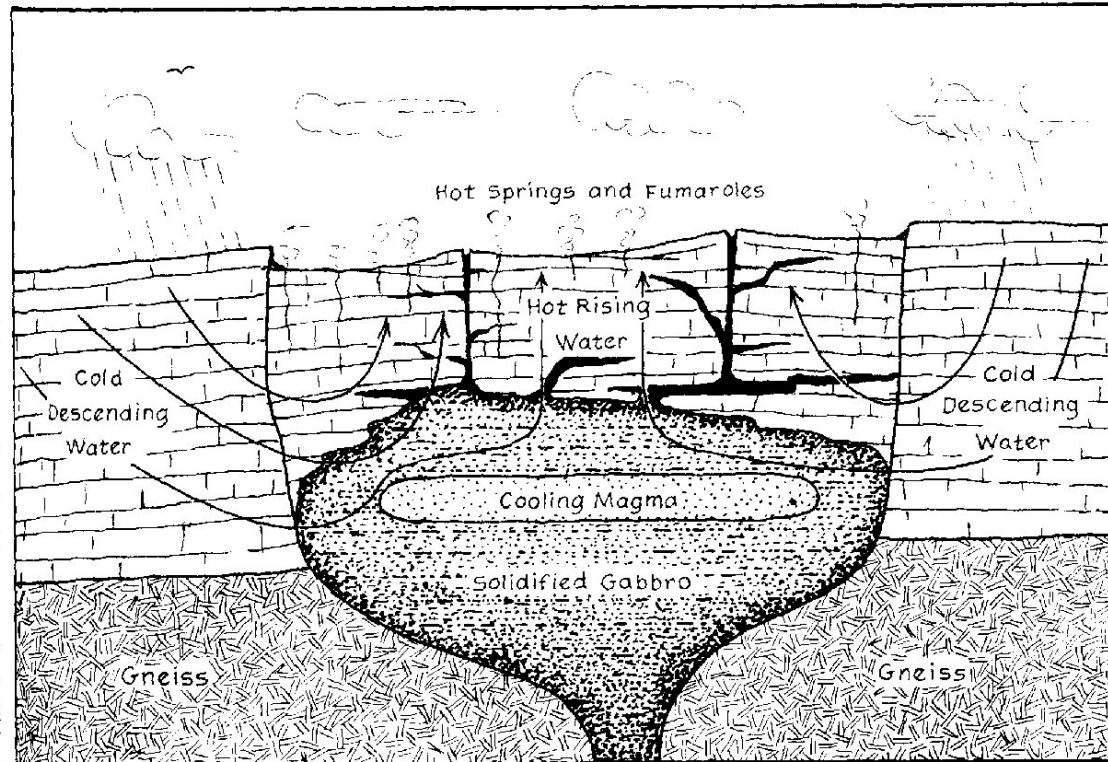
# Cooling of an intrusive body

- Magma convection
- Fractional crystallization results in solidification from bottom up (and top down)





# Cooling of an intrusion



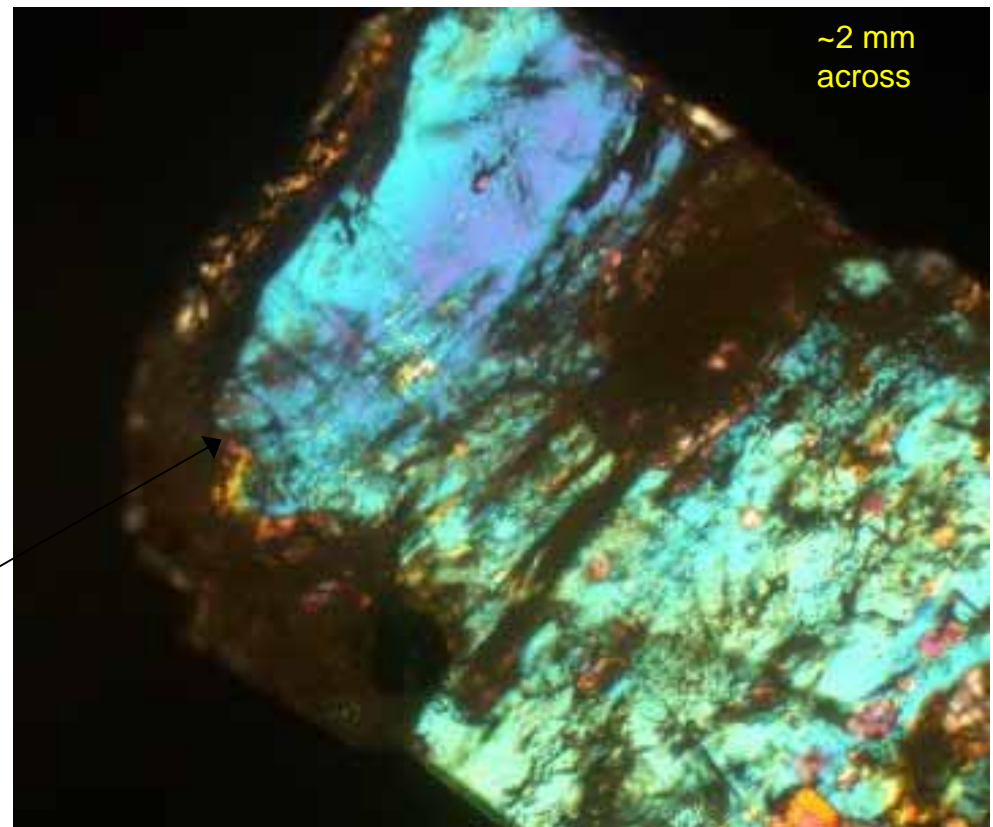
**Figure 6-34** The Skaergaard intrusion set up a large hydrothermal system in which meteoric water heated by the cooling gabbro convected through the permeable basaltic

rocks of the roof and upper walls. (After H. P. Taylor and R. W. Forester, 1979, *Jour. Petr.* 20:355-419).

Groundwater or meteoric water:  
from precipitation

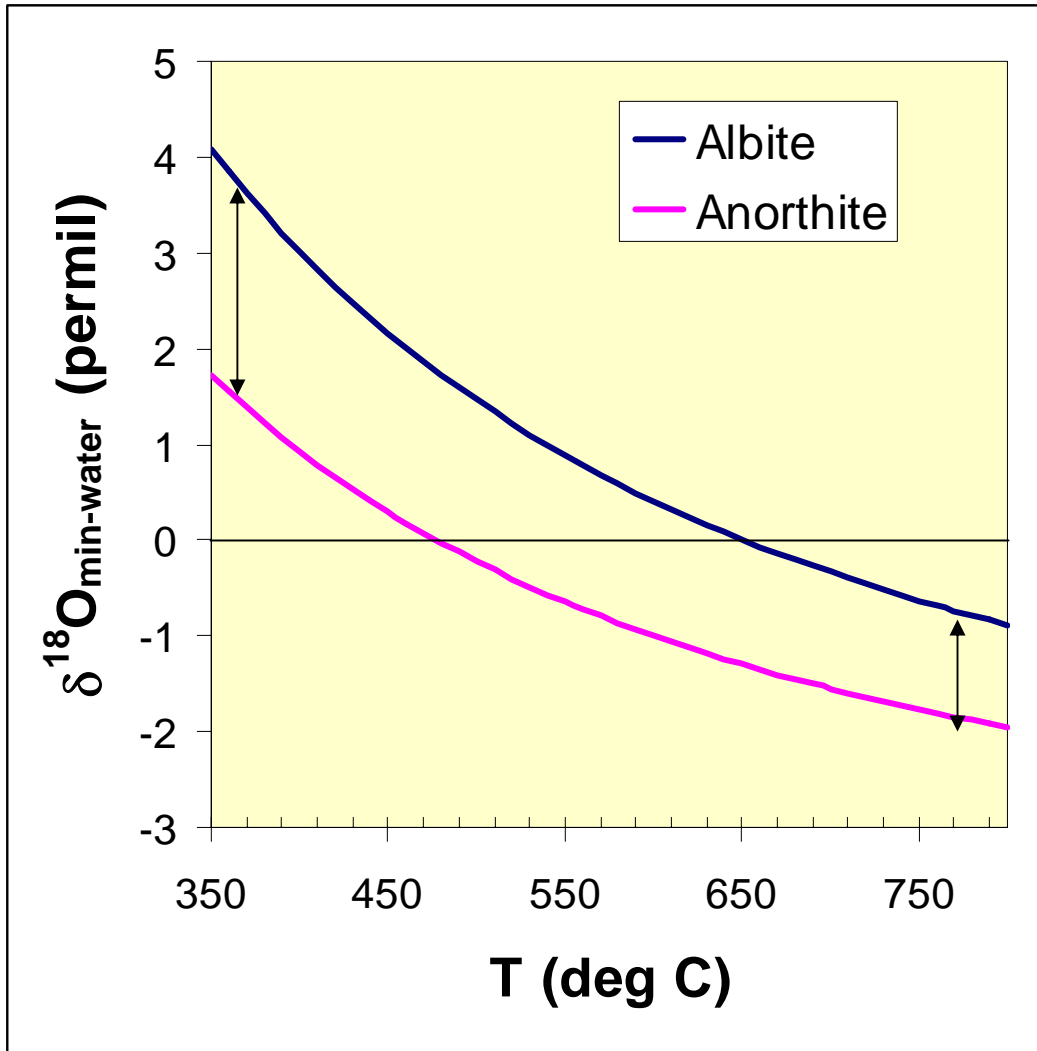
# Hydrothermal exchange of minerals

“...every part of the crystal must have had to ‘swim’ across the ‘river’ in solution in order to reach the other side.” – O’Neil and Taylor, 1967



Solution and redeposition  
in fluid film at interface  
between exchanged and  
unexchanged feldspar.

# Temperature dependence of $\delta^{18}\text{O}$ in feldspar



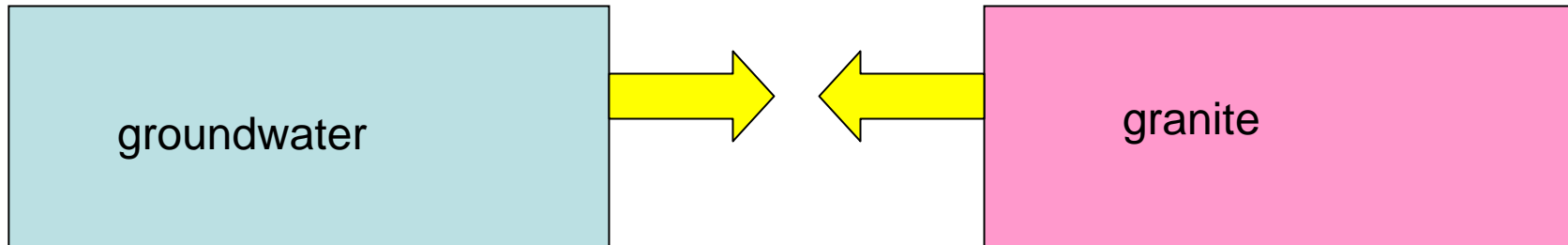
Larger fractionation between two minerals at a LOWER temperature

HOT hydrothermal activity  $>450$  °C will tend to draw down mineral  $\delta^{18}\text{O}$  to lower values

# What is effect of large scale circulation and exchange of water with crystallized but hot intrusion?

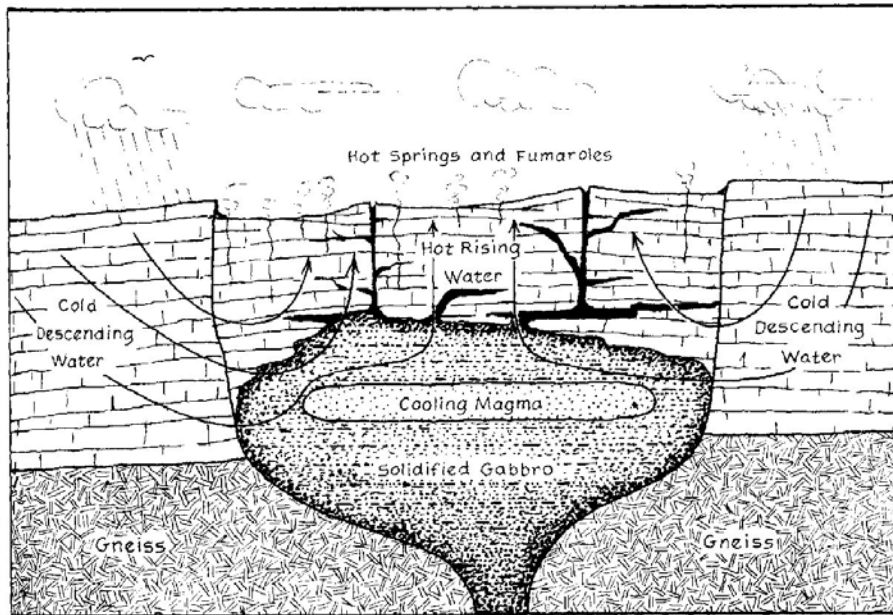
Groundwater: -5 to -25 ‰ in temperate areas

Whole rock granite  $\delta^{18}\text{O}$  ~ feldspar ~ +6 to +13 ‰



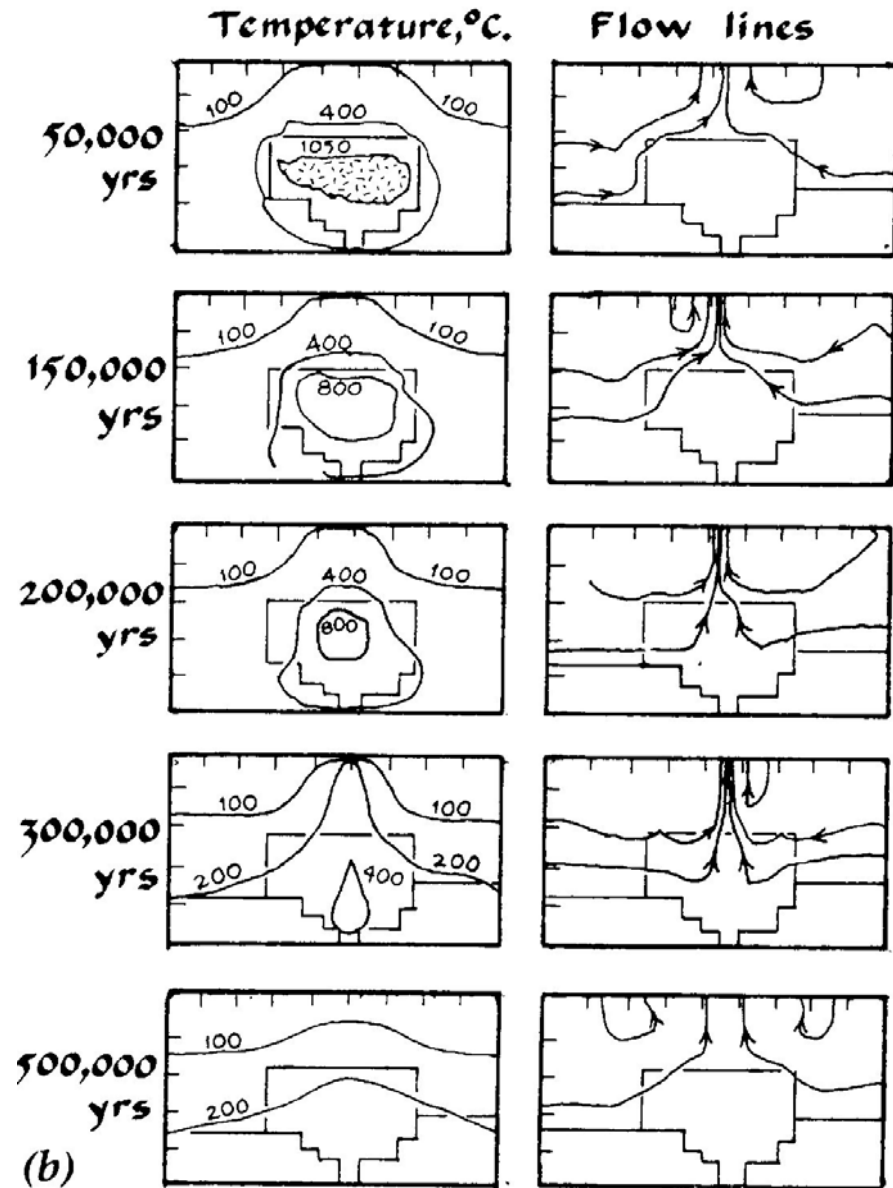
> 450°C;  $\delta^{18}\text{O}_{\text{min-water}}$  is negative

# Cooling of an intrusion: model

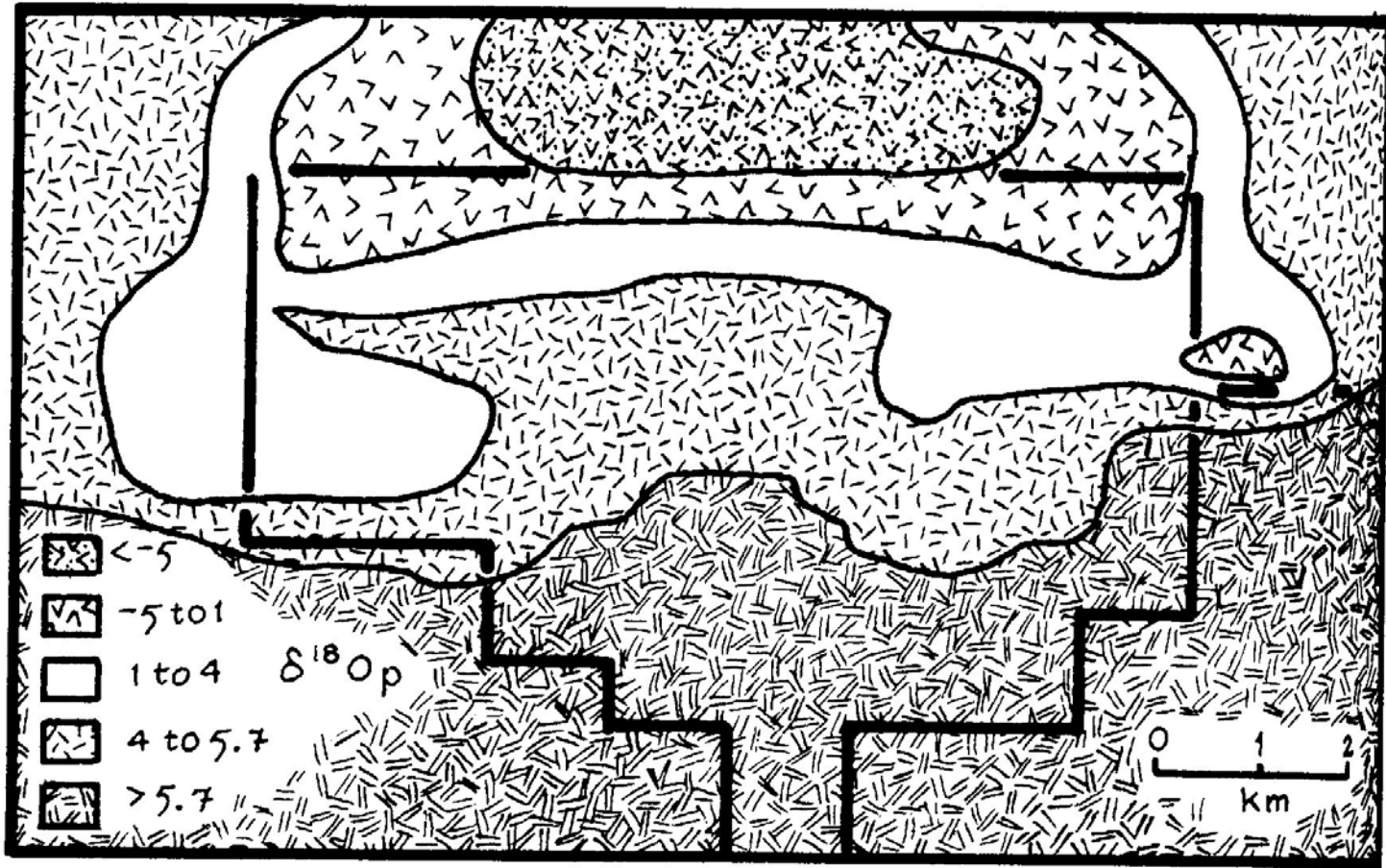


**Figure 6-34** The Skaergaard intrusion set up a large hydrothermal system in which meteoric water heated by the cooling gabbro convected through the permeable basaltic

rocks of the roof and upper walls. (After H. P. Taylor and R. W. Forester, 1979, *Jour. Petr.* 20:355-419).

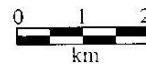
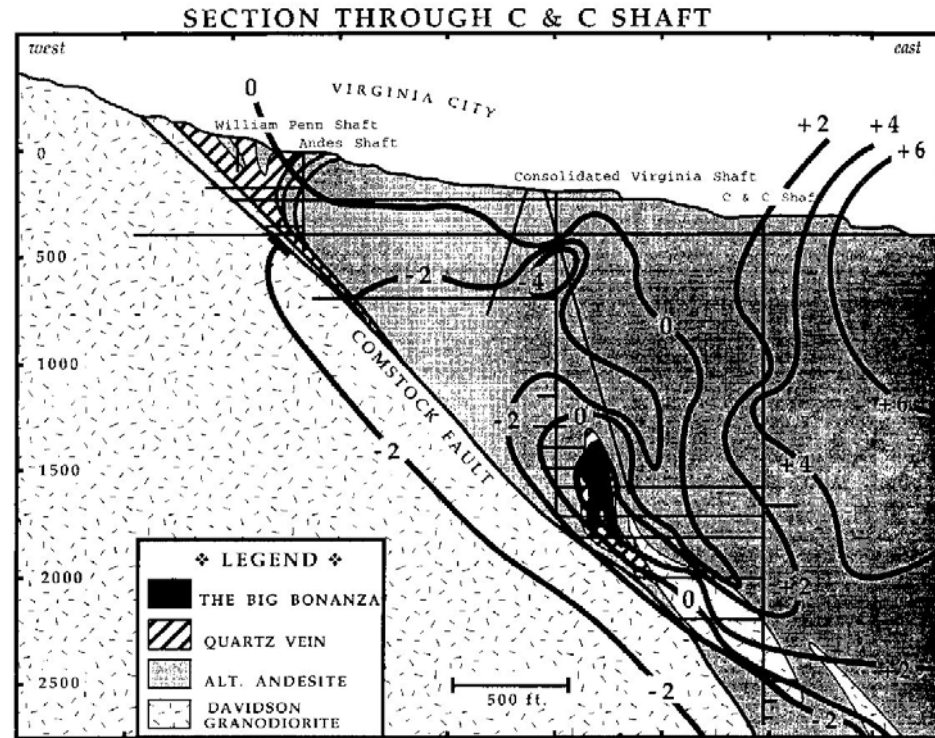
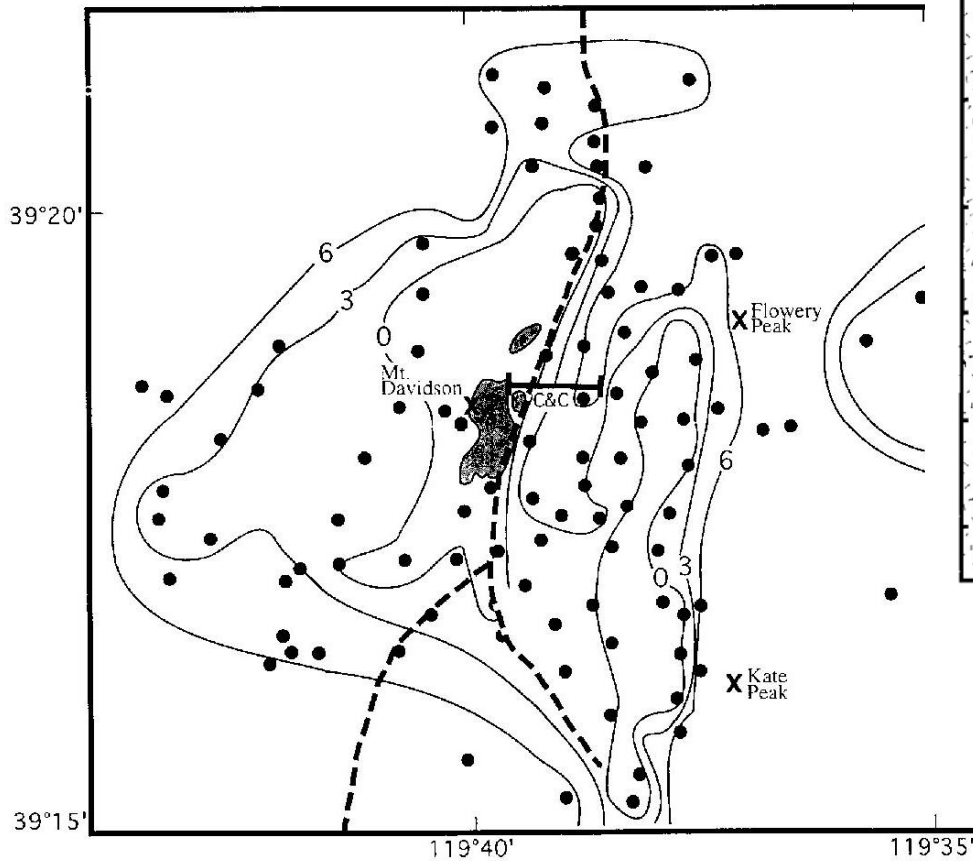


# Cooling and hydrothermal exchange



In map view? Would fractional crystallization produce this picture?

# Famous example: Comstock



# Hydrothermal systems and trace elements





# Fumaroles

- Glass exchanges oxygen with water faster than the feldspar phenocrysts
- In small hydrothermal system = low total water
- Glass oxygen isotopes change while the feldspar  $\delta^{18}\text{O}$  stays pretty much the same.

