

Unlocking the  
Secrets of the  
Mantle Wedge:  
New Insights into  
Melt Generation  
Processes in  
Subduction Zones

T.L. Grove, N. Chatterjee, E. Medard,  
S.W. Parman, C.B. Till







Reginald Aldworth Daly  
(1871 – 1957) :

Bowie Medalist (1946)

*The Quintessential  
AGU member*

- Igneous petrology
- Volcanology
- Paleo-climate
- Mineral Physics
- Geodynamics

*Let us take a moment to  
recount some of Daly's  
thoughts and  
contributions to the  
subject of today's lecture.*

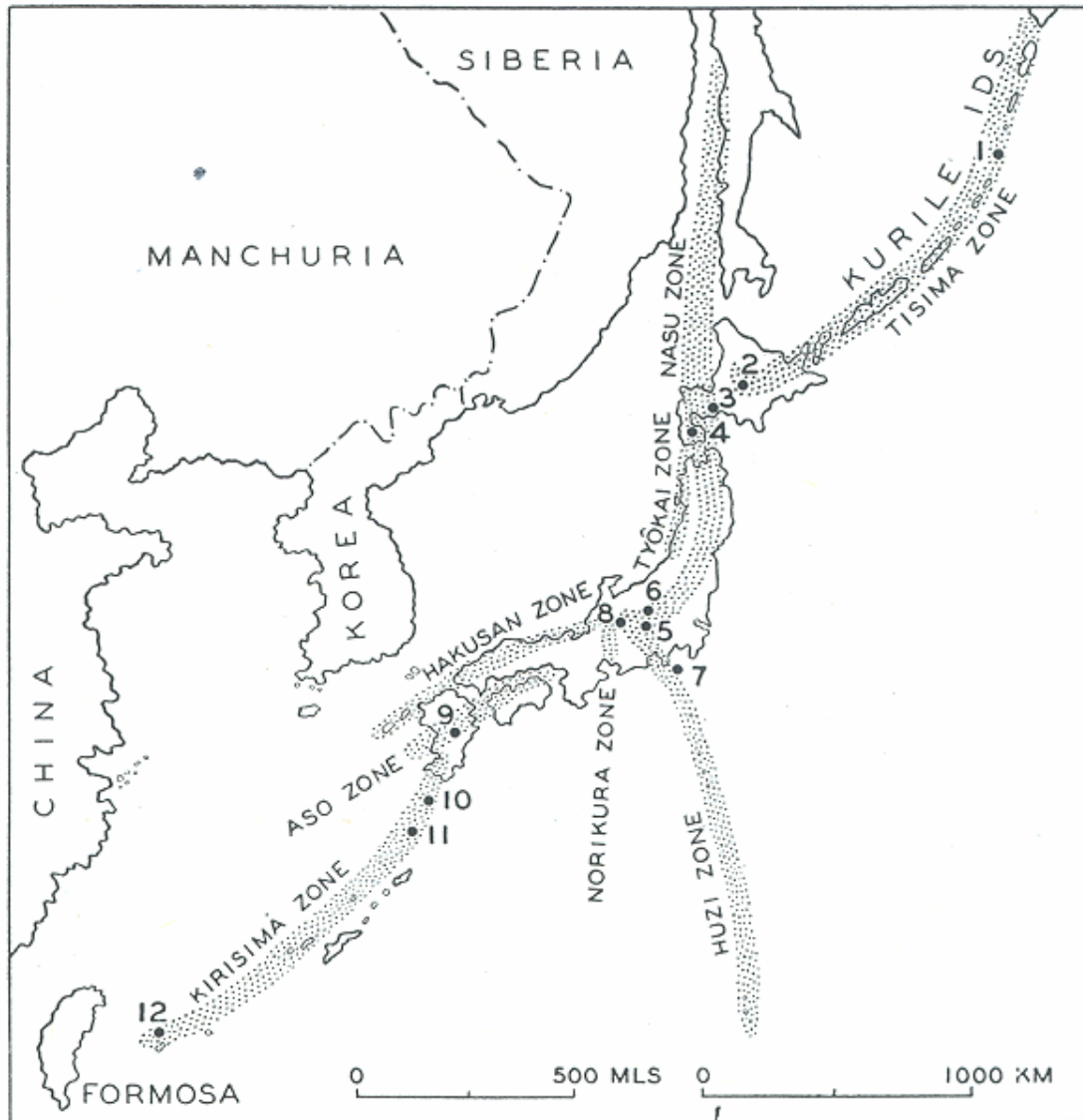


FIG. 74.—Neovolcanic cone chains of Japan and (numbered) localities of volcanic activity during the period 1924–1931. 1, Matuwa Zima; 2, Tokati Dake; 3, Tarumai San; 4, Komaga Take; 5, Asama Yama; 6, Kusatusirane; 7, Osima Mihara Yama; 8, Yake Dake; 9, Aso San; 10, Kutinoerabu Zima; 11, Suwanose Zima; 12, Hatoma Zima. (After H. Tanakadate, *Japanese Jour. Astr. and Geophysics*, vol. 9, No. 1, 1931,

*On island arcs  
in subduction  
zones:*

Daly wrote about these features in his 1933 book “Depths of the Earth and Origin of Magmas”

“Each of these areal groupings of units clearly represents and important genetic problem”

He also endorsed DuToit’s theory of continental drift (1927)

Between 500 bars and 2000 bars the solubility increases nearly as the square root of the pressure, a rule that applies also to the solubility

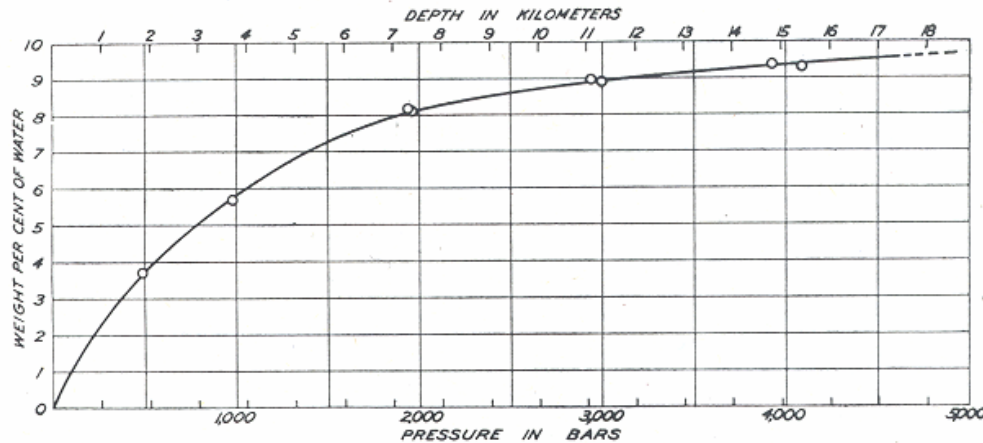


FIG. 123.—Solubility of water in granite glass, as a function of pressure within the earth at the 900° C. isotherm.

of hydrogen in some molten metals, within, however, a different range of pressures. Above 2000 bars the solubility of the water increased

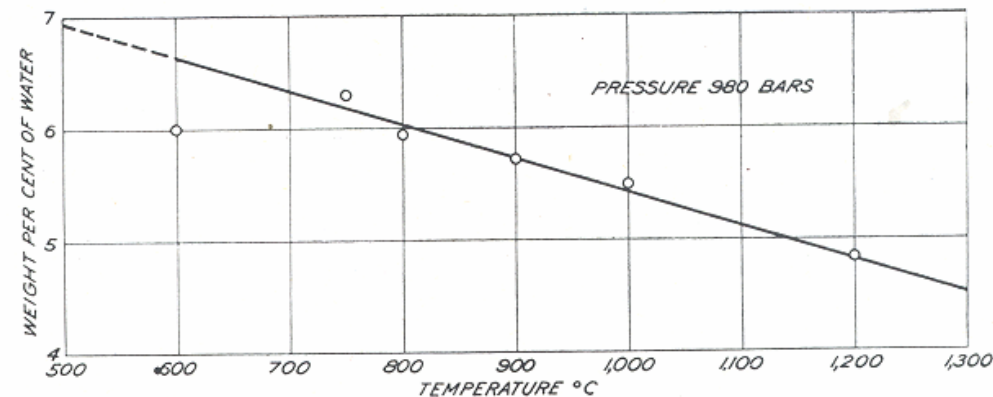


FIG. 124.—Solubility of water in granite glass, as a function of temperature at the pressure of 900 bars (metric atmospheres).

still more slowly with pressure. Goranson's pressure-solubility curve indicates at zero pressure an increase of solubility per bar of 0.0108 per cent of water. Extrapolation to 10,000 bars gives saturation at about 10 per cent of water by weight, instead of about 100 per cent as

## On the importance of H<sub>2</sub>O in magma generation:

Daly wrote the following about the solubility of H<sub>2</sub>O in his 1933 book "Depths of the Earth and Origin of Magmas"

He also noted:

"Other experiments are needed on the solubility of water in basic melts, these representing the dominant magmas in volcanoes of long life."



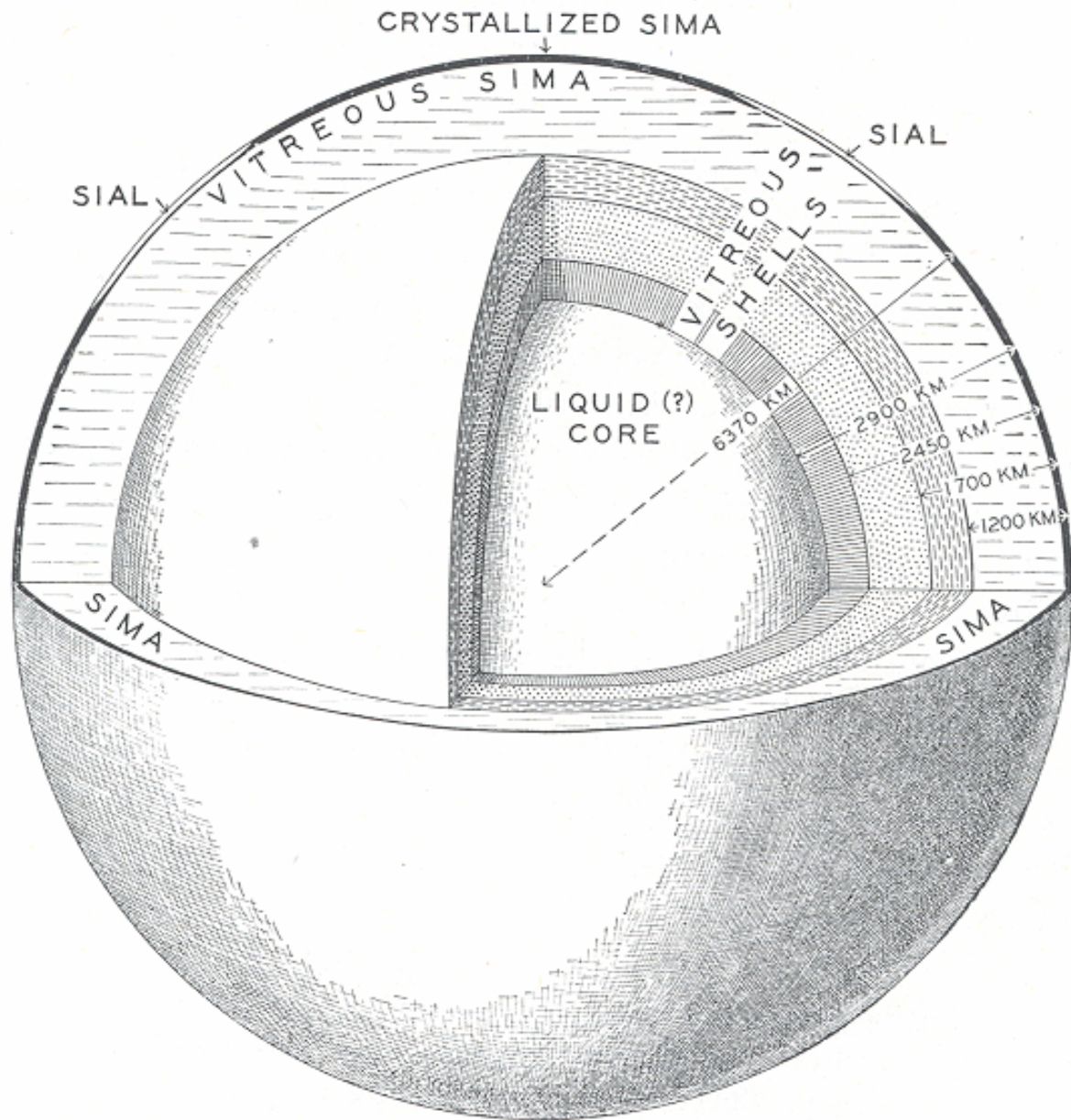


FIG. 91.—Diagram illustrating the shelled nature of the earth. The crystallized Sima below the Sial is thin; thicker elsewhere (heavier exterior line in the sections). Six discontinuities of material are shown, with shell thicknesses nearly to true scale. (After E. A. Hodgson, *Smithsonian Rep. for 1931*, p. 358.)

He also developed a theory of the characteristics of the Earth's deep interior structure –

“The Glassy Shells”

One of the first systematic efforts to relate geophysical measurements to Earth material properties

# Unlocking the Secrets of the Mantle Wedge: New Insights into Melt Generation Processes in Subduction Zones

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S.W. Parman, C.B. Till

- 1) Chemical transport processes from subducted slab to the overlying wedge. Melting from top to bottom in the wedge. Field and experimental evidence.
- 2) Element transport from slab to wedge > Melts? Fluids ? Or more complex processes?
- 3) Insights into subduction zone processes. New experimental constraints.



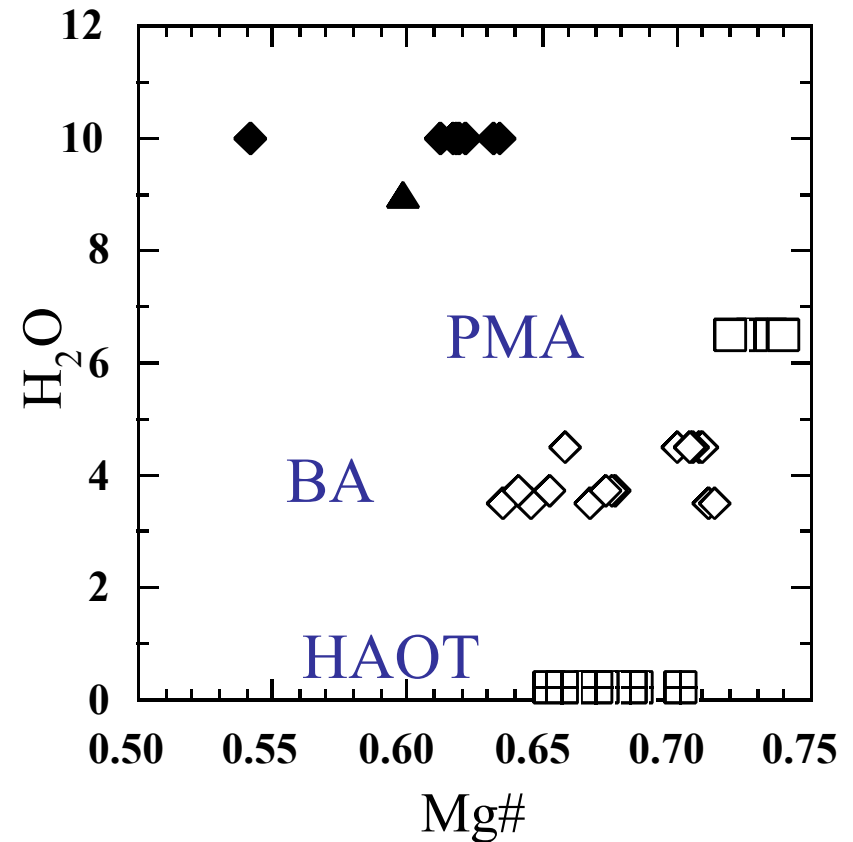
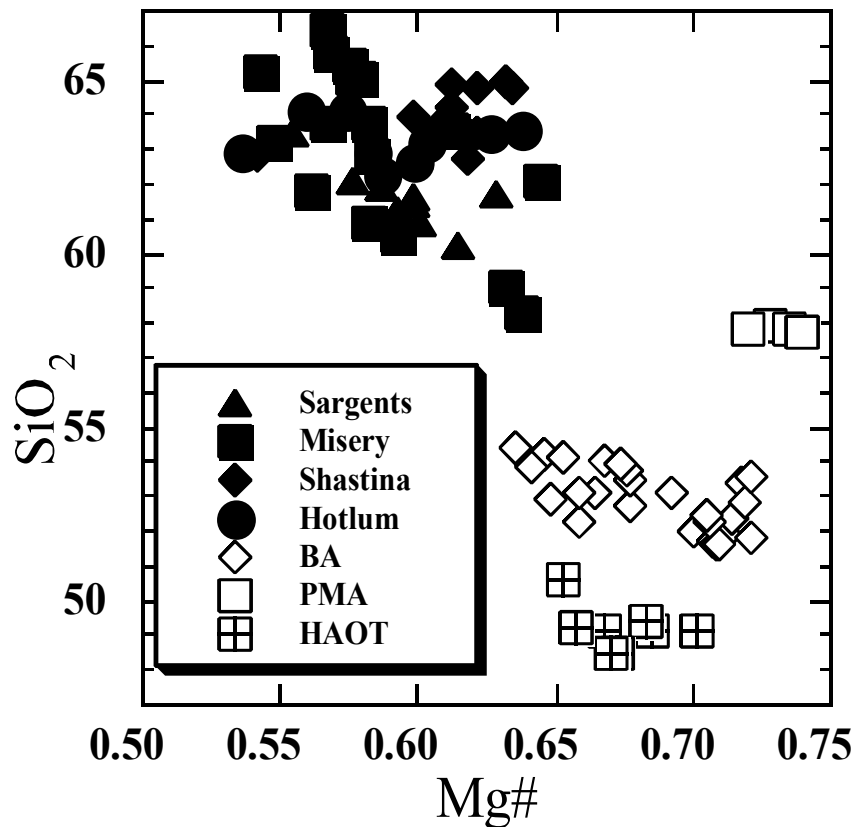
# Topic 1: Chemical transport processes from slab to wedge. Field and experimental evidence from Mt. Shasta region, USA.

- Lavas are high- $\text{H}_2\text{O}$  mantle melts with a significant component added from the subducted slab.
- Where are these melts generated in the mantle wedge?
- What is contributed from the subducted slab?

*Mt. Shasta, N. Calif. – looking W from Med. Lake  
Shasta produced  $\sim 500 \text{ km}^3$  magma in  $\sim 250,000$  years.*

# Major elements and H<sub>2</sub>O

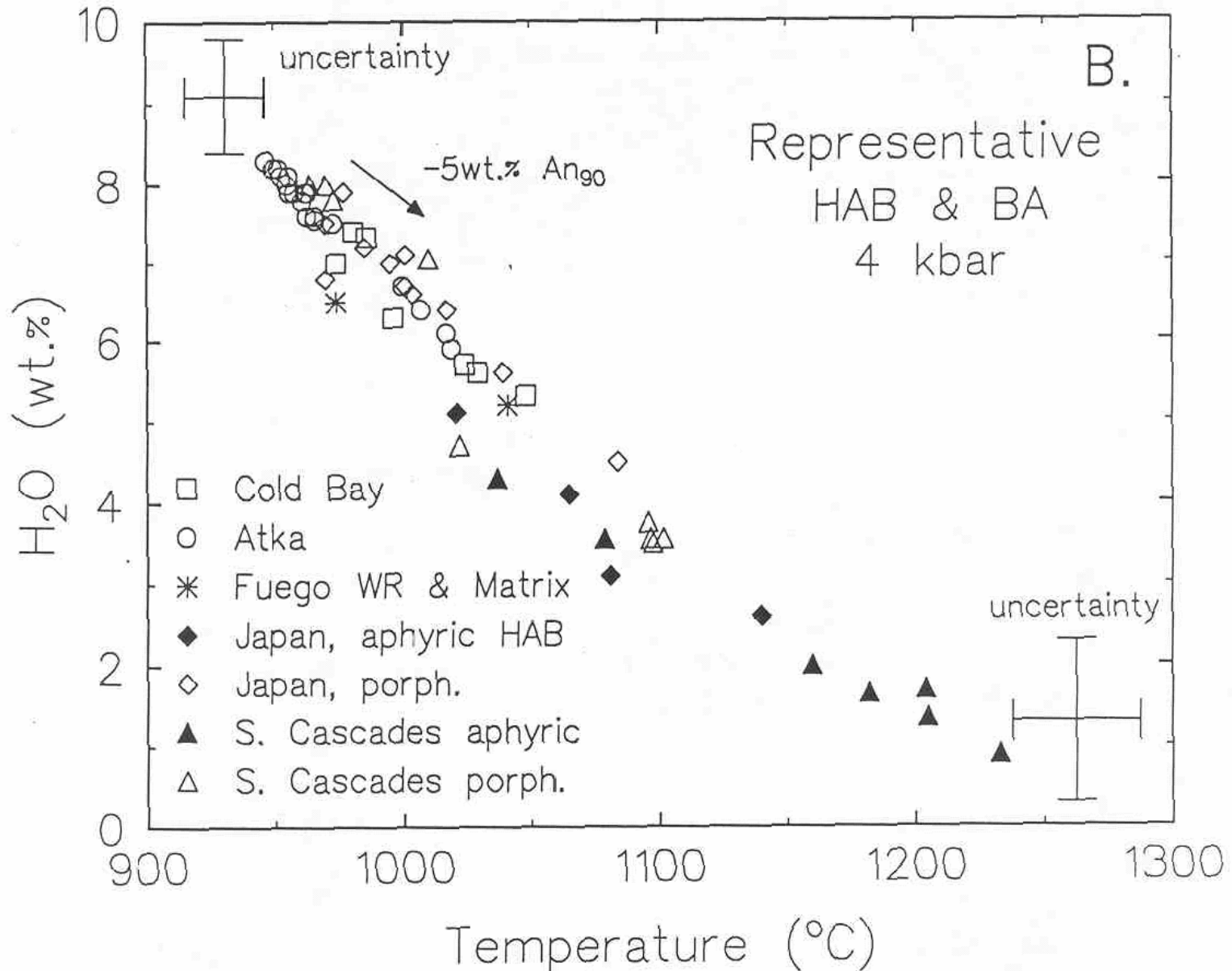
Wet, primitive andesites are in equilibrium with mantle residues = melts of depleted mantle





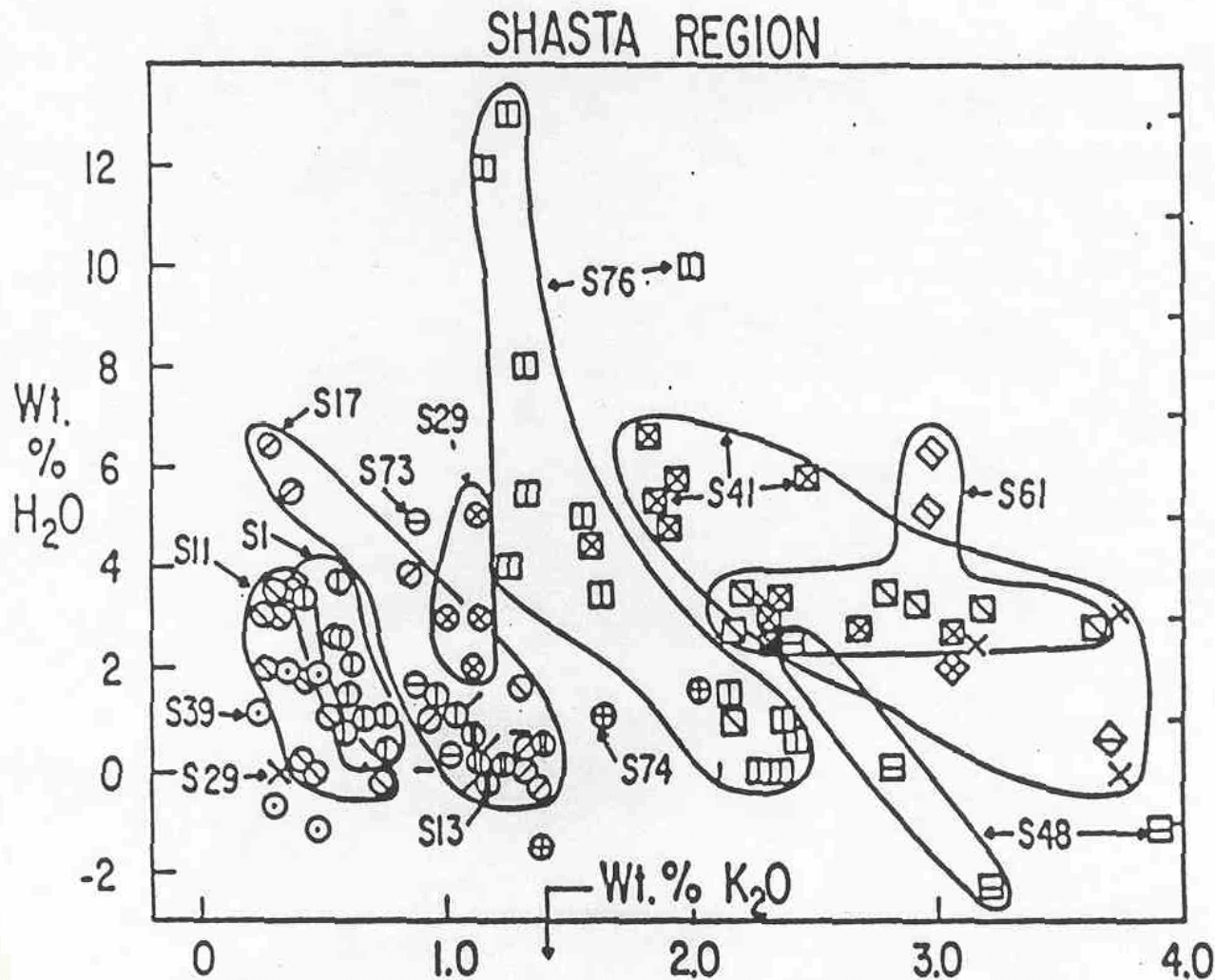
# Estimates of Pre-eruptive H<sub>2</sub>O

- H<sub>2</sub>O solubility in silicate melts is P-dependent and goes to ~ 0 at P = 1 bar.
- So, H<sub>2</sub>O is often lost as a gas phase
- Pre-eruptive H<sub>2</sub>O contents are obtained using:
  - Thermodynamic models of mineral/melt equilibria.
  - Effect of H<sub>2</sub>O on “freezing path” or melt composition produced during fractional crystallization.
  - Direct measurement of H<sub>2</sub>O in melt inclusions.



Estimation of pre-eruptive H<sub>2</sub>O content





Direct measurement of H<sub>2</sub>O in Shasta melt inclusions (Anderson, 1979).

H<sub>2</sub>O-contents of arc magmas seem to be too high to result from any batch melting process of any potential H<sub>2</sub>O-bearing mantle source.

New experimental evidence changes this.


How do these new phase equilibrium constraints help us understand the processes of melting in subduction zones?

S76

Shastina summit – from Mt. Shasta, N. Calif.

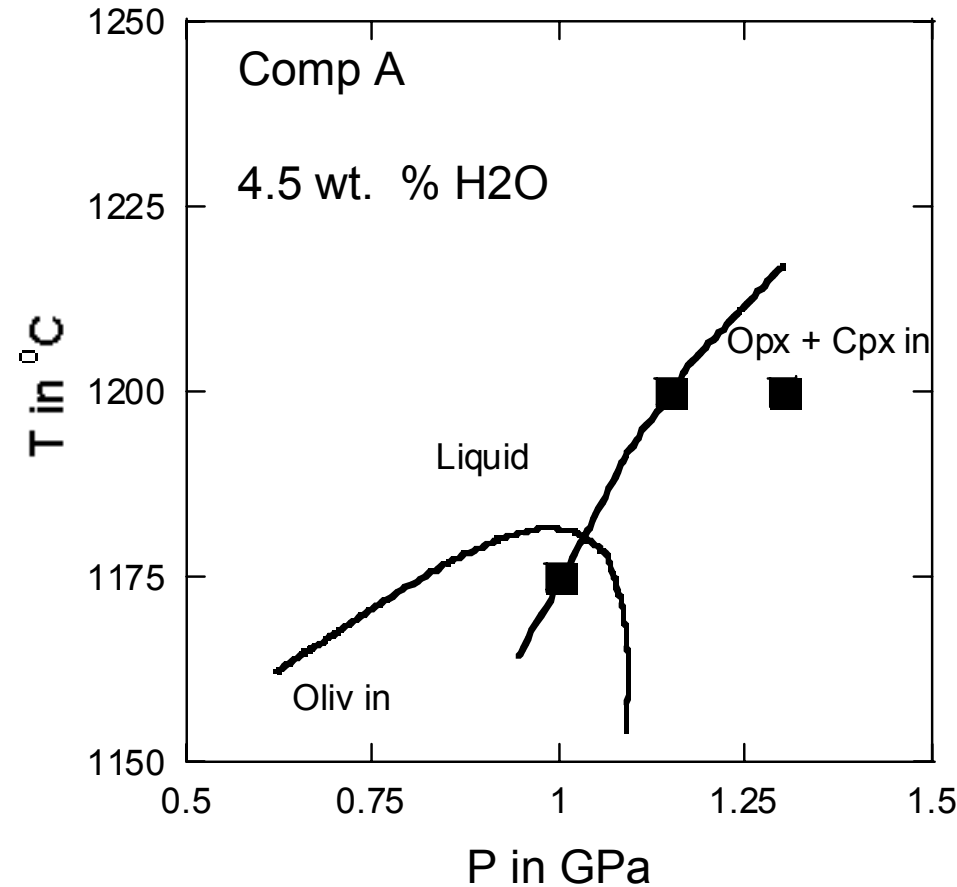
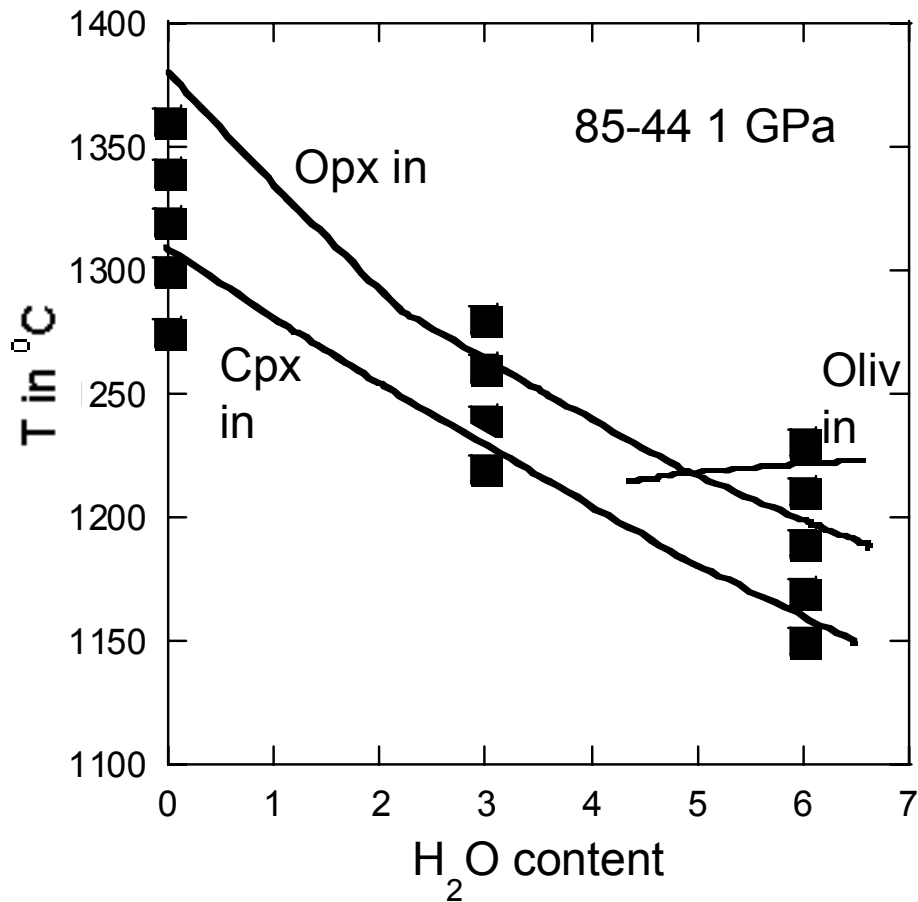






Cinder Cone  
Basaltic Andesite —  
85-44 and S-1

Primitive BA (S-1) and PMA (S-17) – Hydrous melts saturated with a harzburgite residue at top of mantle wedge > 25 % melting.





## Topic 2: Estimating the chemical composition of the fluid-rich component.

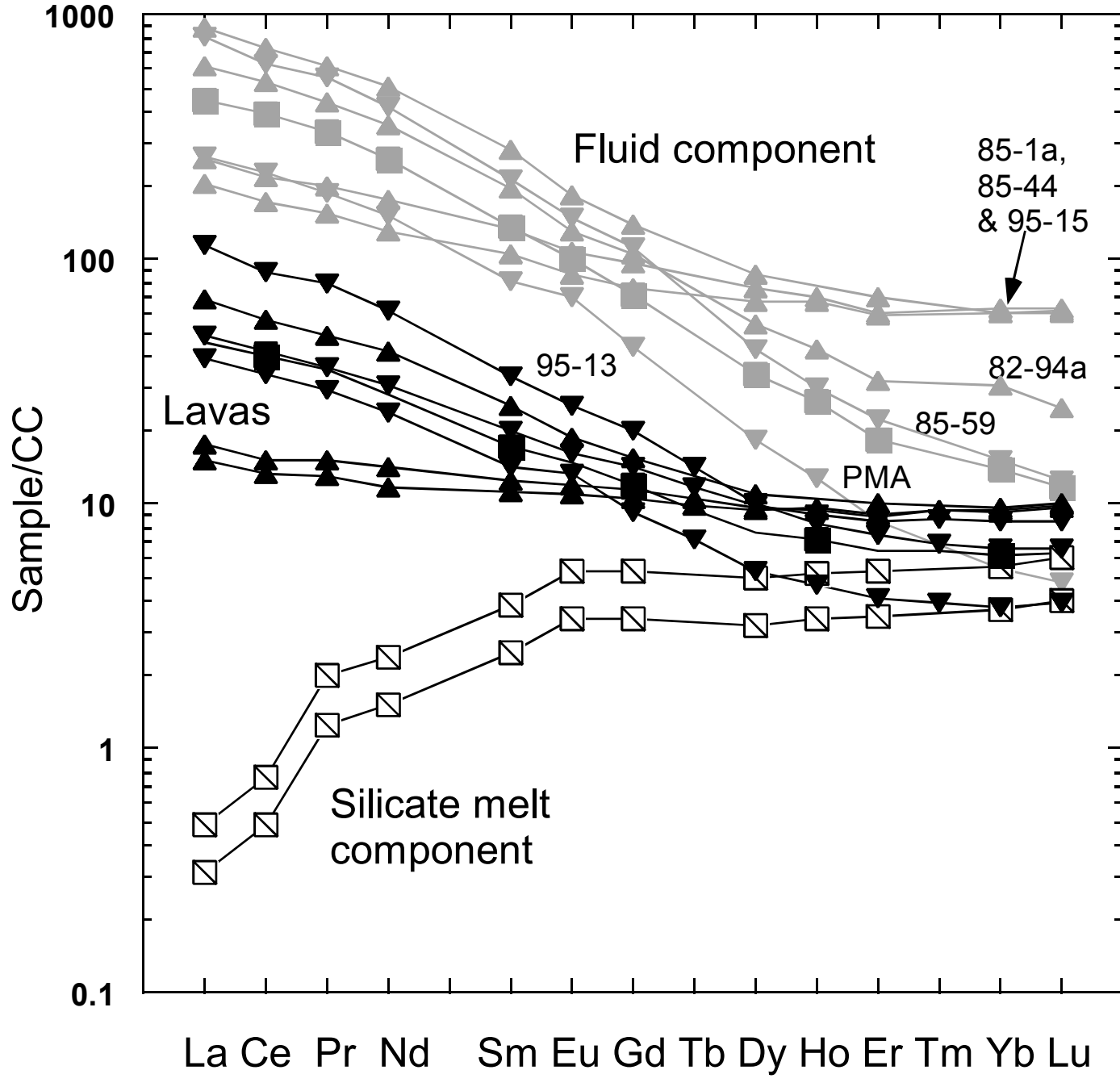
- We will model this by assuming 2 components:  
1) a silicate melt from a harzburgite residue (wedge) 2) a fluid-rich component from the subducted lithosphere (slab).
- Use mass balance. Calculate elemental contribution from mantle melting
- Use  $H_2O$  content of lava to estimate the composition of the  $H_2O$ -rich component.

# Mass Balance Model

$$C_{\text{fluid}} = (C_{\text{lava}} - X_{\text{melt}} C_{\text{melt}}) / (X_{\text{fluid}})$$

- Substitute batch melting equation for  $C_{\text{melt}}$
- $F$  is fraction of mantle melt and
- $D$  is bulk distribution coefficient
- $C_0$  element abundance in mantle source
- $\alpha$  is a correction for other elements in fluid

$$C_{\text{fluid}} = (C_{\text{lava}} - (1 - X_{\text{H}_2\text{O}}/\alpha)C_0 / [F + D(1 - F)]) / (X_{\text{H}_2\text{O}}/\alpha)$$



Estimated fluid component = gray

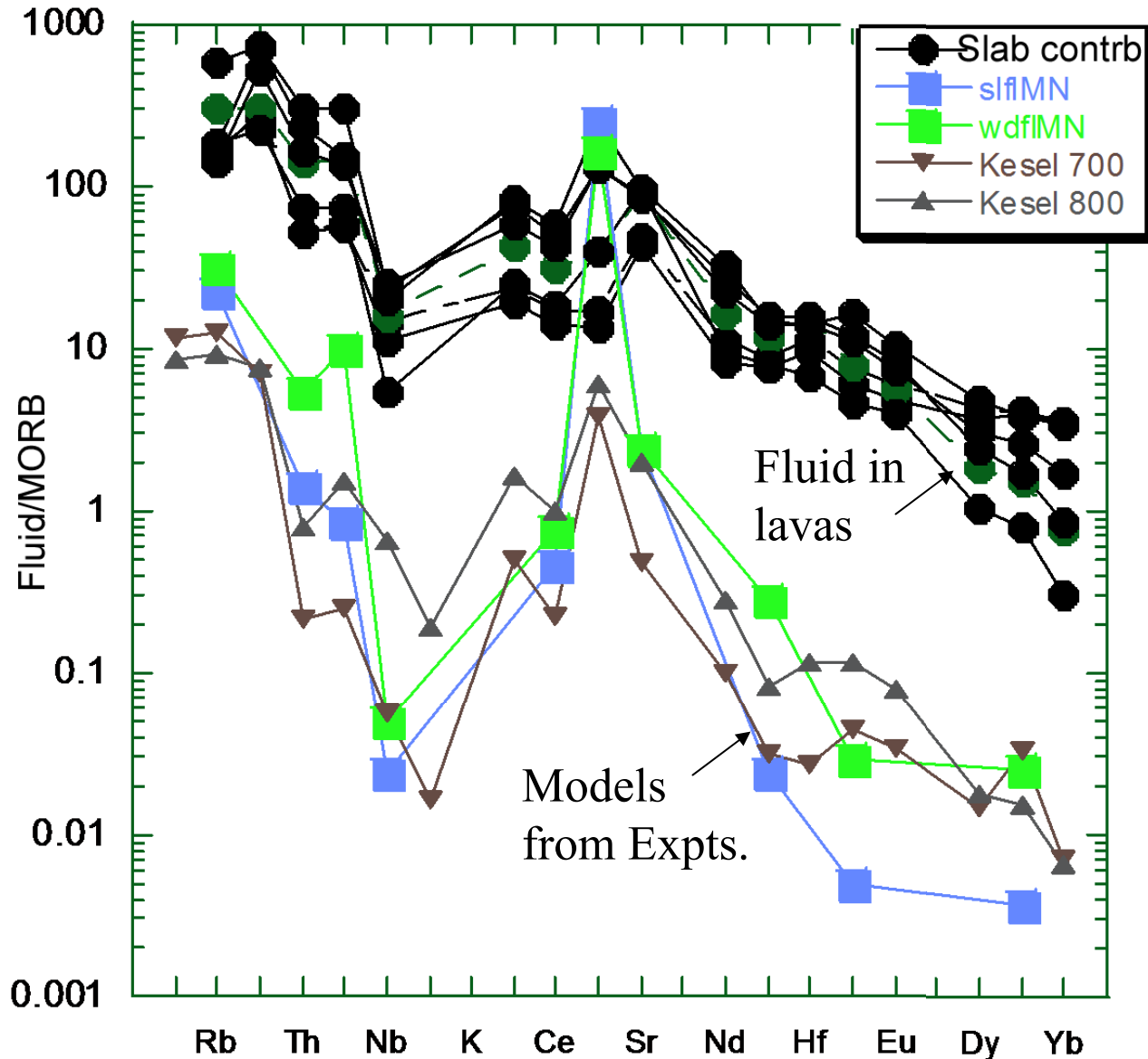
Lavas = solid black

Silicate melt of mantle = open square.

Note the dichotomy in  $\text{La}/\text{Sm}_N$  and  $\text{Dy}/\text{Yb}_N$



H<sub>2</sub>O-rich component a fluid? No...



Estimated fluid-rich component (black circles)

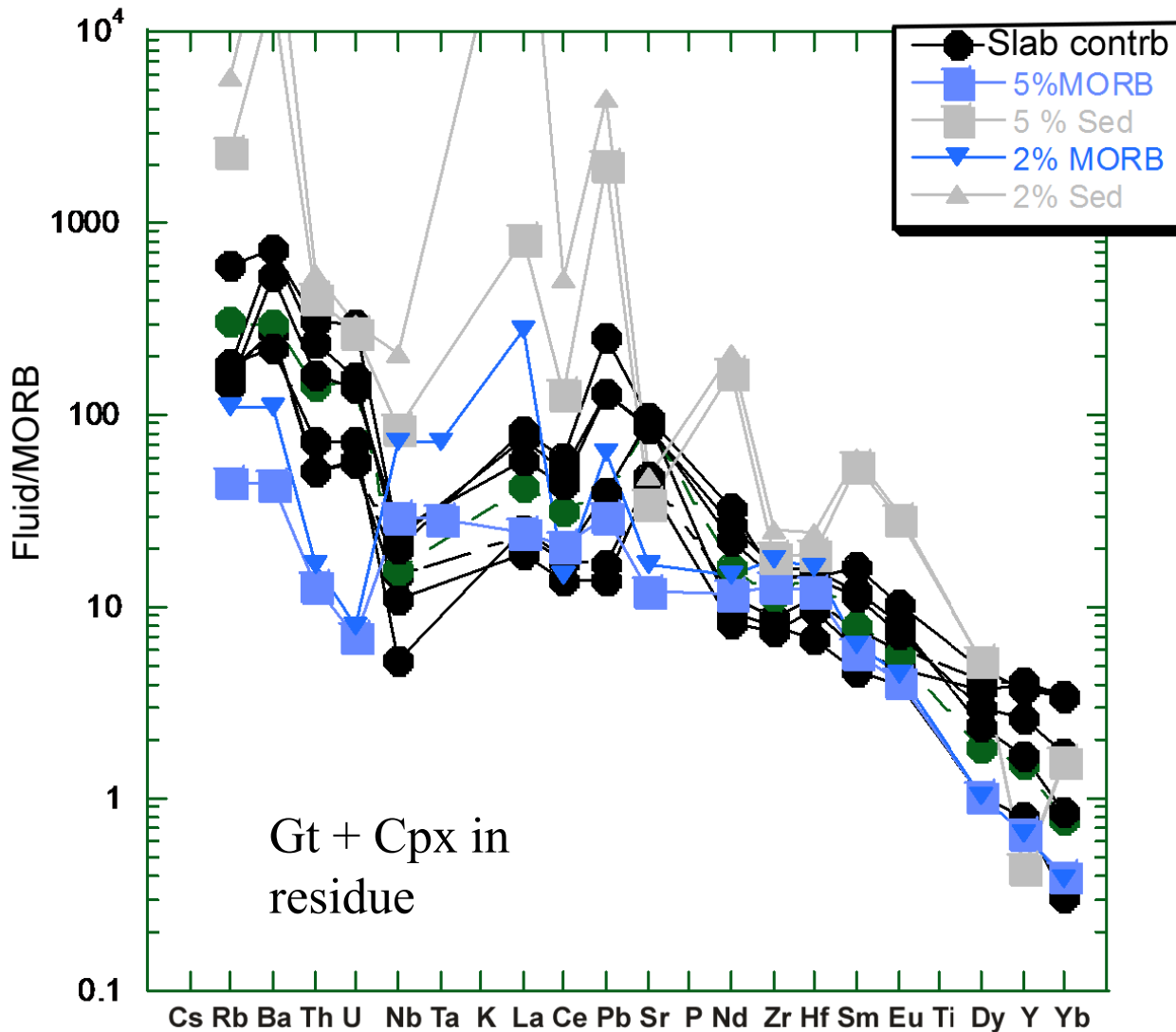
Least similar to a hydrous fluid saturated with eclogitic residue

Slf = slab fluid  
Ds from Ayers, Brenan, Kogiso, Stalder, etc.

Wdfl = wedge fluid

Kesel (2005) = fluid in MORB at 4 GPa

H<sub>2</sub>O-rich component a silicate melt?  
 Much closer....

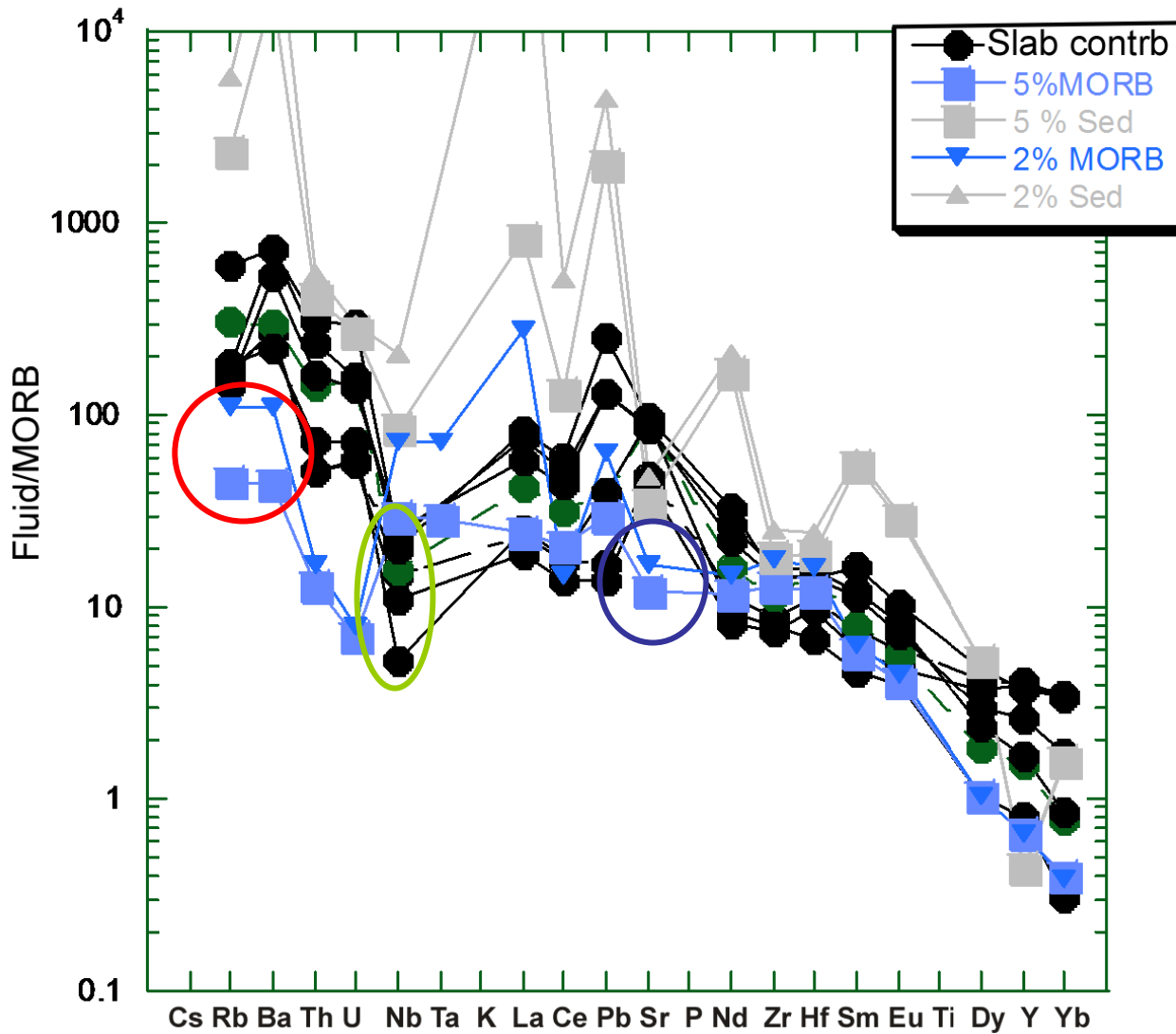


Estimated fluid-rich component (black circles)

Most similar to a mix of hydrous low degree melt of eclogitic residue

*n*-MORB (Hofmann) and Sediment (Ben Otham)

eclogite melt Ds from Green et al. (2000)



But the eclogite melt model of MORB & Sediment are not perfect fits.

Misfits:

Highly incompatible elements

& HFSE

& Fluid mobile



# Any H<sub>2</sub>O rich slab fluid/melt is likely to interact with the wedge

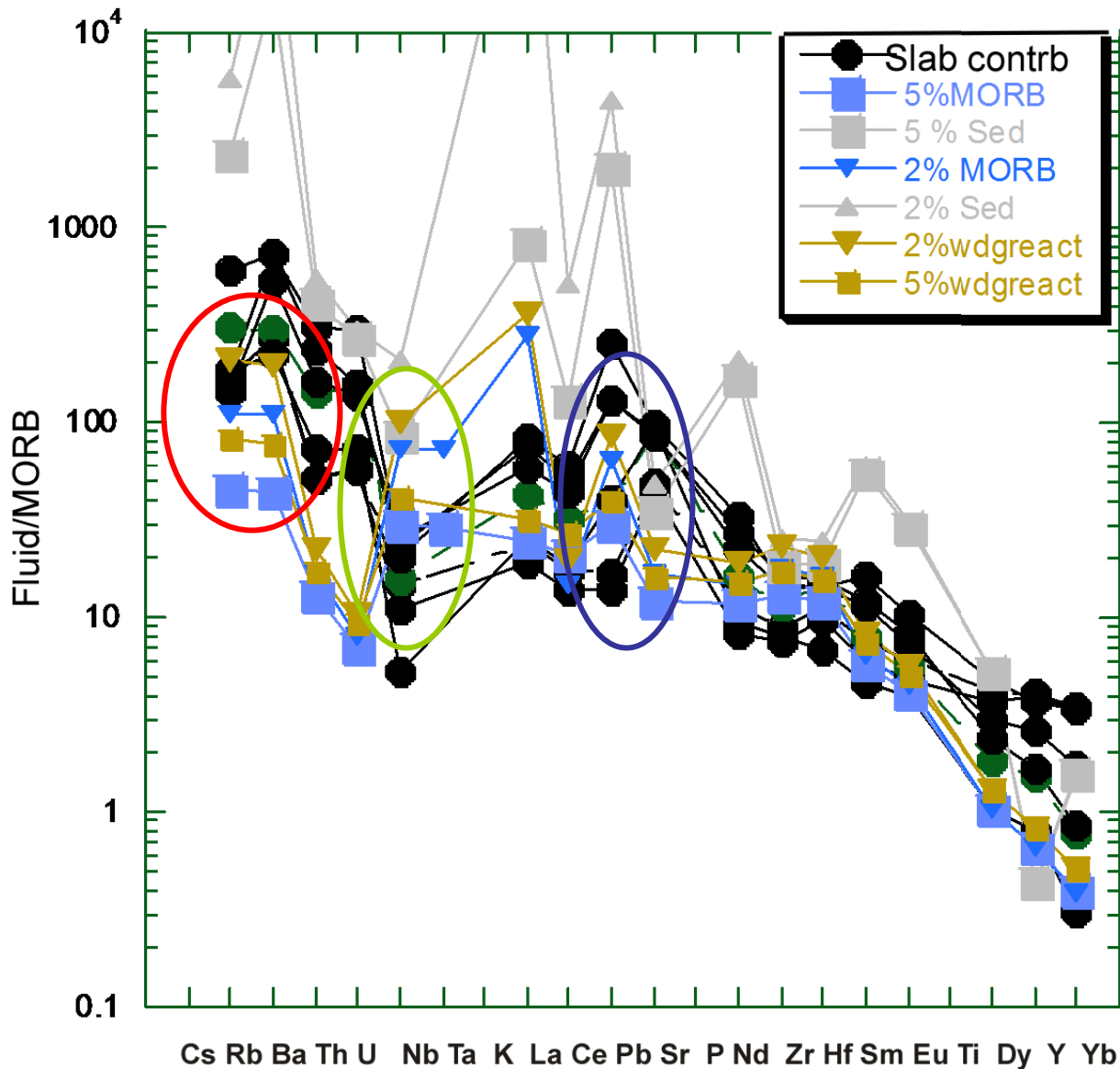
- SiO<sub>2</sub> solubility in an H<sub>2</sub>O-rich fluid will be low -Zhang & Franz (2000) Newton and Manning (2003)



- Bell et al (2005) characterize chemical interaction between wedge & subduction added component in Kaapvaal harzburgites. Metasomatic reaction is:



Let's further react the slab melt with the wedge. The result is Distilled Essence of Slab Melt.

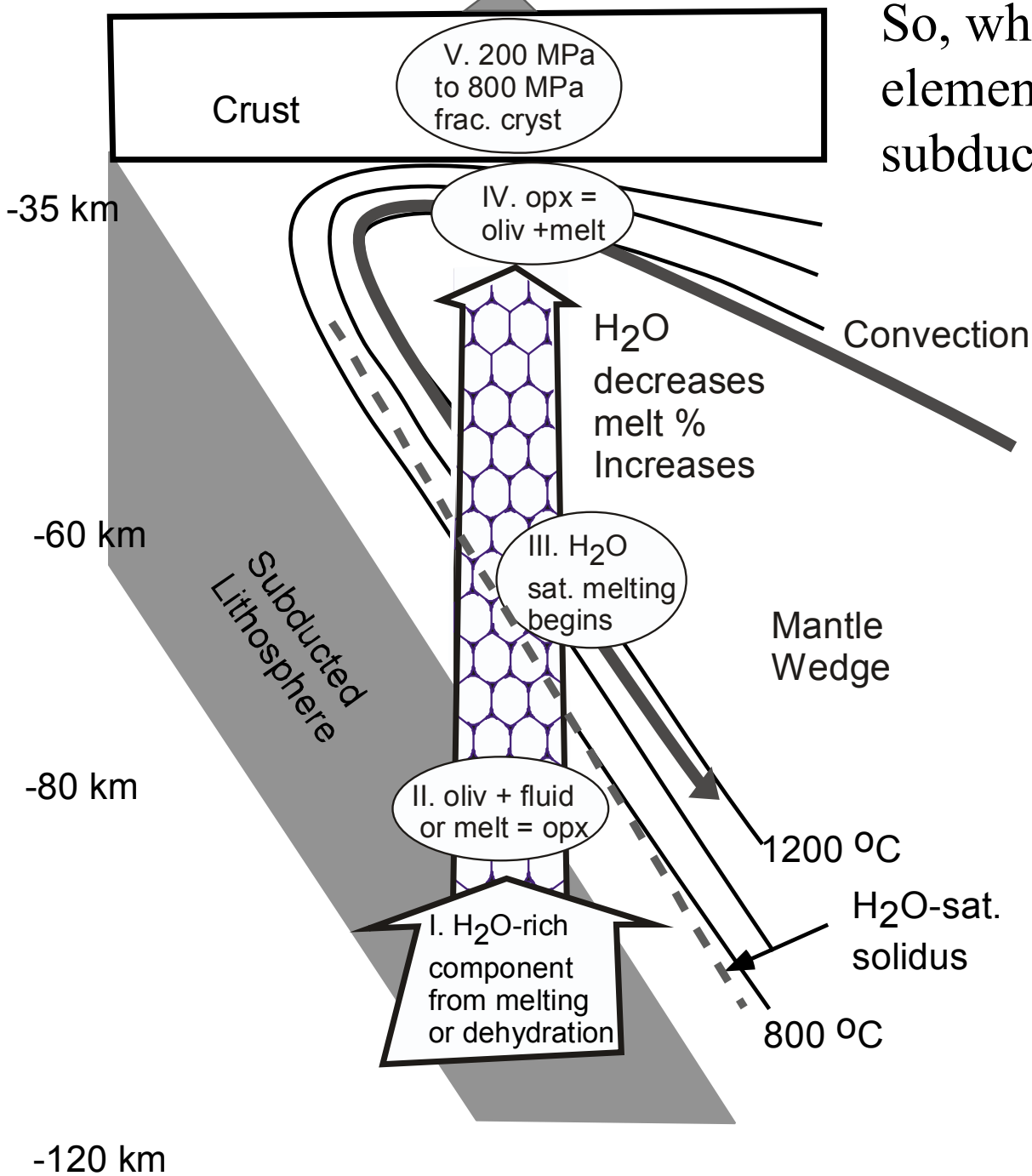


Brown symbols show effect of wedge peridotite + slab melt interaction at base of wedge using reaction inferred by Bell et al. (2005).

highly incompatible elements - better

HFSE - worse

Fluid mobile - better



So, what medium transfers elements from hot, young subducted lithosphere?

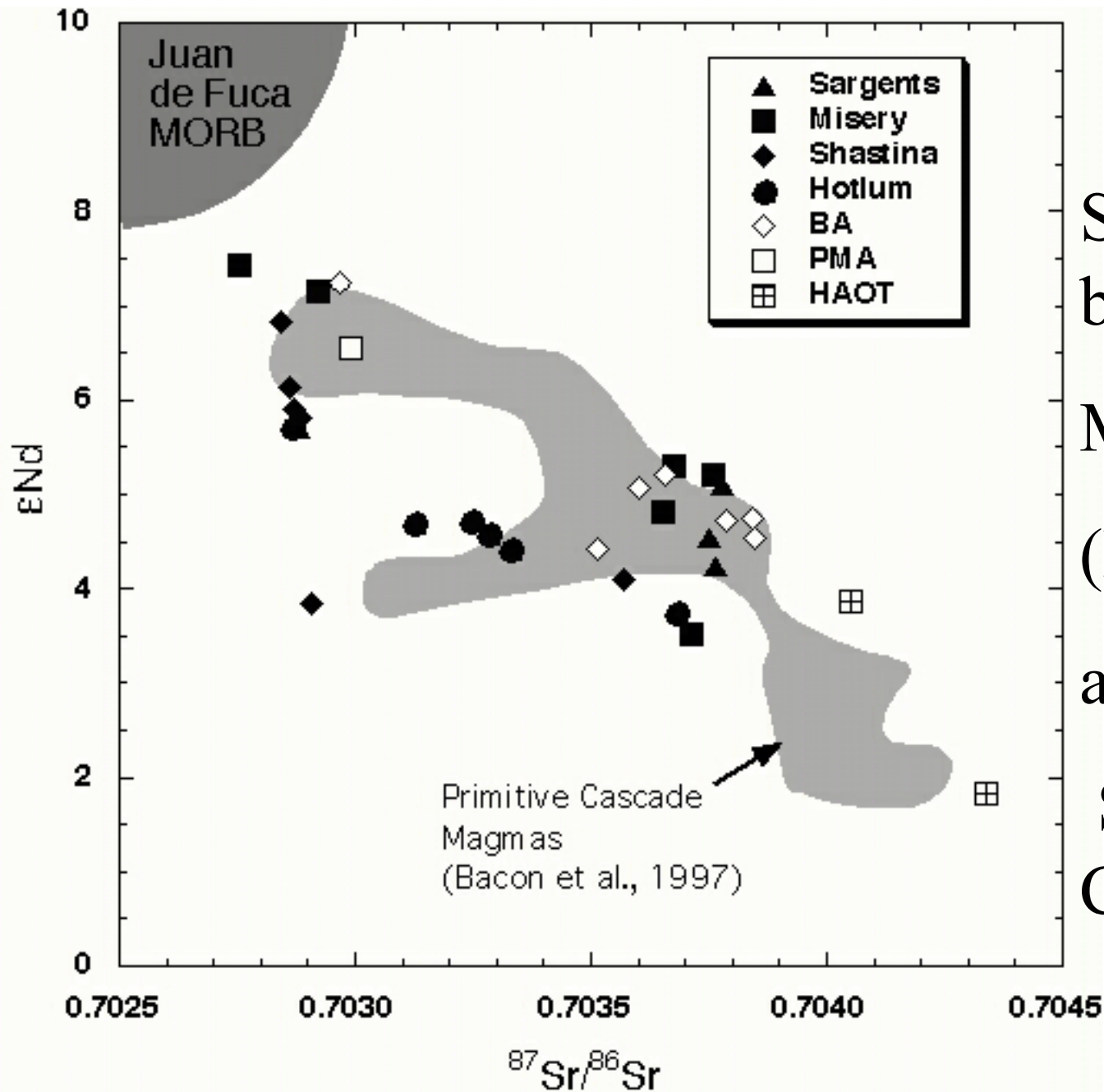
Is it a melt?? A fluid??

Looks most like a low degree melt of sediment/MORB eclogite.

Fluid – not a good match.

Mantle wedge / slab melt interaction improves model.





Signatures of both:  
 MORB  
 (High La/Sm)  
 and  
 Sediment  
 Components

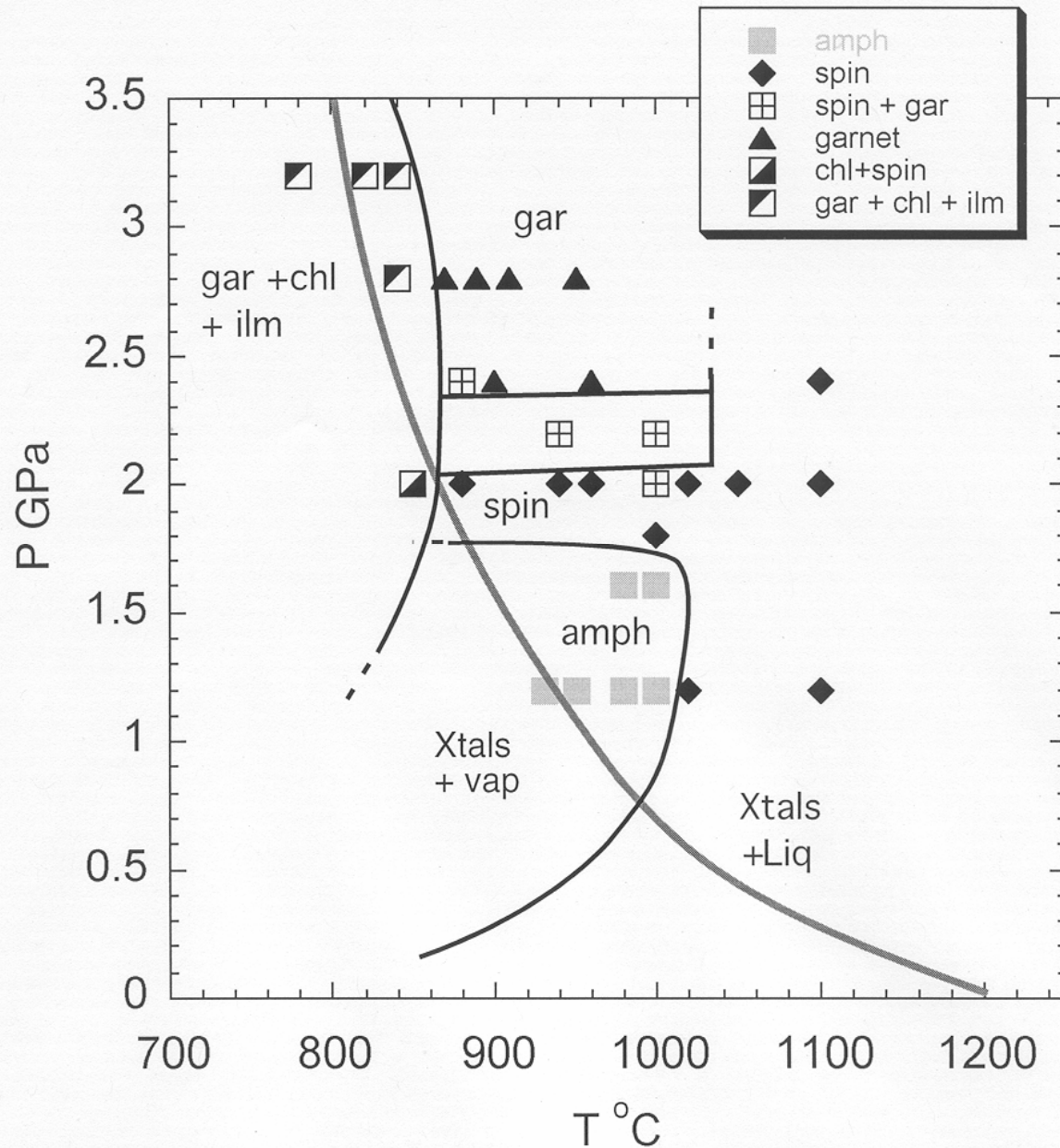
### Topic 3: New experimental constraints on subduction zone melting processes.

- Can the slab and the wedge BOTH melt?
- Can we understand the high pre-eruptive  $H_2O$  contents of arc magmas?



Mt. Shasta, Calif. – looking South from US 97.



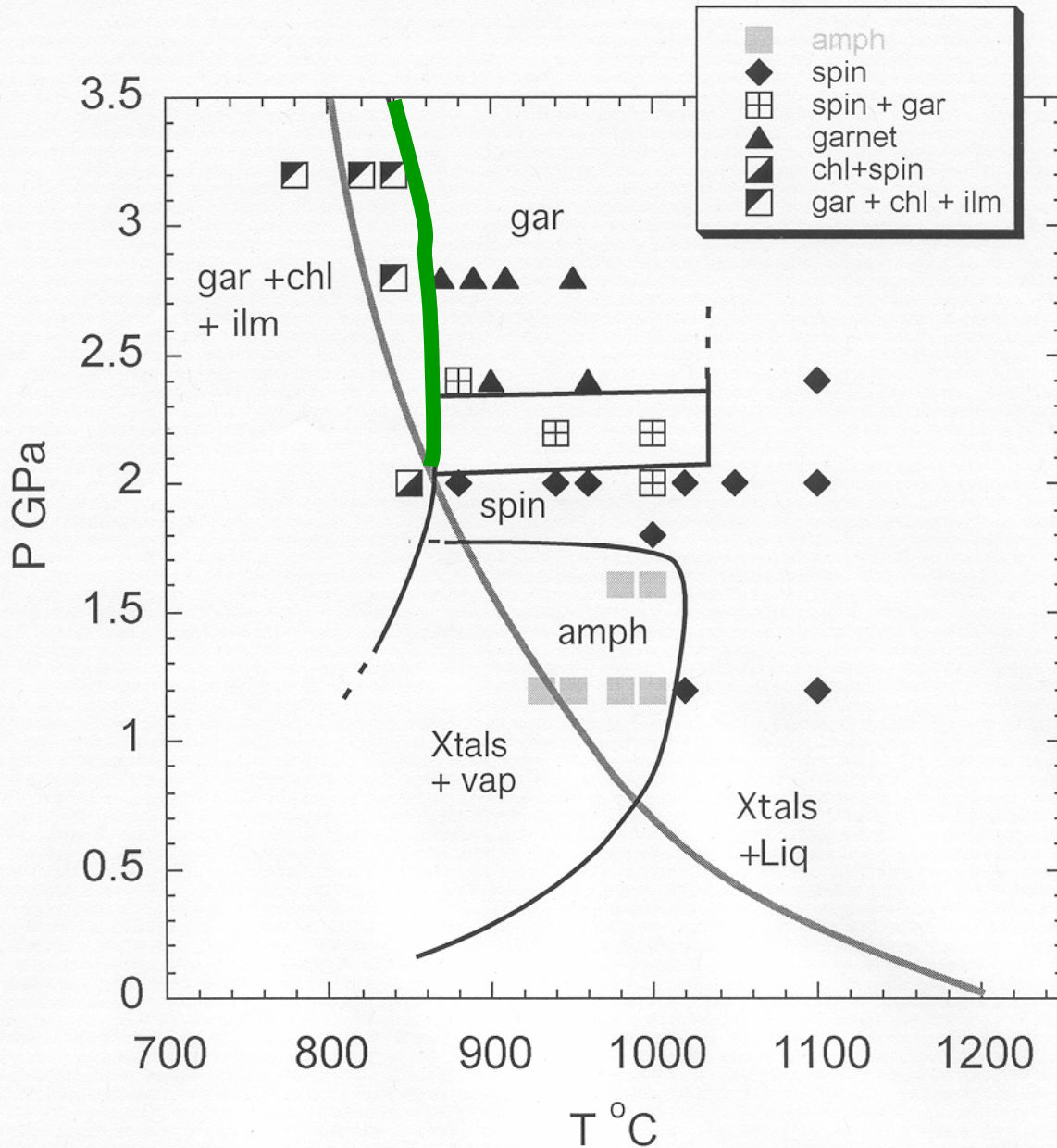


3) New experimental data from Grove et al. (2006) EPSL 249: 74-89

Shows that hydrous phases are stable on the vapor-saturated mantle solidus.

We will use this data to develop a model for melting in the mantle wedge.



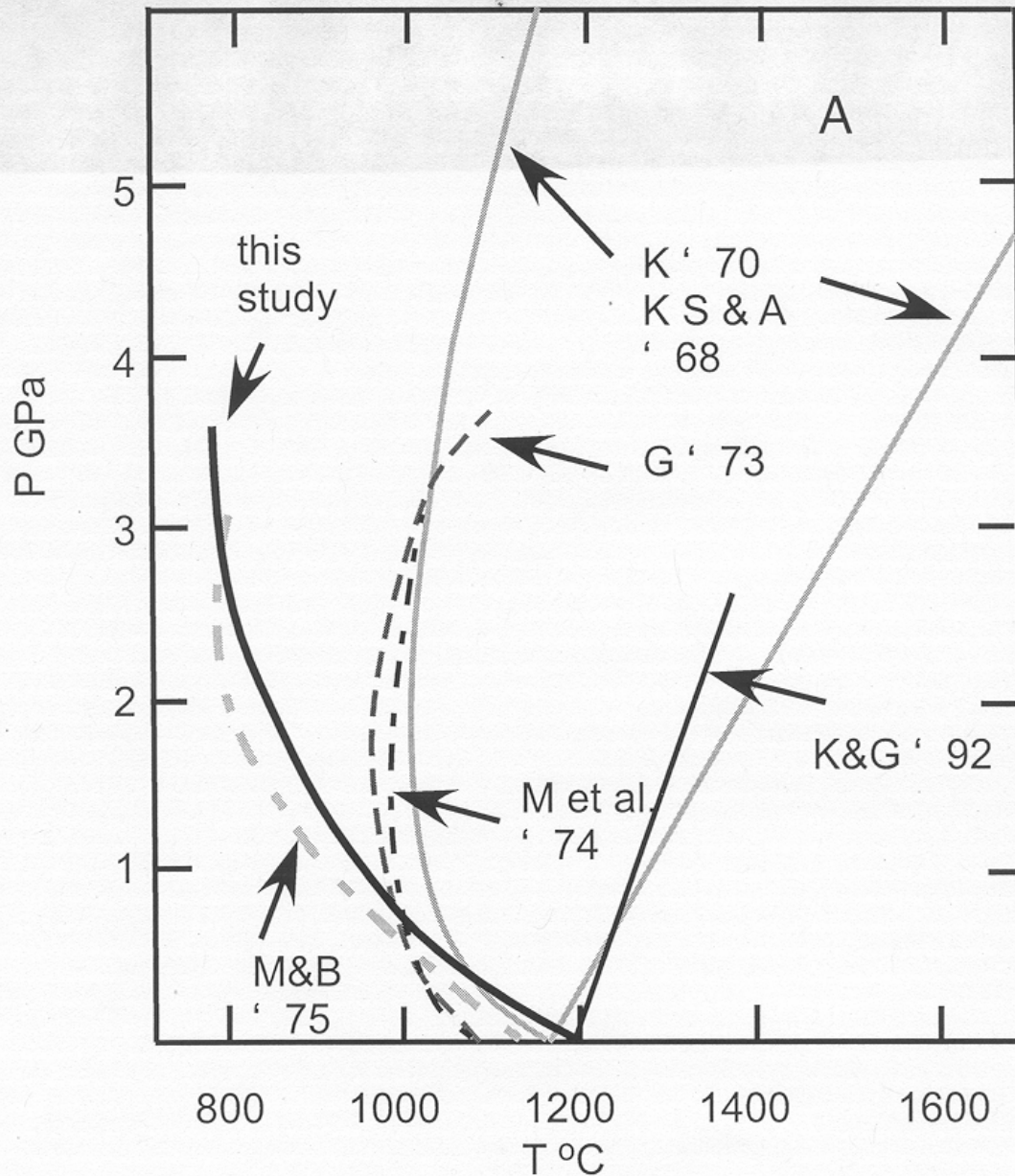


Chlorite on the  
vapor - saturated  
solidus

– a way to  
transport H<sub>2</sub>O  
deep into the  
wedge

Also, Ilmenite,  
Rutile &  
Ti-clinohumite  
are stable.

# Old & New Expts. Why the difference? – melting kinetics

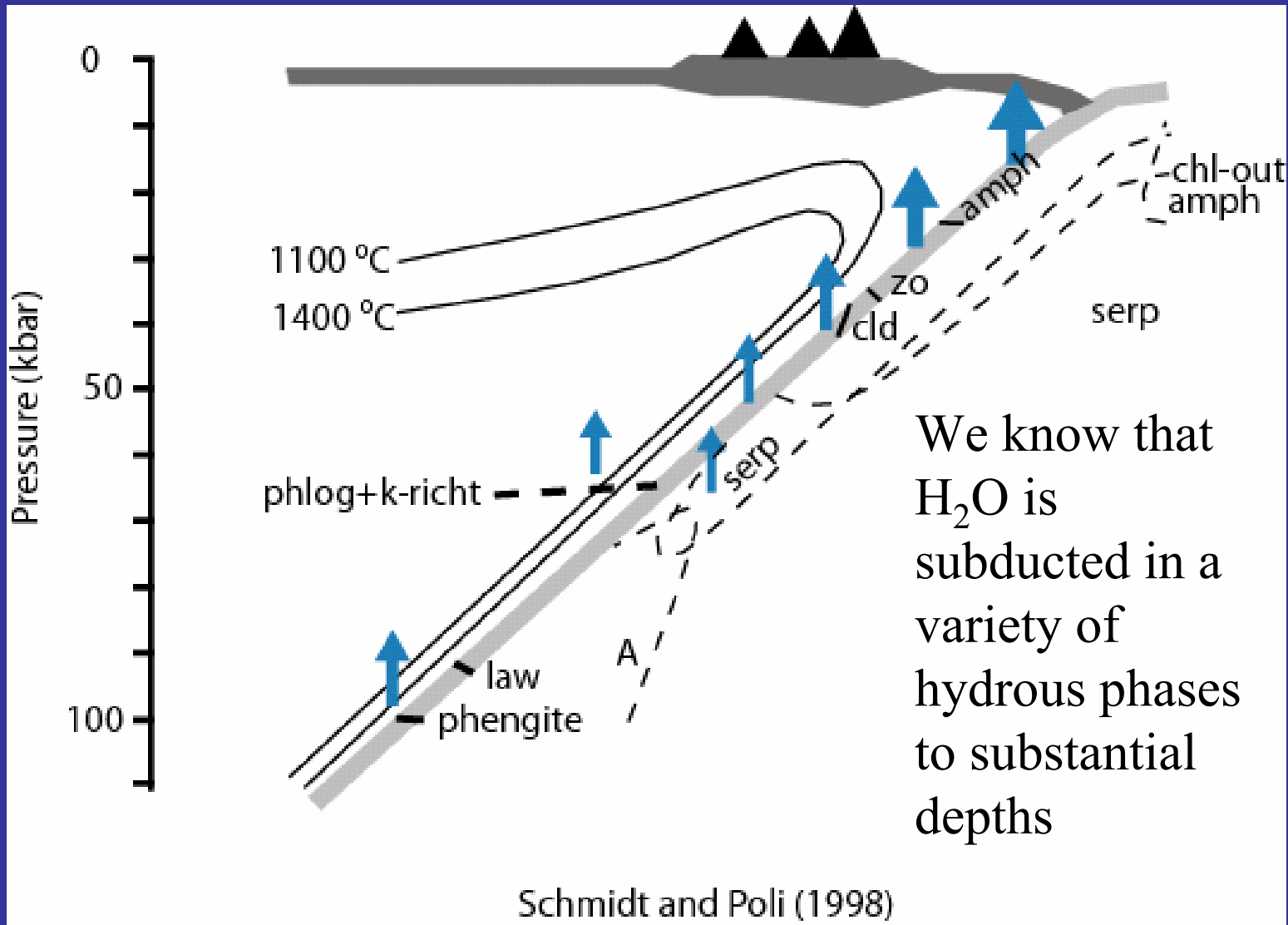


Olivine melting rate is slower than that of pyroxene

-Olivine also melts at a lower Temperature by about 200 °C

- In the short run time expts pyroxene melted first

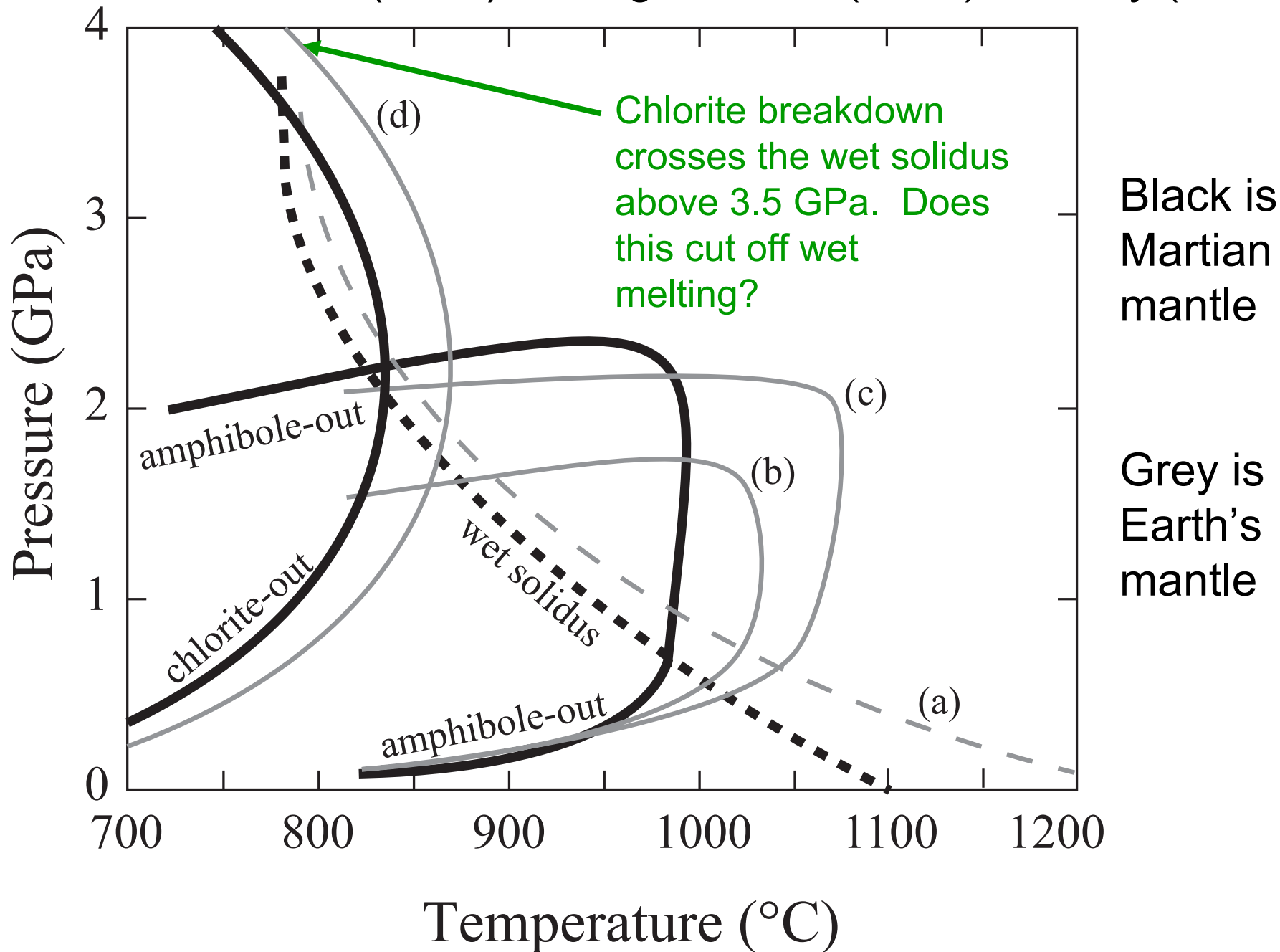
# Serpentine and chlorite dehydration as a source for H<sub>2</sub>O.



How do these phases interact with the wet peridotite solidus?

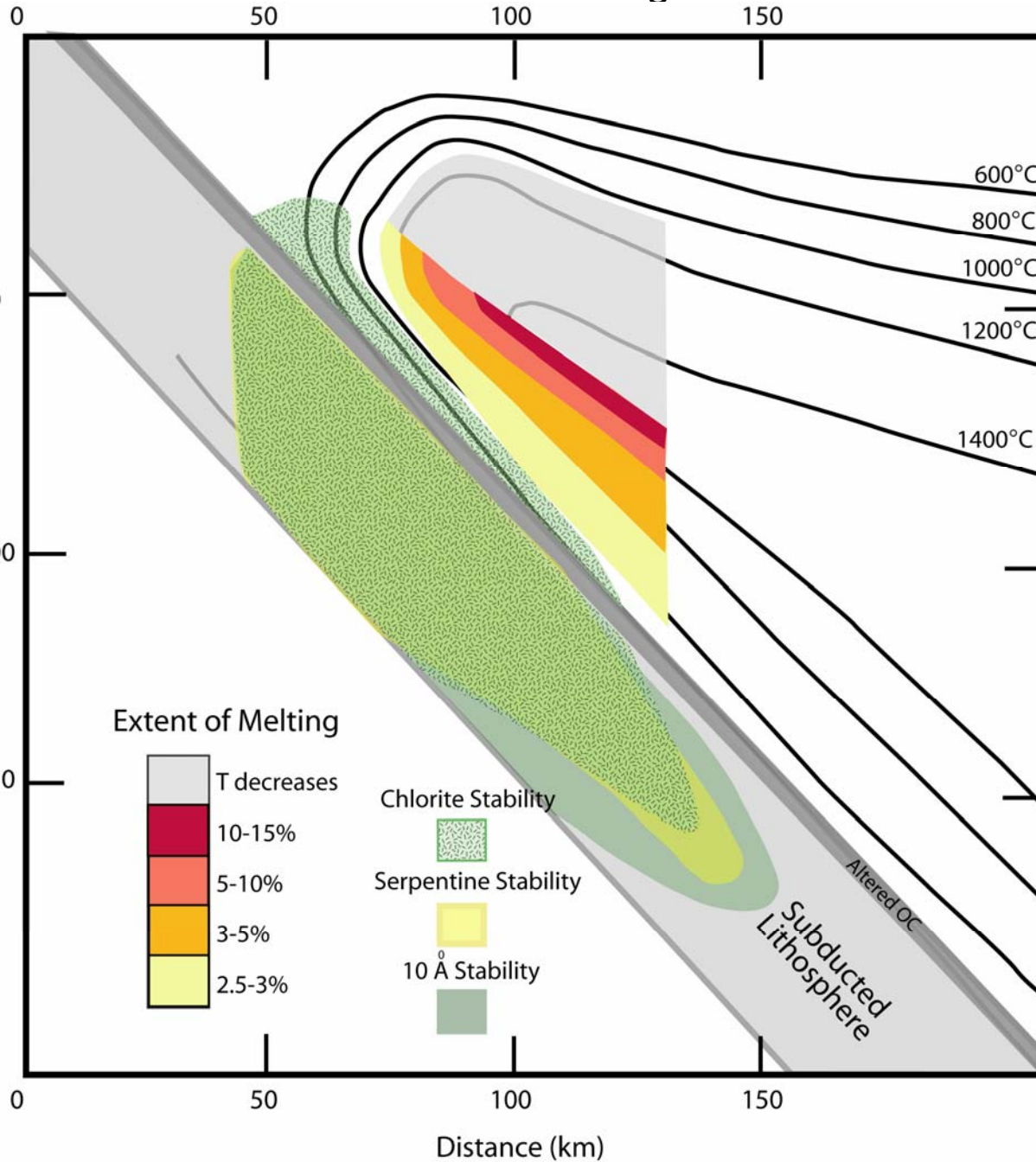


Medard & Grove (2006), Fumigali & Poli (2005), Pawley (2003)



Where is water stored in the wedge?

Thermal model of Kelemen et al. (2003)



Hydrous phases in the mantle wedge & subducted slab.

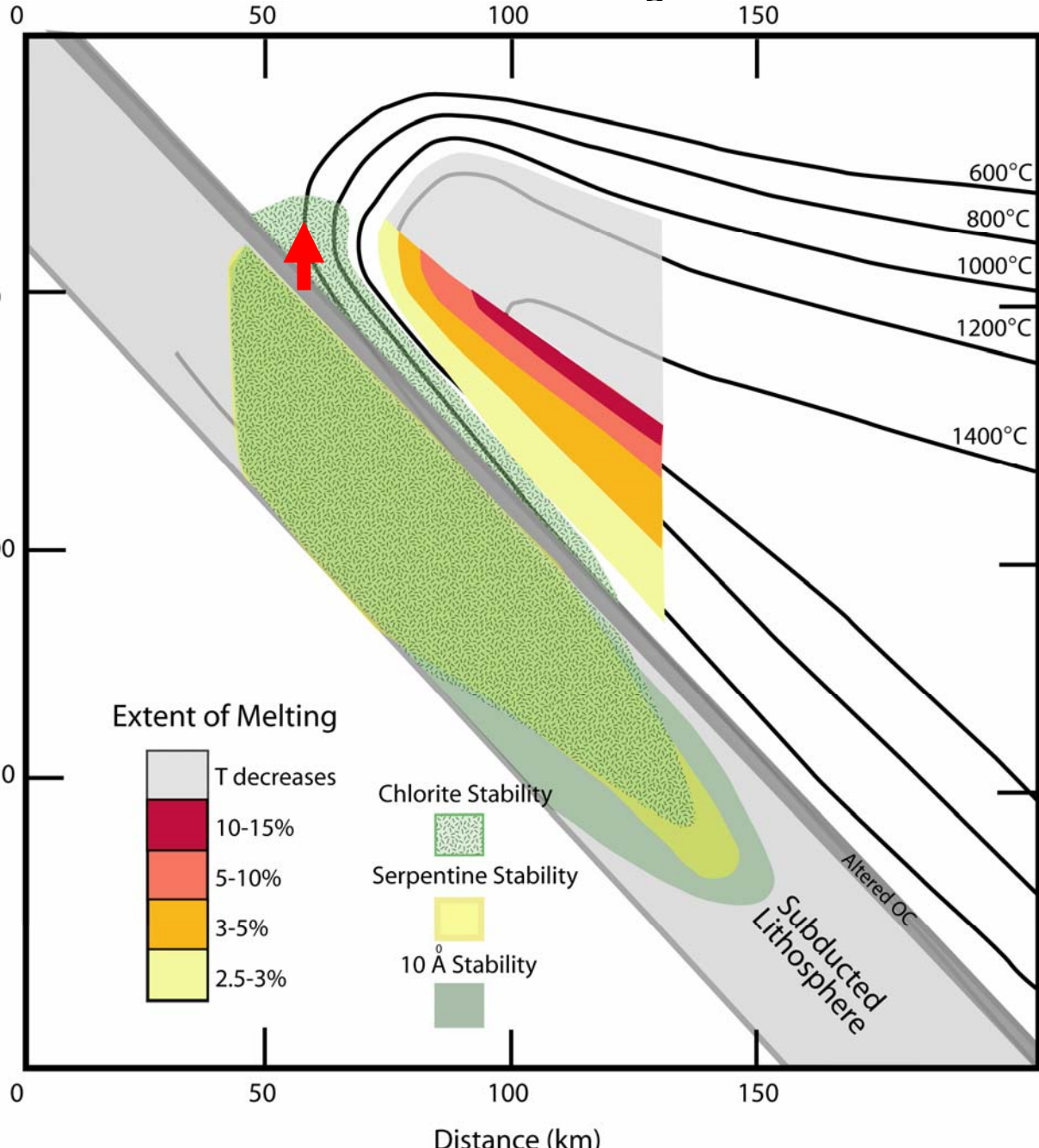
Chlorite provides a source of  $H_2O$  for wet arc melting that is above the slab.

Produced by fluid released from the slab at shallow depths.

$H_2O$  is stored even when the slab is too hot.

Chlorite also stable below the slab-wedge interface in the cool core of the slab.

# Where is water stored in the wedge?



Hydrous phases in the mantle wedge & subducted slab.

Chlorite provides a source of H<sub>2</sub>O for wet arc melting that is above the slab.

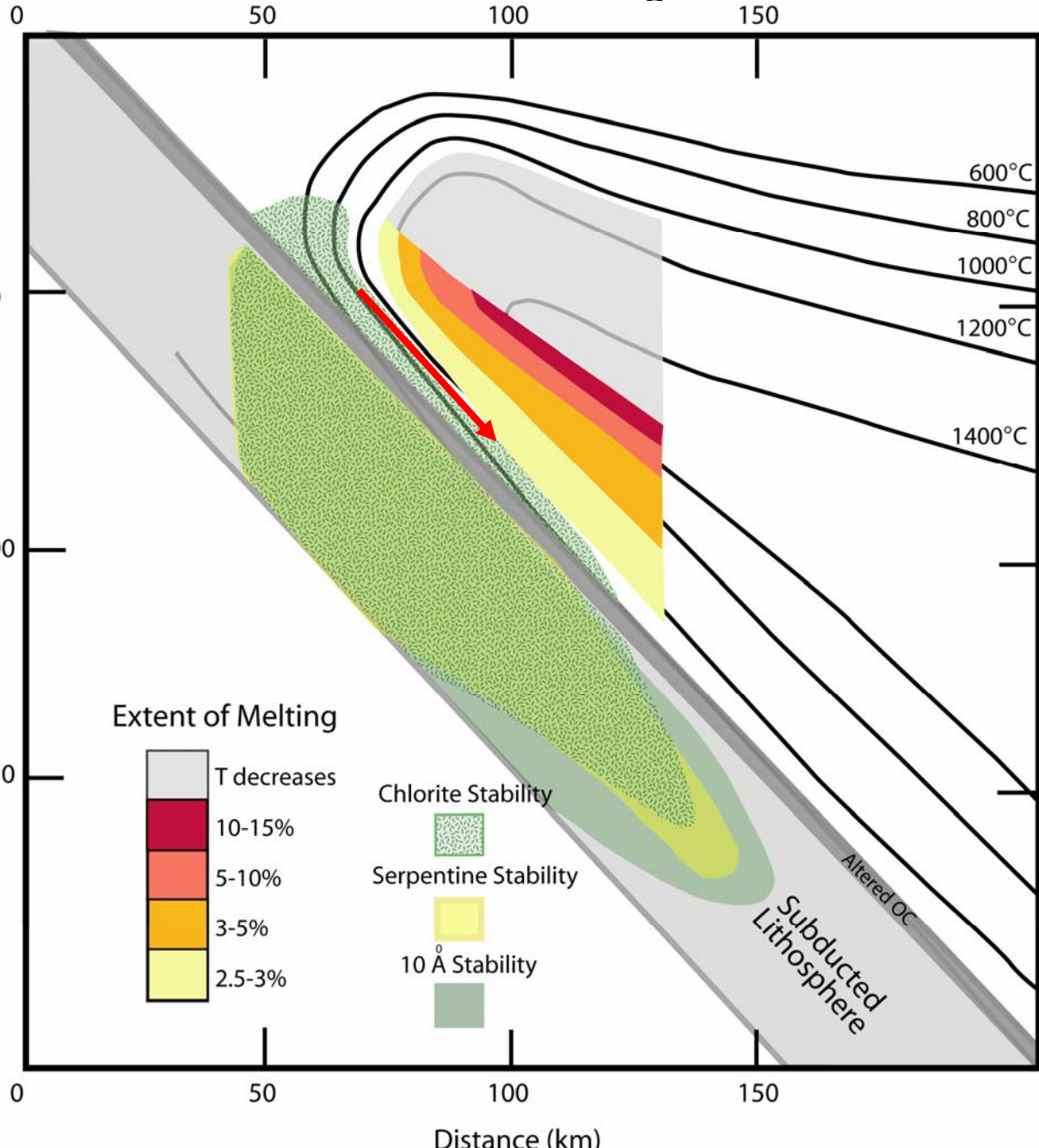
Produced by fluid released from the slab at shallow depths.

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# Where is water stored in the wedge?



Hydrous phases in the mantle wedge & subducted slab.

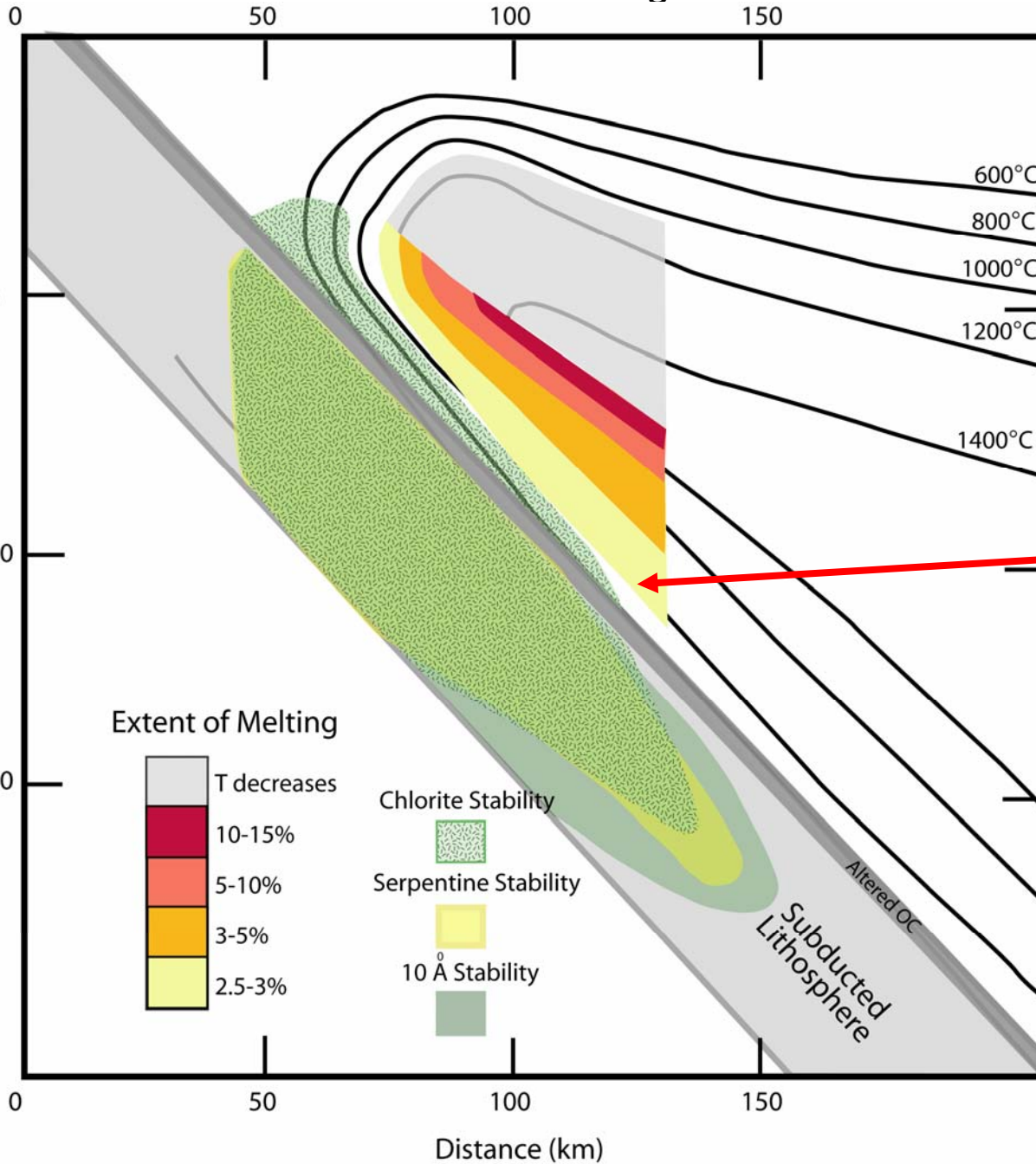
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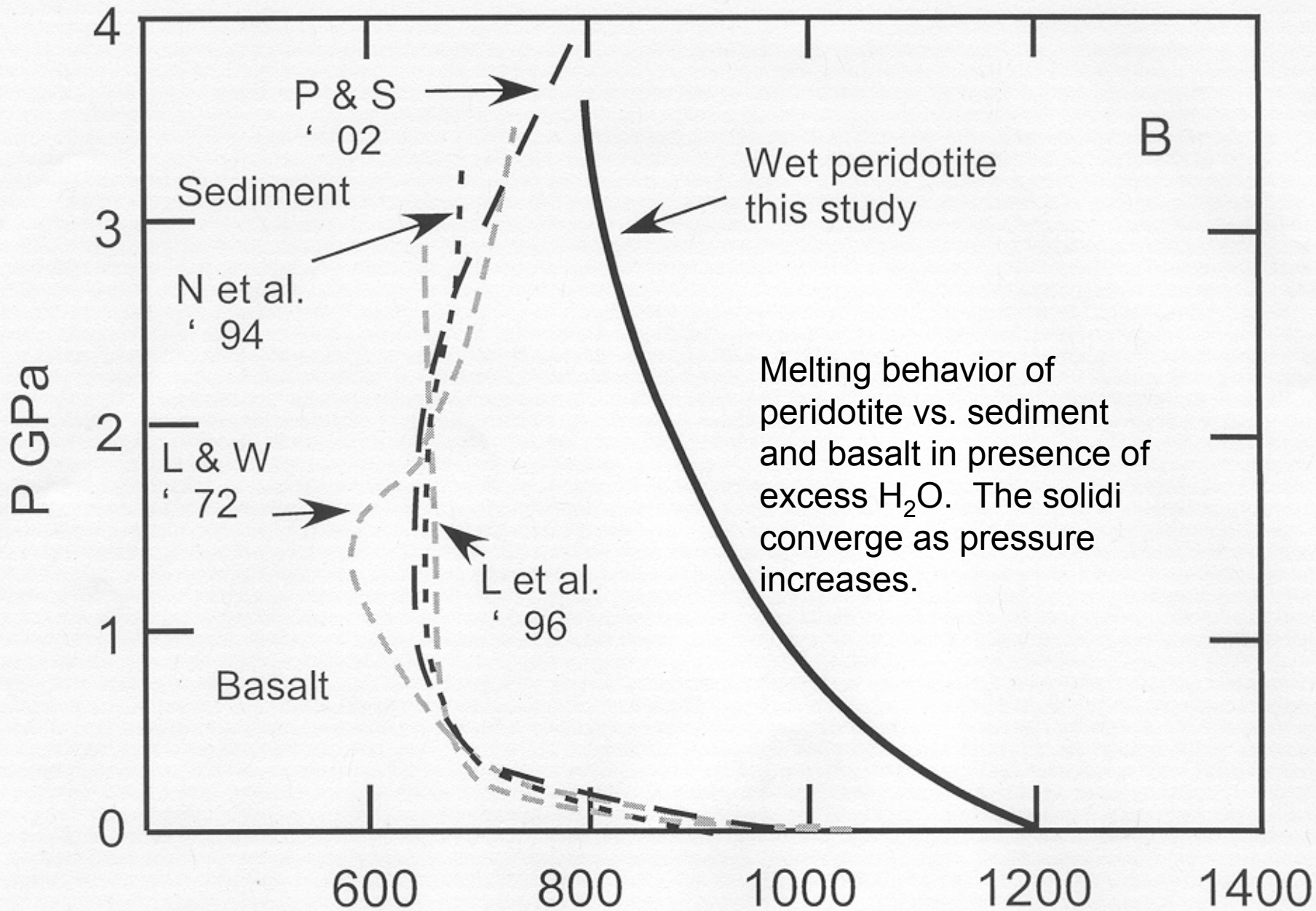
# Where is water stored in the wedge?



Chlorite breaks down as pressure increases.

Thus, the H<sub>2</sub>O supply through chlorite breakdown will stop at slab depths of 120 – 150 km.

Chlorite is stable below the slab-wedge interface in the cool core of the slab and this could supply H<sub>2</sub>O.





The melting model:

Melting paths calculated wherever vapor-saturated melting could occur – no assumptions about melt connectivity Mibe et al. (1999).

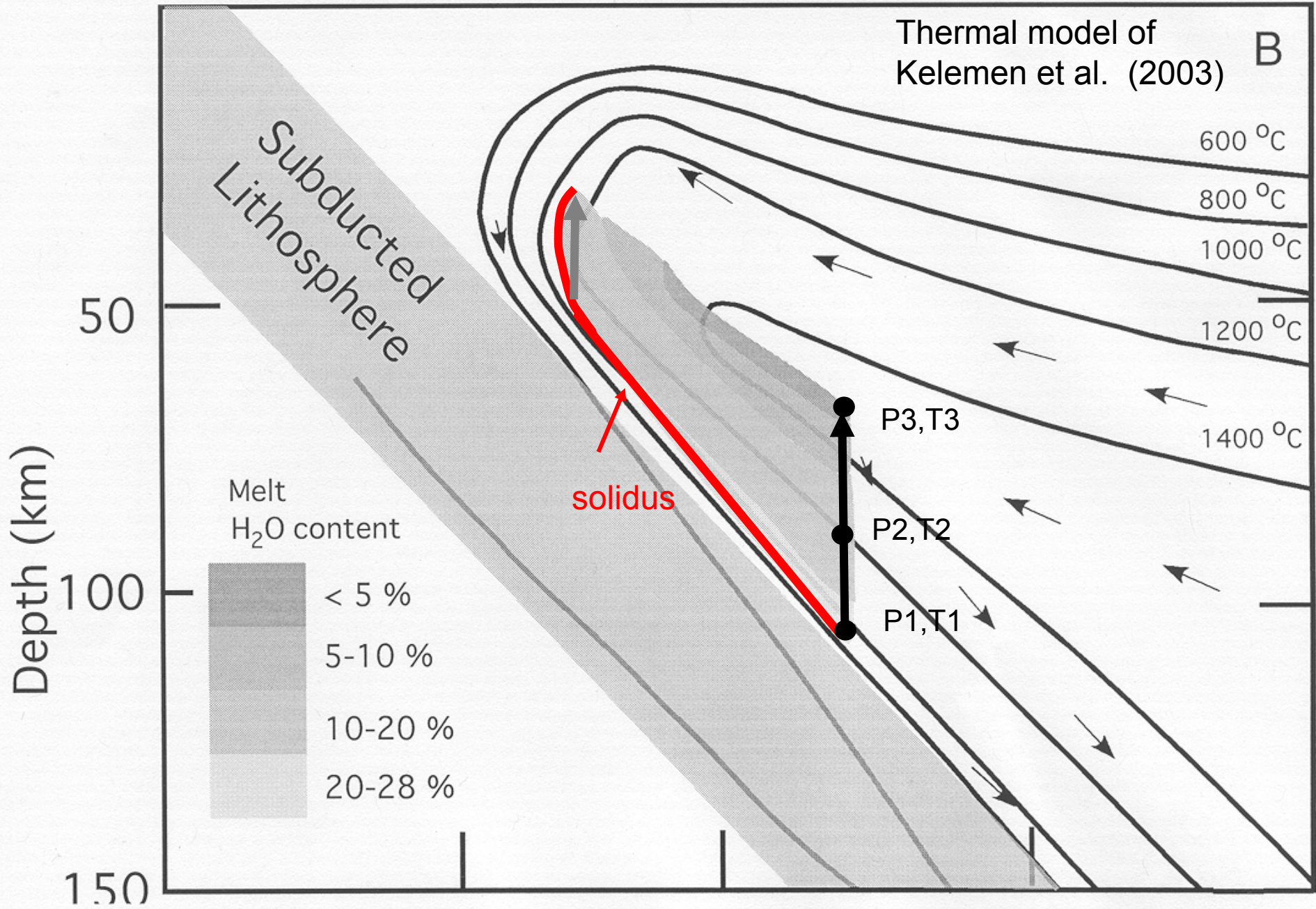
Buoyant hydrous melts leave the base of wedge and ascend into the overlying mantle by porous flow.

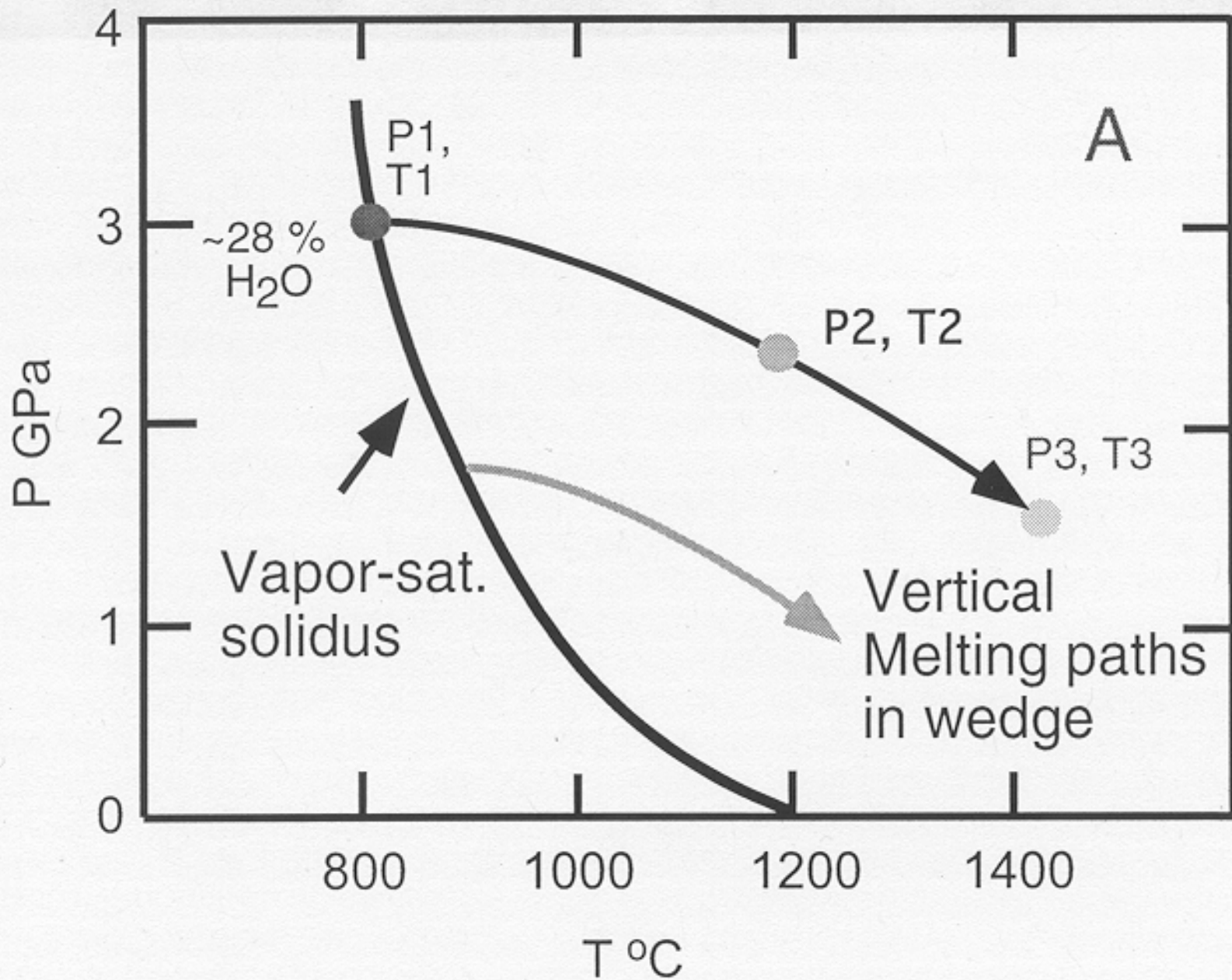
Melt volume equilibrates with mantle at each step –both thermally and chemically – **reactive porous flow**

Assumptions: initial critical melt fraction –

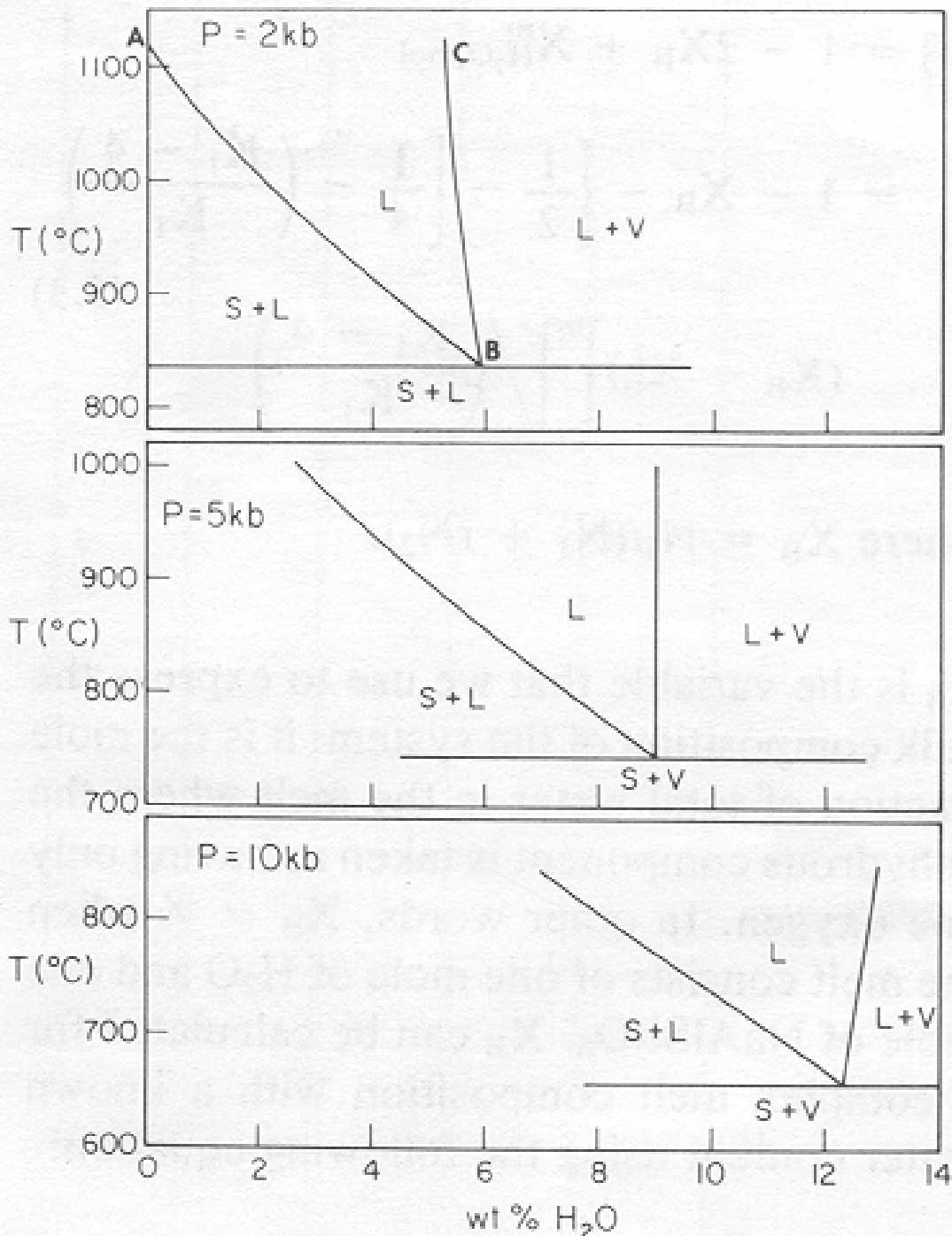
$F_{crit} = 2.5$  wt. % values range from  $< 0.1$  (Kohlstedt, 1992) to 8 % Fujii et al. (1986)

Thermal model of  
Kelemen et al. (2003) **B**







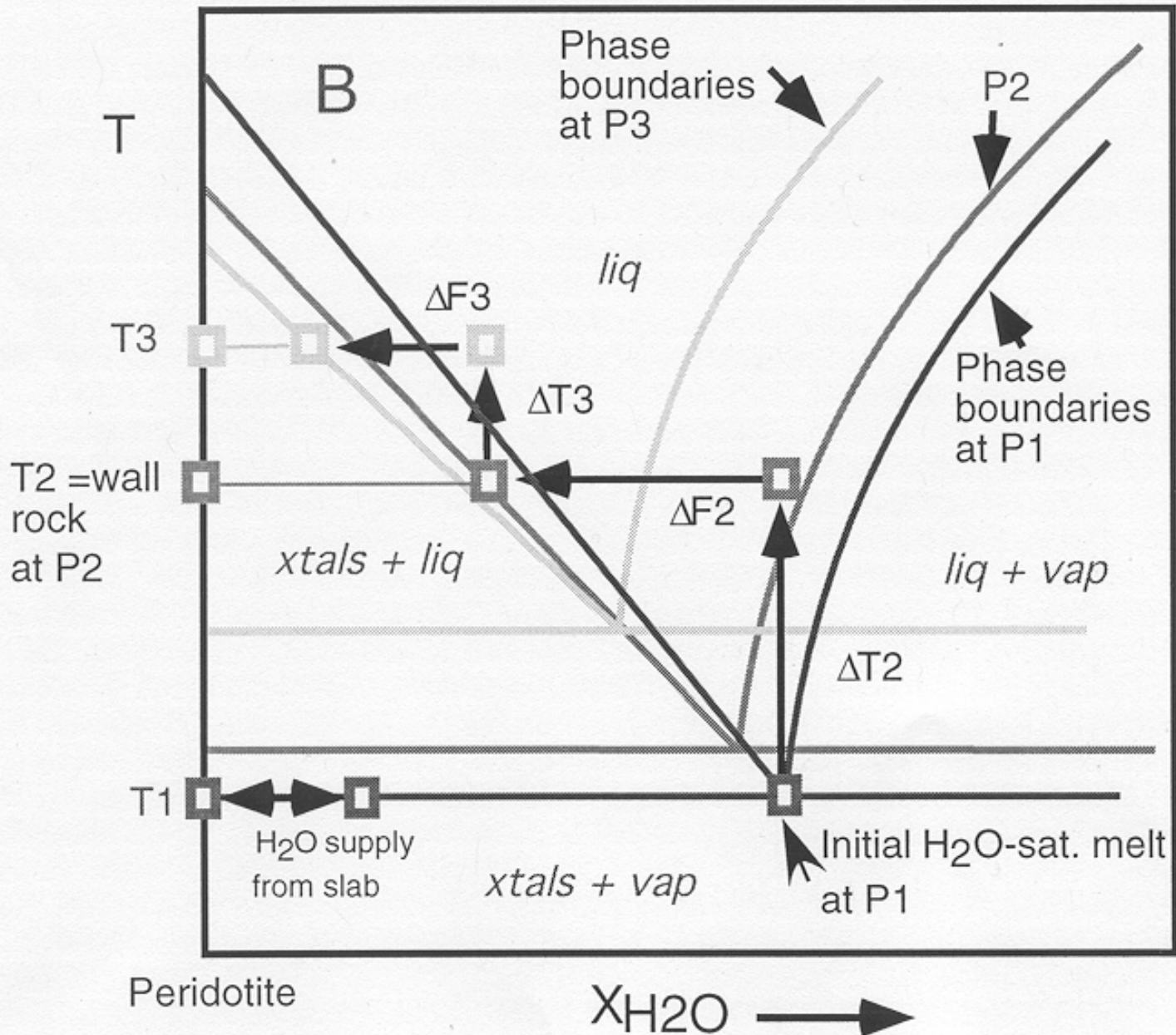


Silver and Stolper (1985) speciation model for melting in simple two component systems mineral – H<sub>2</sub>O

Includes molecular H<sub>2</sub>O – OH speciation and leads to a planar T – P – X<sub>H<sub>2</sub>O</sub> solid – melt boundary

Note linearity of liquidus boundary.

This melting behavior is “adjusted” for peridot.



Melt H<sub>2</sub>O content is strongly influenced by the lever rule effect –

melt % increases rapidly as bulk composition

Approaches liquidus.

Reactive porous flow melting = melt volume must come into equilibrium with mantle at each step –both thermally and chemically

The melting model:

We use our phase diagram & measured H<sub>2</sub>O solubility vs. pressure in forsterite – H<sub>2</sub>O to predict the peridotite – melt boundary in T – P – XH<sub>2</sub>O space. The expression is:

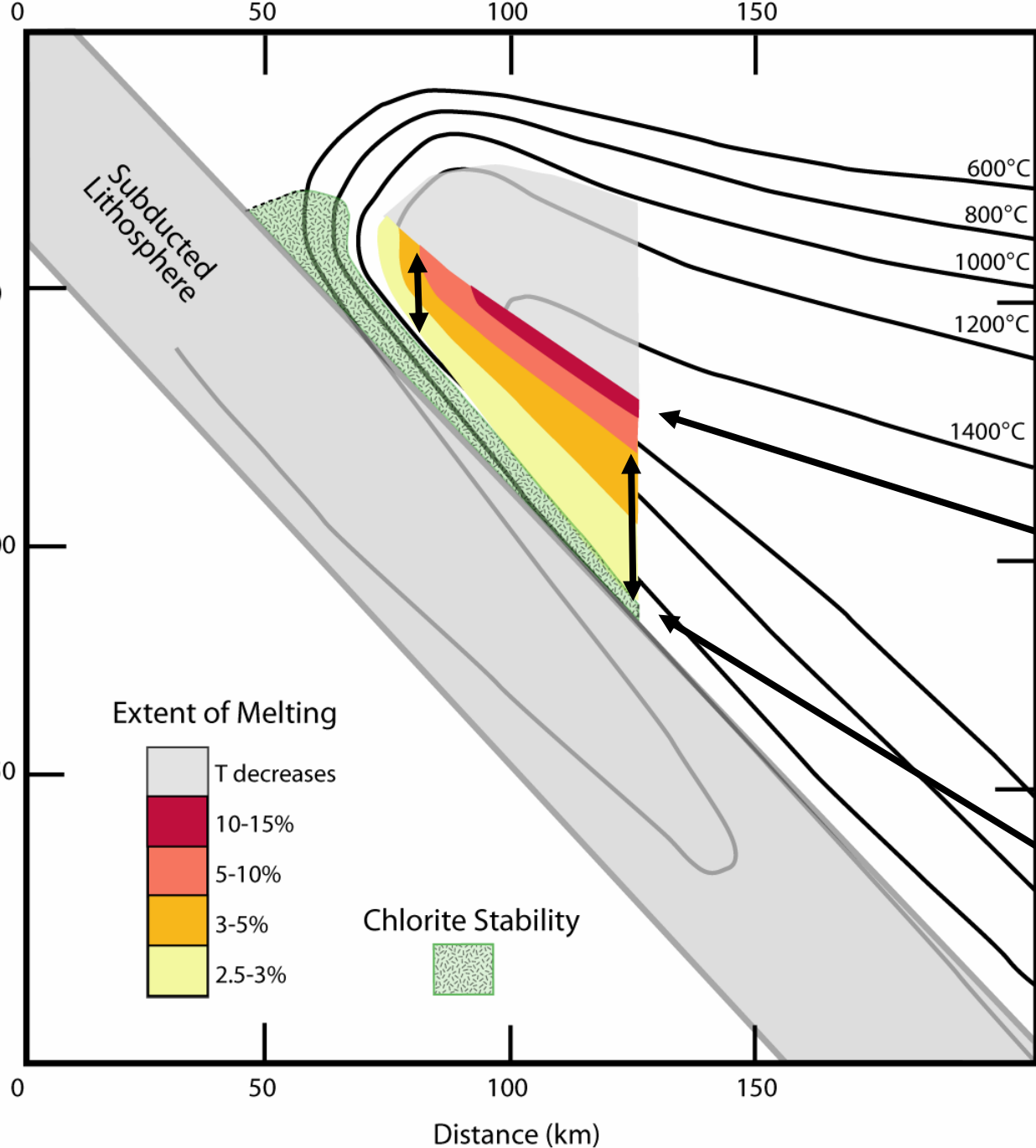
$$7290 * P - 810 * T - 24600 * H_2O + 1093500 = 0$$

where T is in °C, P is in kilobars and H<sub>2</sub>O content is in wt. %.

At P<sub>2</sub>, T<sub>2</sub> the amount of melt (F<sub>P<sub>2</sub>,T<sub>2</sub></sub>) is given by:

$$F_{P_2, T_2} = ((X_{init} - X_{P_2, T_2}) / X_{init}) * F_{init} + F_{init}$$



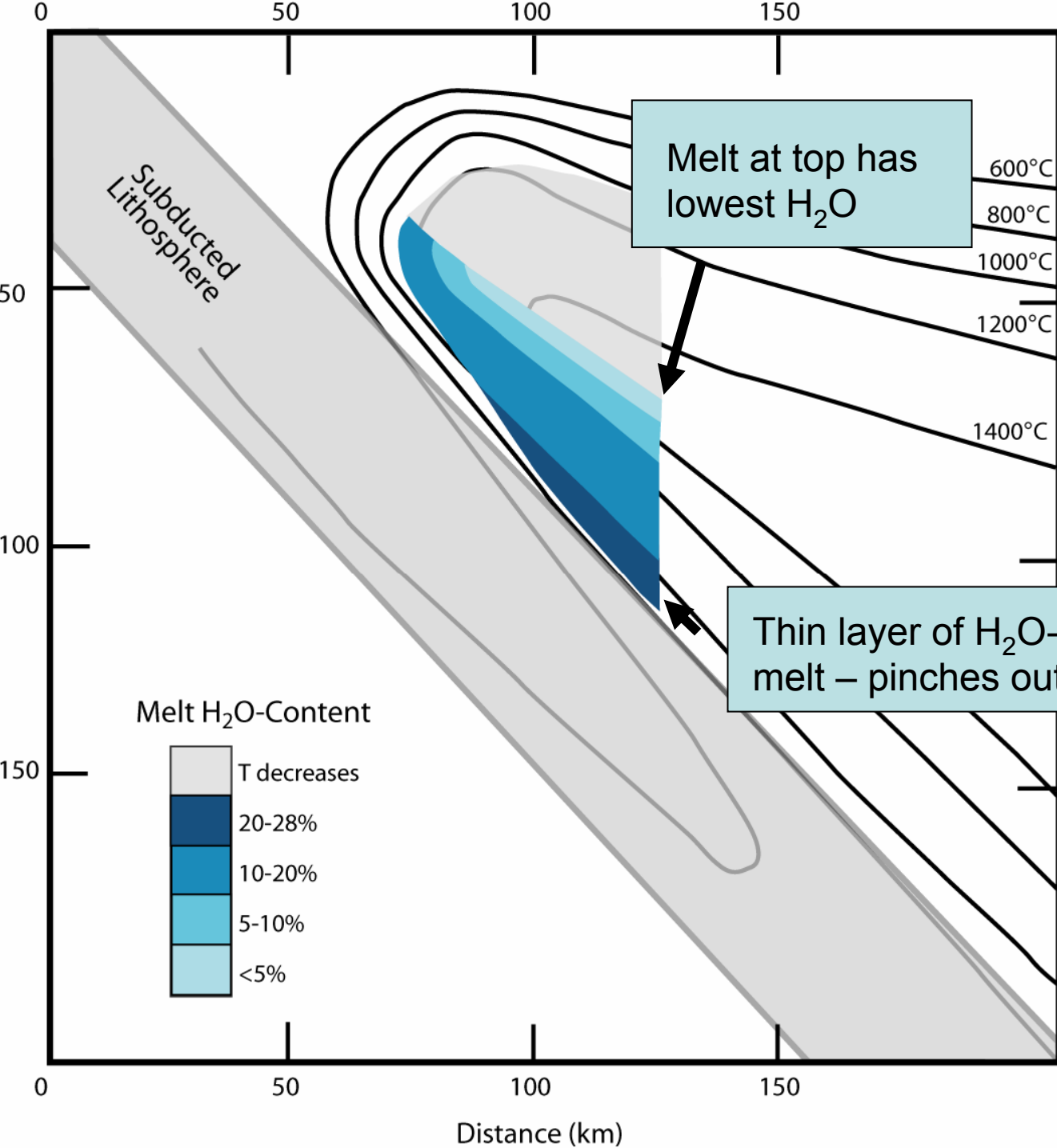


Maximum melt % occurs in a thin layer in hot core of the wedge

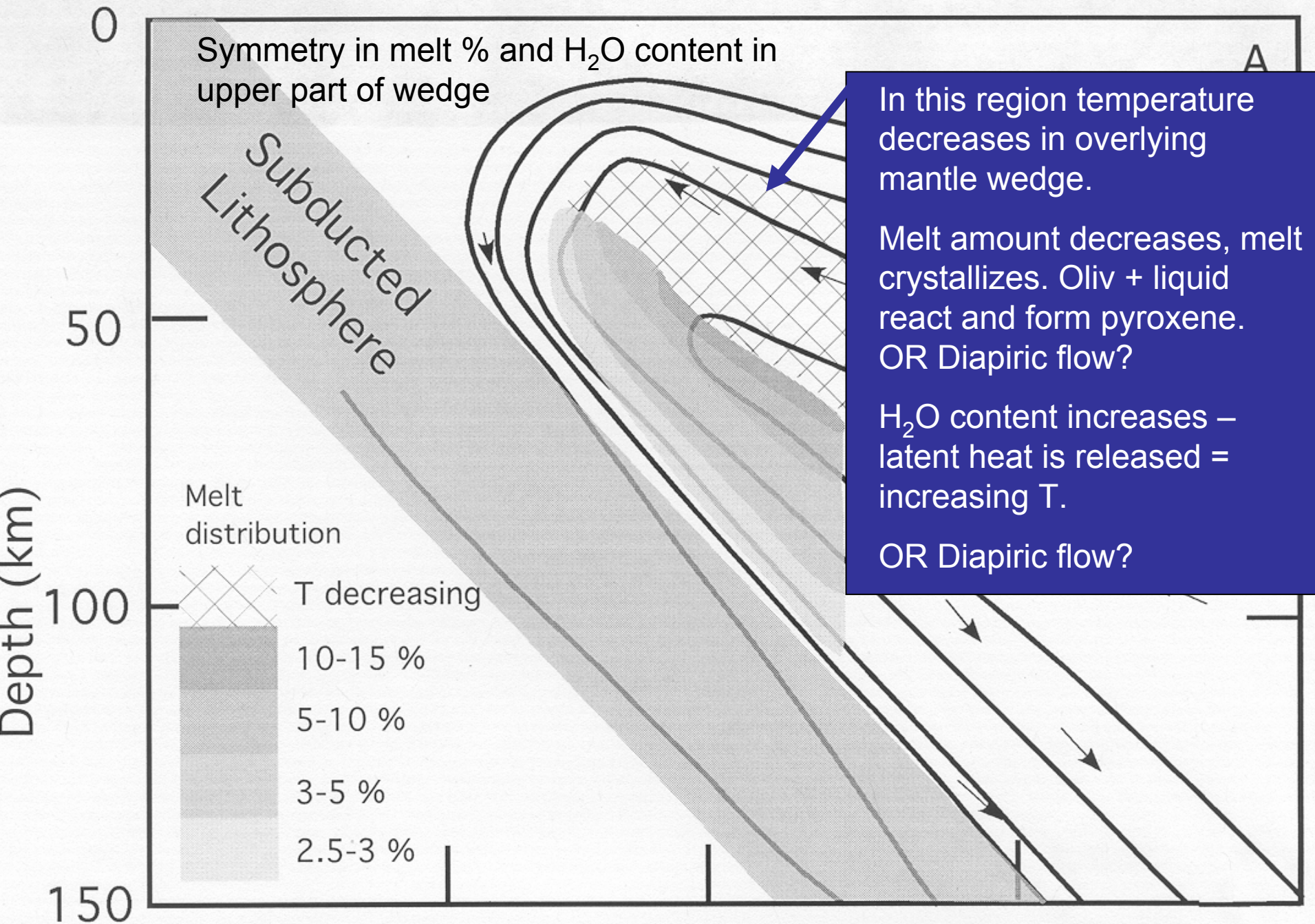
Highest melt fraction achieved in a very thin layer.

Most of wedge contains < 5% melt (double arrows).

Chlorite is transported down into the wedge ABOVE the slab and gives up its H<sub>2</sub>O at solidus.

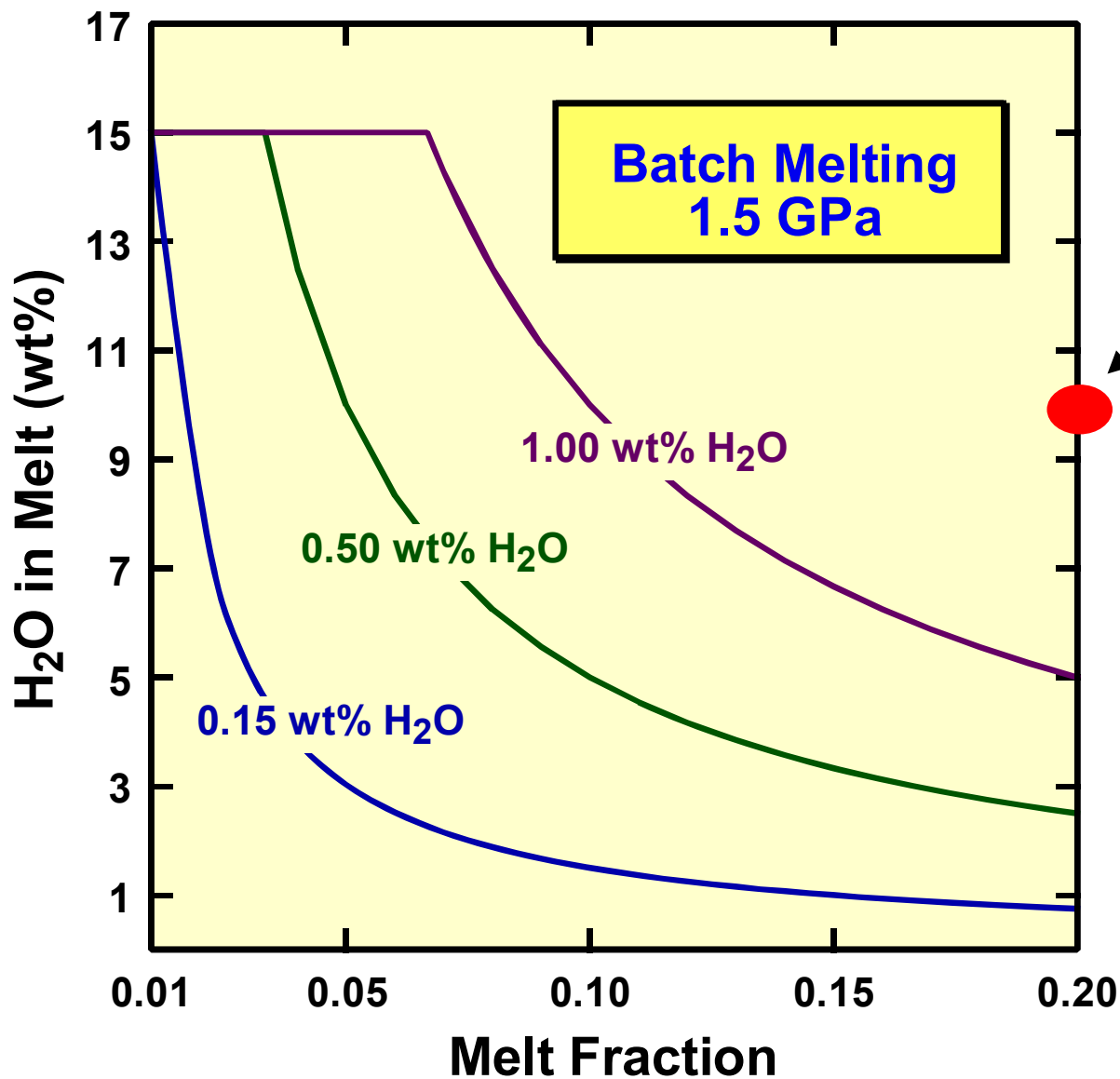


Melt H<sub>2</sub>O content continuously decreases as melt ascends and reacts w/ hotter mantle





When mantle peridotite is hydrated it contains 13 % chlorite. Bulk H<sub>2</sub>O of solid is 2 wt. % .



Bulk perid. = 2 wt. % H<sub>2</sub>O. H<sub>2</sub>O content of melt could exceed 10 wt. %.

High H<sub>2</sub>O contents are possible.

- Hydrous flux melting explains the shallow final equilibration depth of arc magmas AND provides a mechanism for creating  $\text{SiO}_2$  – rich crust through arc magmatic inputs.
- Stable chlorite in the mantle wedge allows for high  $\text{H}_2\text{O}$  content in arc magmas.
- At the same time subducted sediment and basalt can melt and transfer key trace element signatures to melts of mantle wedge.

Mt. Shasta, N. Calif. – looking W from Med. Lake

## *Experimental Details*

Au capsules – Piston cylinder - 1.2 – 3.2 GPa

Hart & Zindler Primitive Mantle

Oxide starting mix

With MgO added as  $\text{Mg}(\text{OH})_2 = 14 \text{ wt. \% H}_2\text{O}$

## *Run Duration*

96 – 140 hours ( a few at 24 hrs)

## *Experimental Products*

Homogeneous olivine, opx, cpx, spinel and/or garnet

Melt or vapor phase (supercritical fluid)

## *Equilibrium*

QUILF used to check Temp. from Opx-Cpx – within 1 sigma of uncertainty

and  $f_{\text{O}_2}$  from oliv-opx-spinel (Ballhaus et al., 1994) = QFM + 0.8



## PREVIOUS STUDIES

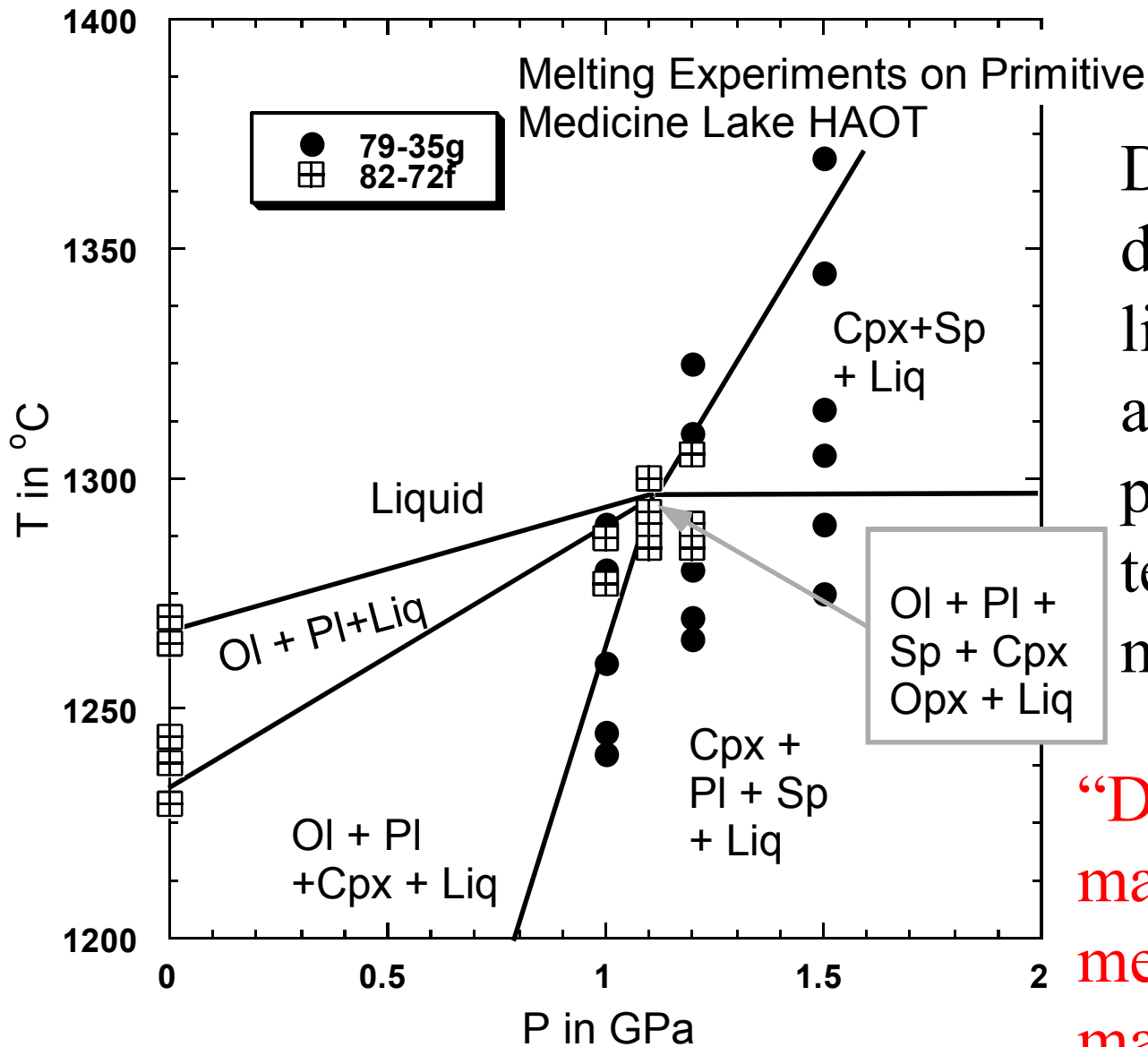
<i>Run Times</i>	<i>H<sub>2</sub>O added</i>	<i>Capsule</i>
<b>High – T melting</b>		
Kushiro et al. (1968)		
<i>5 – 30 min</i>	<i>30%</i>	<i>Mo &amp; Pt</i>
Millhollen et al. (1974)		
<i>0.5 – 3 hrs.</i>	<i>5.7 %</i>	<i>Pt</i>
Green (1973)		
<i>1 – 6 hrs.</i>	<i>10 %</i>	<i>AgPd alloy</i>

## **Low – T melting**

Mysen and Boettcher (1975)		
<i>24 – 64 hrs.</i>	<i>20- 30 %</i>	<i>AgPd alloy</i>

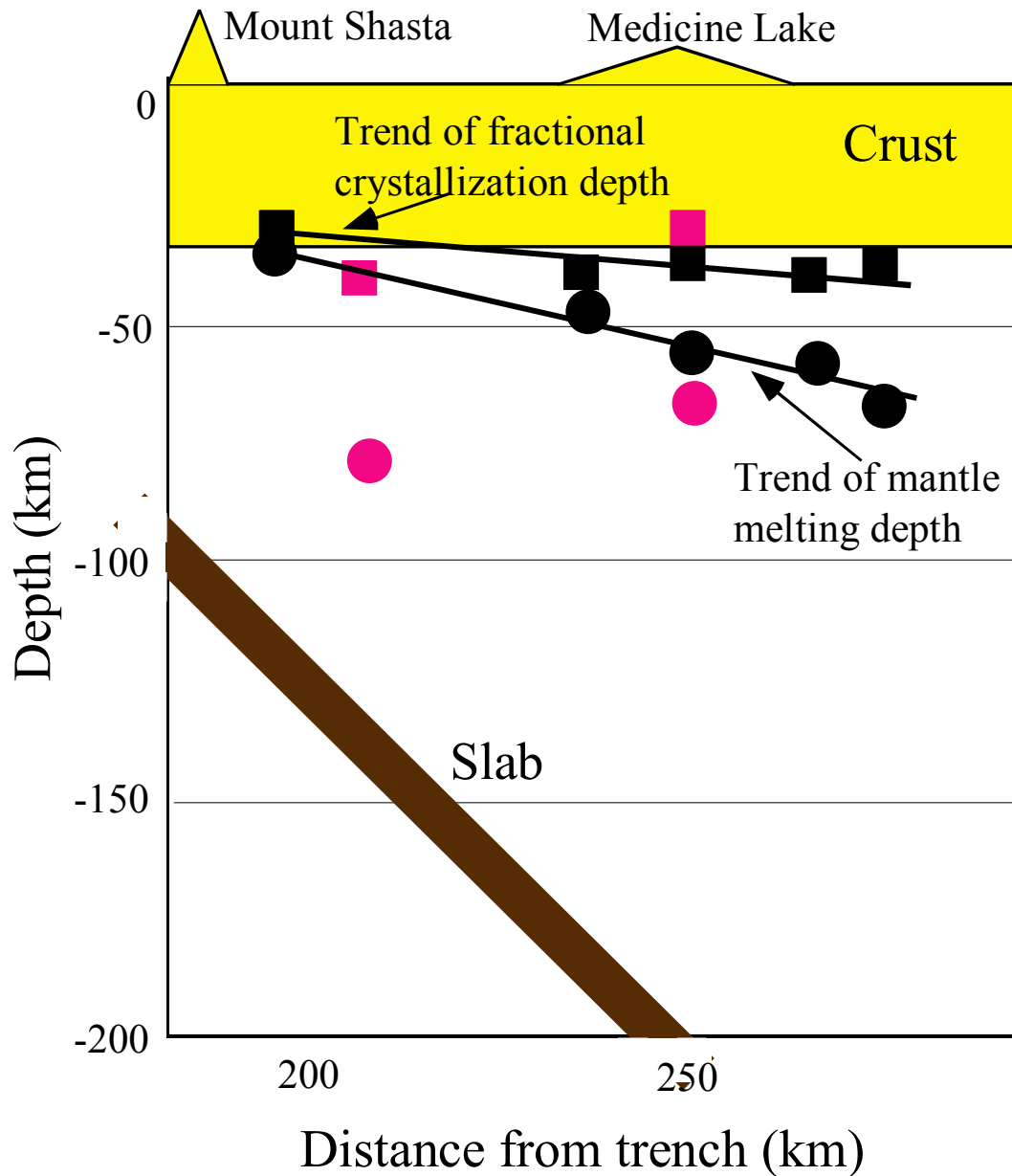
## THIS STUDY

<i>48 – 120 hrs</i>	<i>14 –30 %</i>	<i>Au</i>
---------------------	-----------------	-----------



Direct  
determination of  
liquidus mineral  
assemblage at high  
pressure and  
temperature =  
mantle conditions

“Dry” HAOT  
magmas record  
melting as convecting  
mantle is drawn into  
the wedge



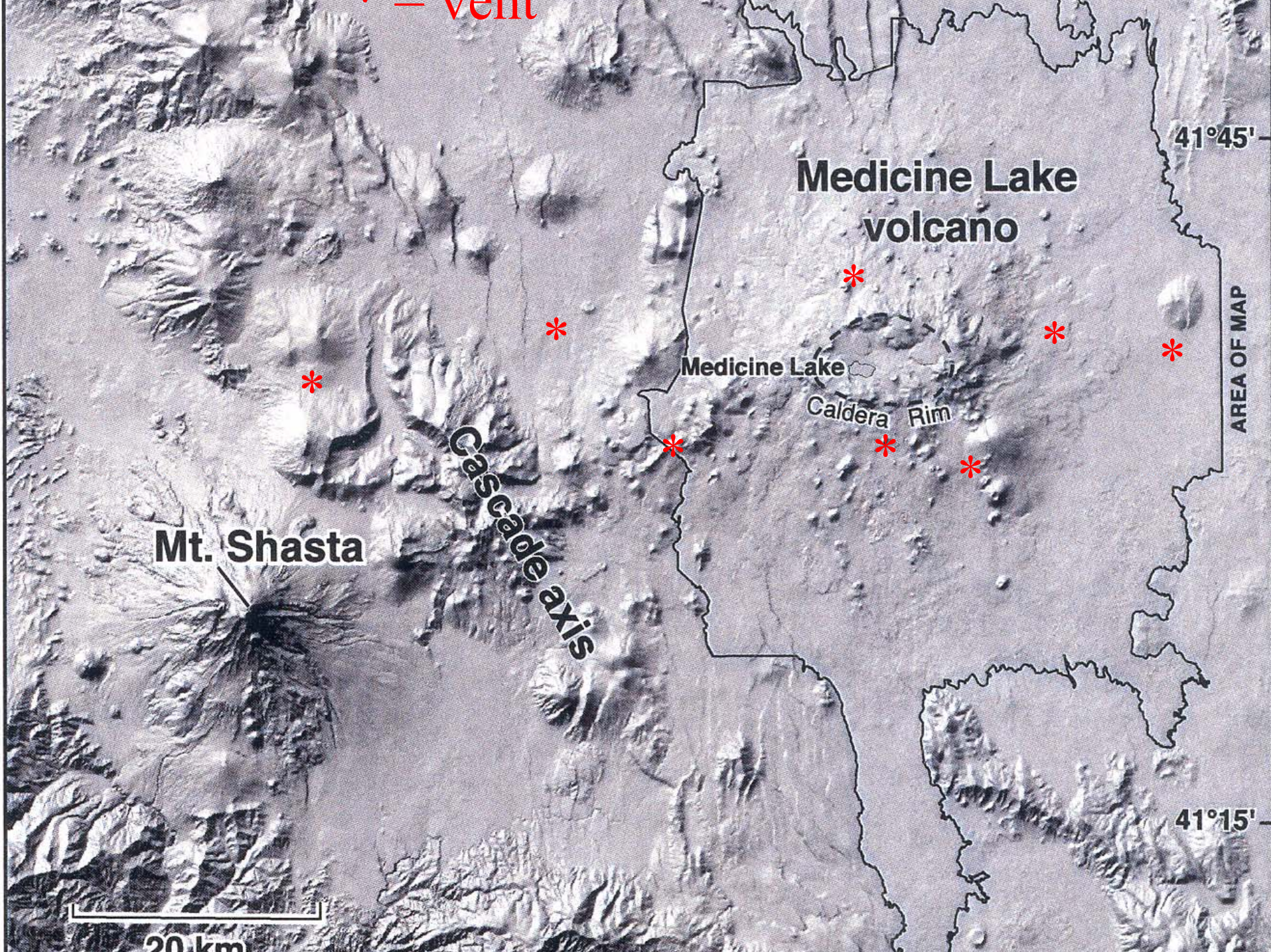
Shallow, hot mantle melting beneath the Cascades

Inferred from Pressure of multiple saturation.

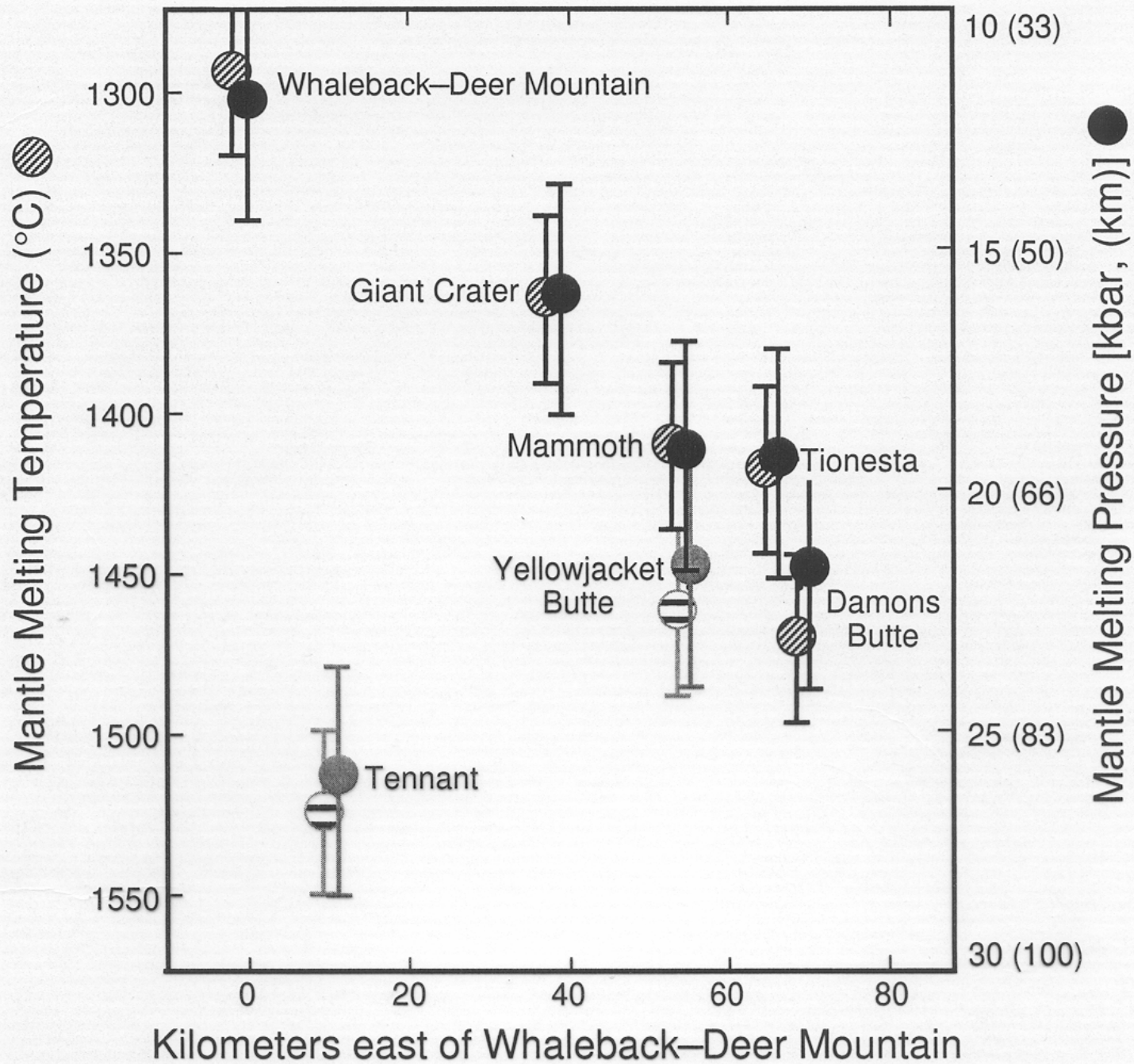
$T = 1300 - 1450 \text{ }^{\circ}\text{C}.$

Elkins-Tanton et al. (2001)









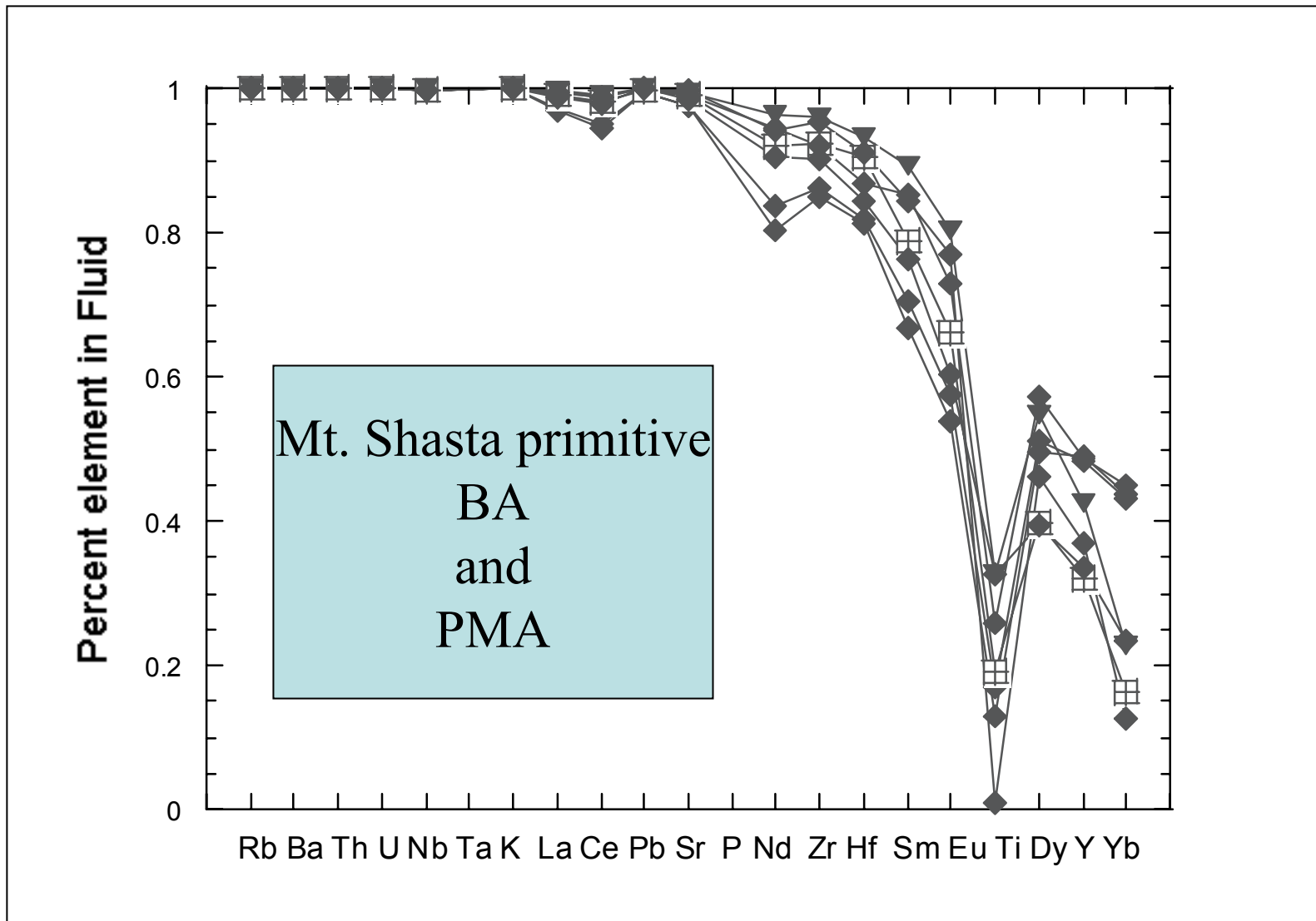
# Major element characteristics of the fluid-rich Mt. Shasta component

- $\text{Na}_2\text{O}$  = 25 to 33 wt.% of the “fluid”
- $\text{K}_2\text{O}$  = 5 to 13 wt. % of “fluid”
- $\text{SiO}_2$  = 0 wt. %
- $\text{H}_2\text{O}$  = 54 to 70 wt. %
- Similar to finding of Stolper & Newman (1994).

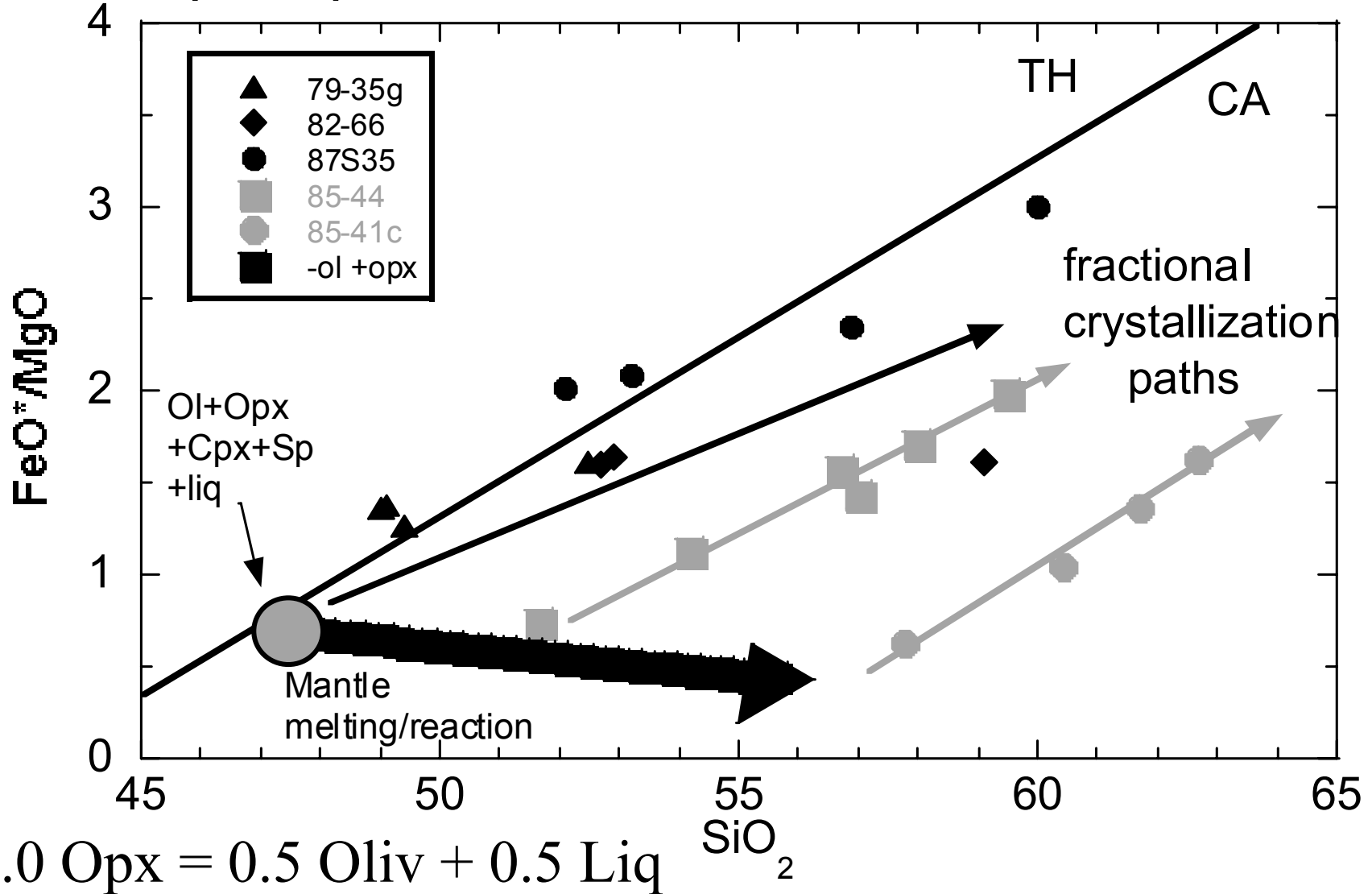
So, what is it? A melt or a fluid?

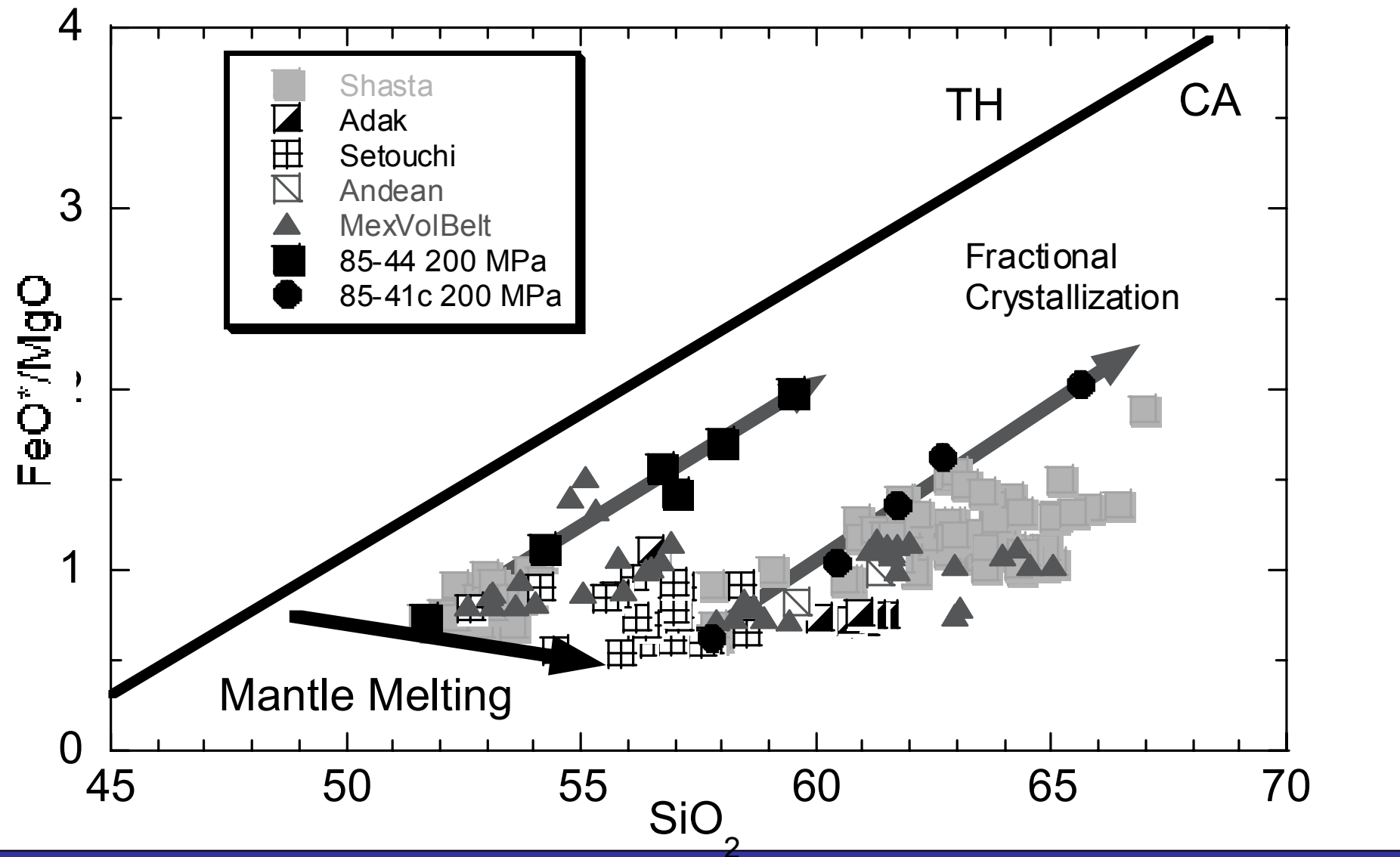


# Estimated Slab Contribution



Mantle melting trend to high-SiO<sub>2</sub> - low FeO\*/MgO is controlled by the reaction relation  
 oliv + liq → opx.





Shasta  $\text{H}_2\text{O}$ -rich lavas have high  $\text{SiO}_2$  and low  $\text{FeO}^*$ , similar to adakites and Japanese sanukitoids: characteristics inherited from low-P mantle melting.