

Curriculum Vitae

1988	Baccalauréat C (Science)
1989 – 1991	Math Sup - Math Spe P (Highschool preparatory course)
1991 – 1993	European Highschool of Industrial Chemistry of Strasbourg
1993 – 1994	Licence of Chemistry – University Paris VI
1994 – 1995	Maitrise of Chemistry – University Paris VI Options: Analytical chemistry and Environment
1995 – 1996	D.E.A. of analytical chemistry
1996 – 1997	Ph'D - University of Orleans – Geological Department
1997 – 2002	Ph'D – University of Bayreuth – BGI
2003 – 2006	Pharma consultant
2006 – 2009	Maternity leave

Practicals

EHIC'S practical

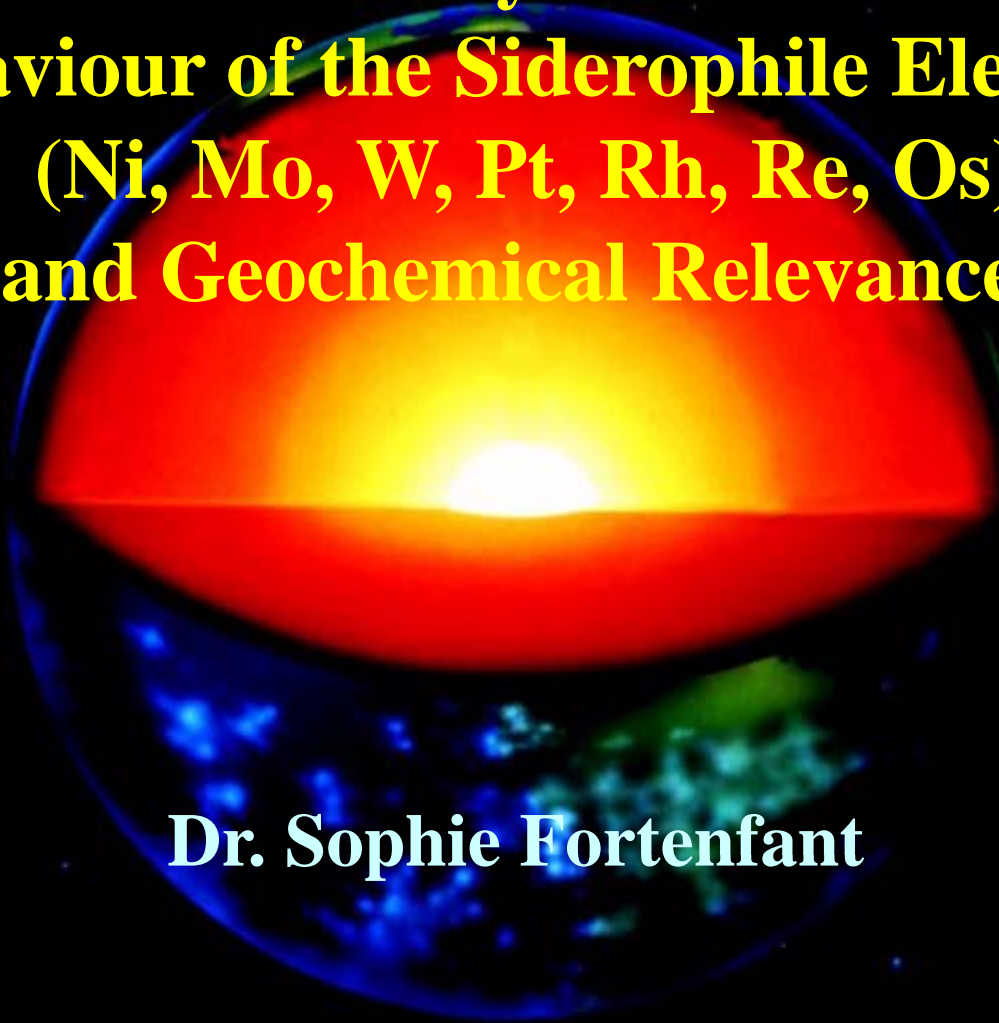
- Qualitative and quantitative analysis of a slag (Analytical laboratory - EHIC'S)
- Quality control of the secondary water at the nuclear plant of Dampierre

Maitrise practical

- Optimisation of an electrodeposition technique for zinc and cadmium (Ecole Centrale Paris)

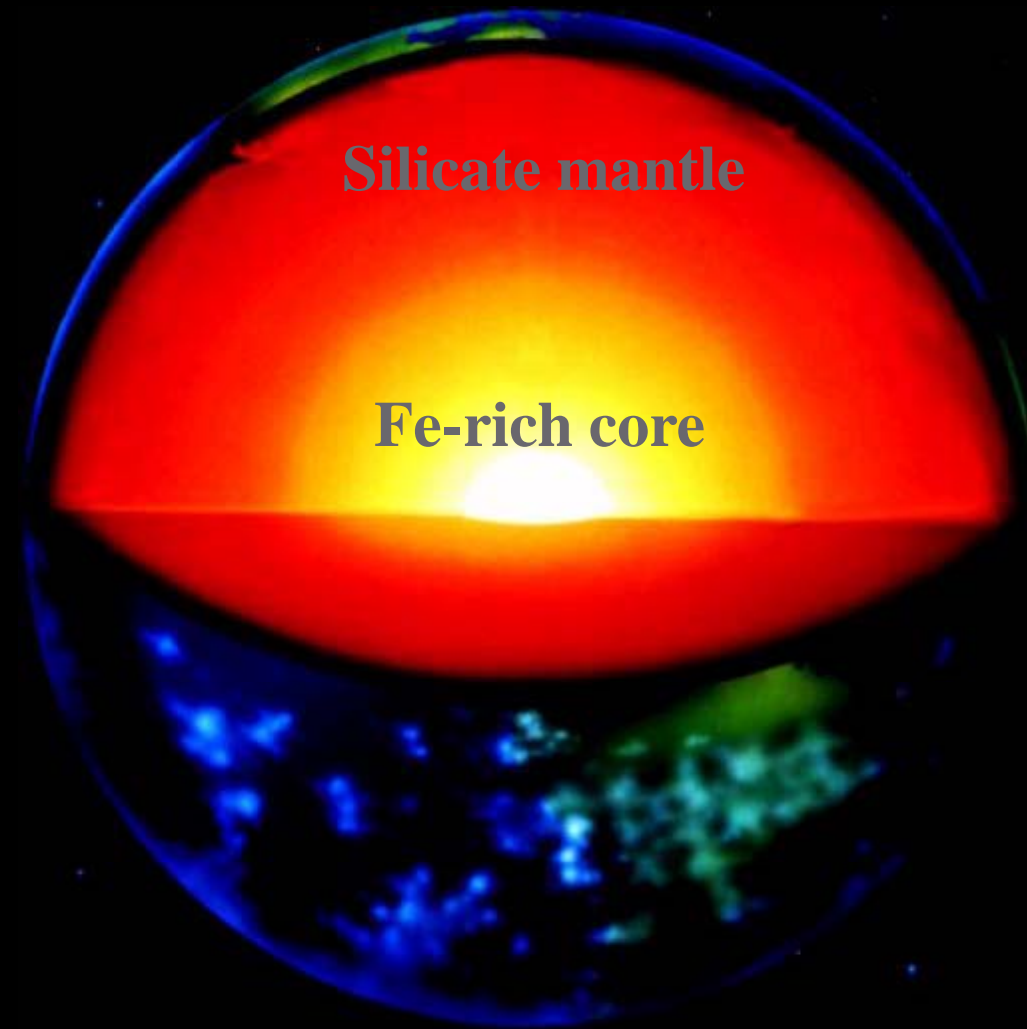
D.E.A practical

- Investigation of the optimal physico-chemical conditions for the electrodeposition of actinides in the presence of boron and calcium (Ecole Centrale Paris)



**Experimental Study on the Partitioning
Behaviour of the Siderophile Elements
(Ni, Mo, W, Pt, Rh, Re, Os)
and Geochemical Relevance**

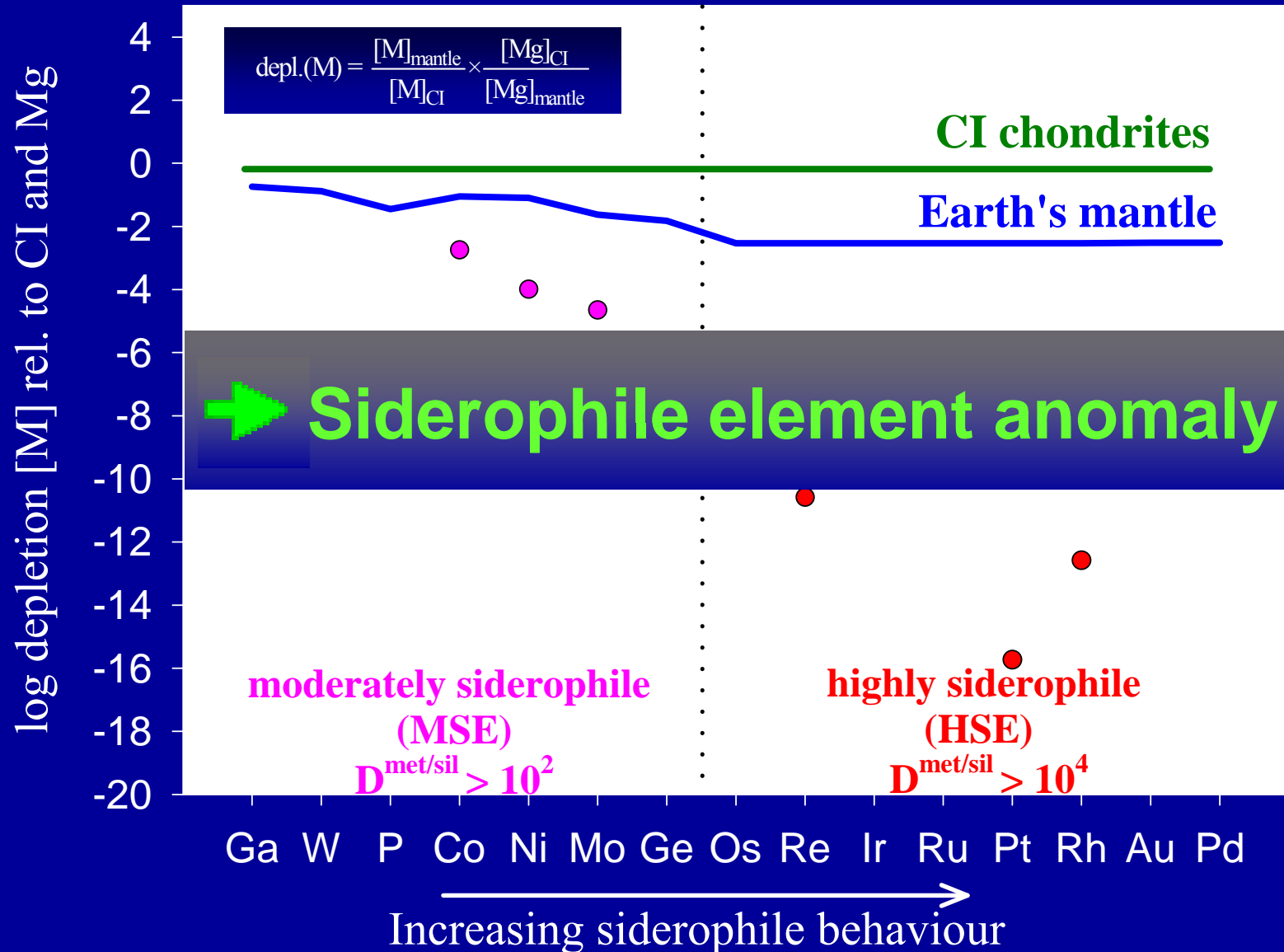
Dr. Sophie Fortenfant



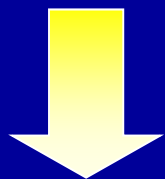
Silicate mantle

Fe-rich core

Siderophile element depletion



Earth's accretion and core formation models



Late Veneer

O'Neill (1991)

- ✚ Heterogeneous accretion
- ✚ The mantle abundances of the MSE is the consequence of a multi-stage core-mantle equilibrium process.
- ✚ The elevated and chondritic mantle abundances of the HSE is explained by the addition of a late veneer of chondritic material after core segregation.

Magma Ocean

Li and Agee (1996)
Righter and Drake (1997)

- ✚ Homogeneous accretion
- ✚ The abundances of the siderophile elements in the Earth's mantle result from core-mantle equilibrium at high pressure and high temperature in a deep magma ocean



Experimental solubility studies

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graph TD; A[Experimental solubility studies] --> B[One-bar studies]; A --> C[HP-HT studies]; B --> D[Loop technique]; B --> E[Stirred crucible technique]; C --> F[Multi-Anvil]; C --> G[Piston cylinder];
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The diagram is a hierarchical flowchart on a dark blue background. At the top is a yellow box with the text 'Experimental solubility studies'. Two yellow arrows point downwards from this box to two more yellow boxes: 'One-bar studies' on the left and 'HP-HT studies' on the right. From 'One-bar studies', two yellow arrows point to 'Loop technique' and 'Stirred crucible technique'. From 'HP-HT studies', two yellow arrows point to 'Multi-Anvil' and 'Piston cylinder'.

**One-bar
studies**

**HP-HT
studies**

**Loop
technique**

**Stirred
crucible
technique**

**Multi-
Anvil**

**Piston
cylinder**

$$\log [M] = \frac{x}{4} \times \log fO_2 + \frac{a}{T} + \frac{b \times P}{T} + c \times nbo / t + d$$



Degree of polymerisation of the melt
($0 < nbo/t < 4$)

Oxygen fugacity

Temperature

Pressure

Previous work

-  MSE: the effects of fO_2 , T and P on the solubility of these elements in melts are well defined at both low and high pressure, but the compositional effect is not yet well constrained.
-  HSE: because of their low solubilities in melts, experimental studies are more difficult. Low-pressure results are often scattered and need to be improved. High-pressure results are rare in the literature.

Experimental solubility studies

One-bar studies

HP-HT studies

Loop technique

Stirred crucible technique

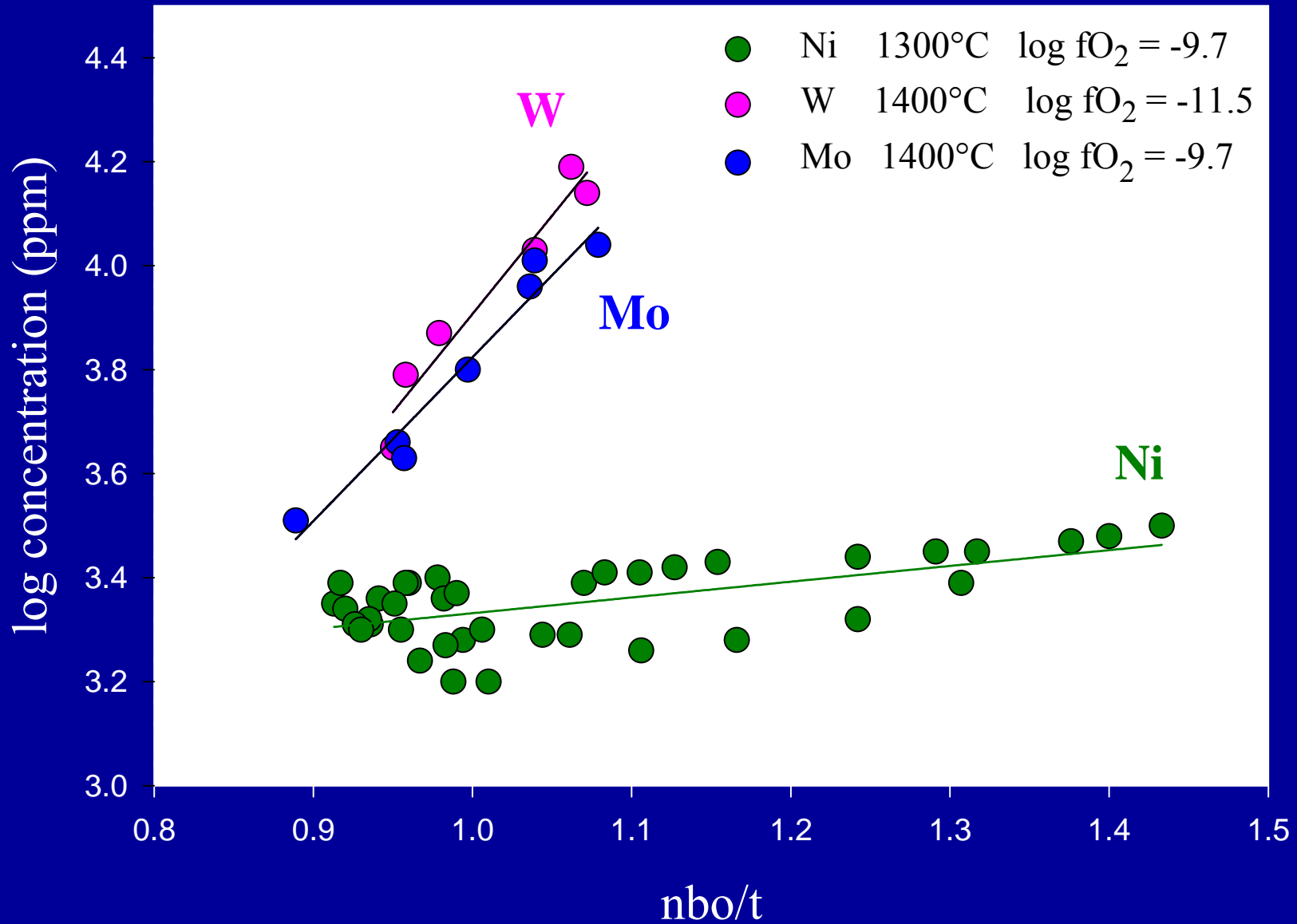
Multi-Anvil

Piston cylinder

Compositional effect on the f_{O_2} dependence of the T dependence of the

Effects of P and T on the partitioning of Re and Os between liquid metal and magnesio-wüstite

Compositional effect on the solubilities of Ni, Mo and W at 1 bar



Compositional effect on the solubilities of Ni, Mo and W at 1 bar

Basaltic melt
nbo/t ~ 0.9

$$D_{\text{Ni}}^{\text{met/sil}} = \approx 10^2$$

$$D_{\text{Mo}}^{\text{met/sil}} = \approx 10^3$$

$$D_{\text{W}}^{\text{met/sil}} = \approx 10^2$$

Peridotitic melt
nbo/t ~ 2.5

$$D_{\text{Ni}}^{\text{met/sil}} \downarrow 0.5 \text{ log-unit}$$

$$D_{\text{Mo}}^{\text{met/sil}} \downarrow 2 \text{ log-units}$$

$$D_{\text{W}}^{\text{met/sil}} \downarrow 2 \text{ log-units}$$

Assuming core-
mantle equilibrium



$$D_{\text{Ni}}^{\text{met/sil}} = 15$$

$$D_{\text{Mo}}^{\text{met/sil}} = 53$$

$$D_{\text{W}}^{\text{met/sil}} = 8$$

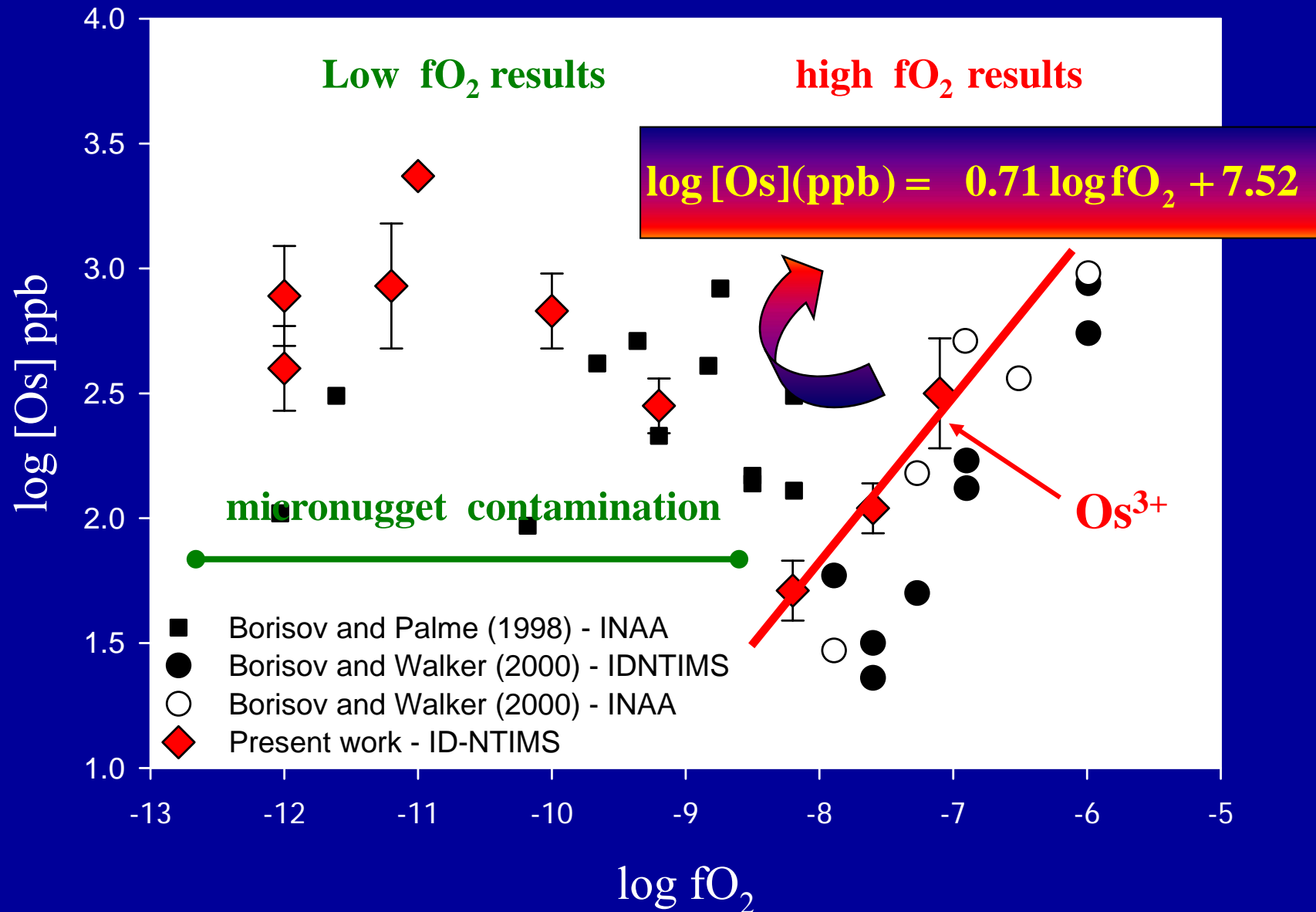


Ni, Mo and W abundances in the Earth's mantle could result from HP-HT equilibrium between peridotitic melt and metal.

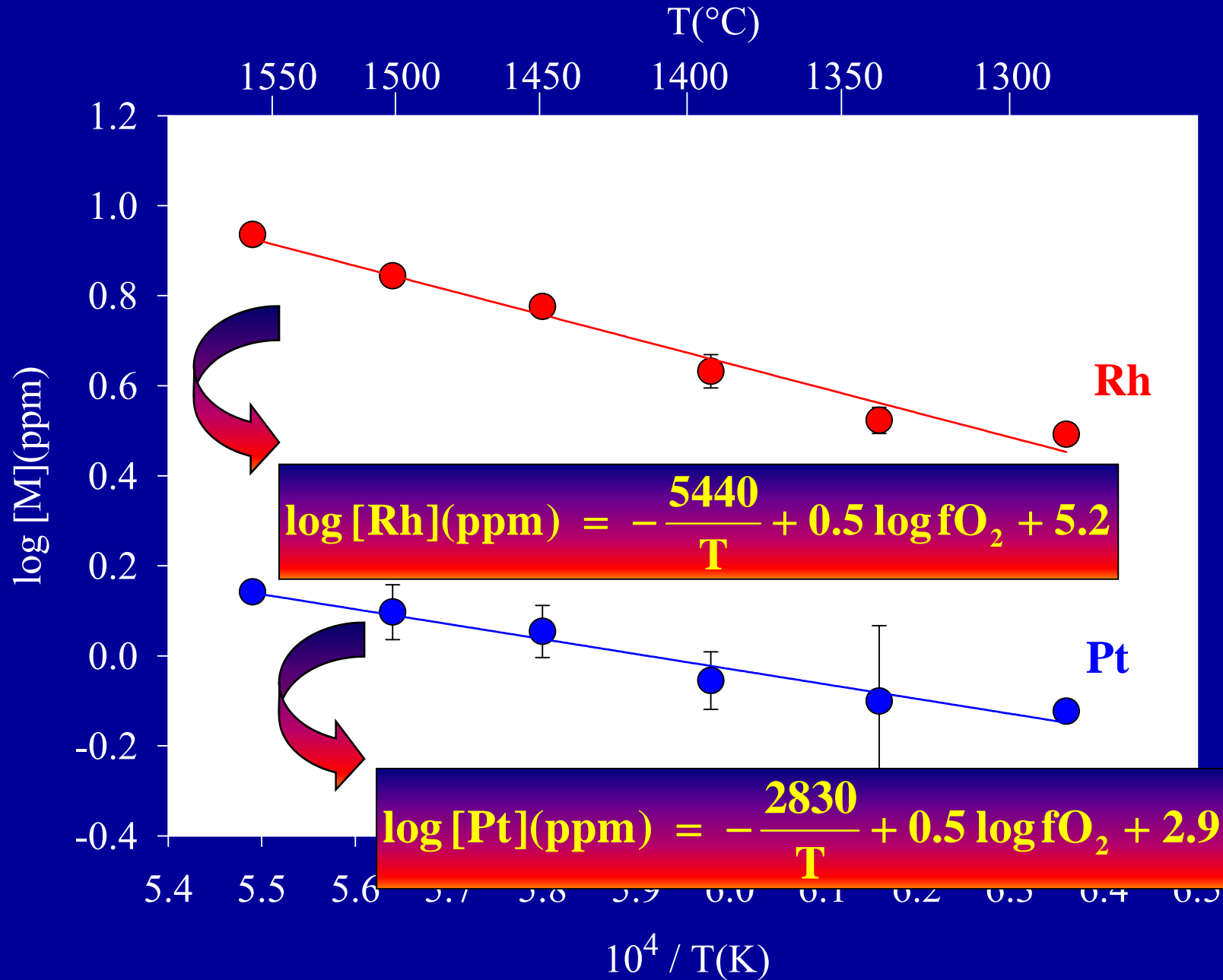


Magma ocean

Oxygen fugacity dependence of the solubility of Os at 1 bar



Temperature dependence of the solubilities of Pt and Rh at 1 bar



1-bar solubility results for Os, Pt and Rh

$$\log [\text{Os}](\text{ppb}) = 0.71 \log f\text{O}_2 + 7.52$$

$$\log [\text{Pt}](\text{ppm}) = -\frac{2830}{T(\text{K})} + 0.5 \times \log f\text{O}_2 + 2.9$$

$$\log [\text{Rh}](\text{ppm}) = -\frac{5440}{T} + 0.5 \times \log f\text{O}_2 + 5.2$$



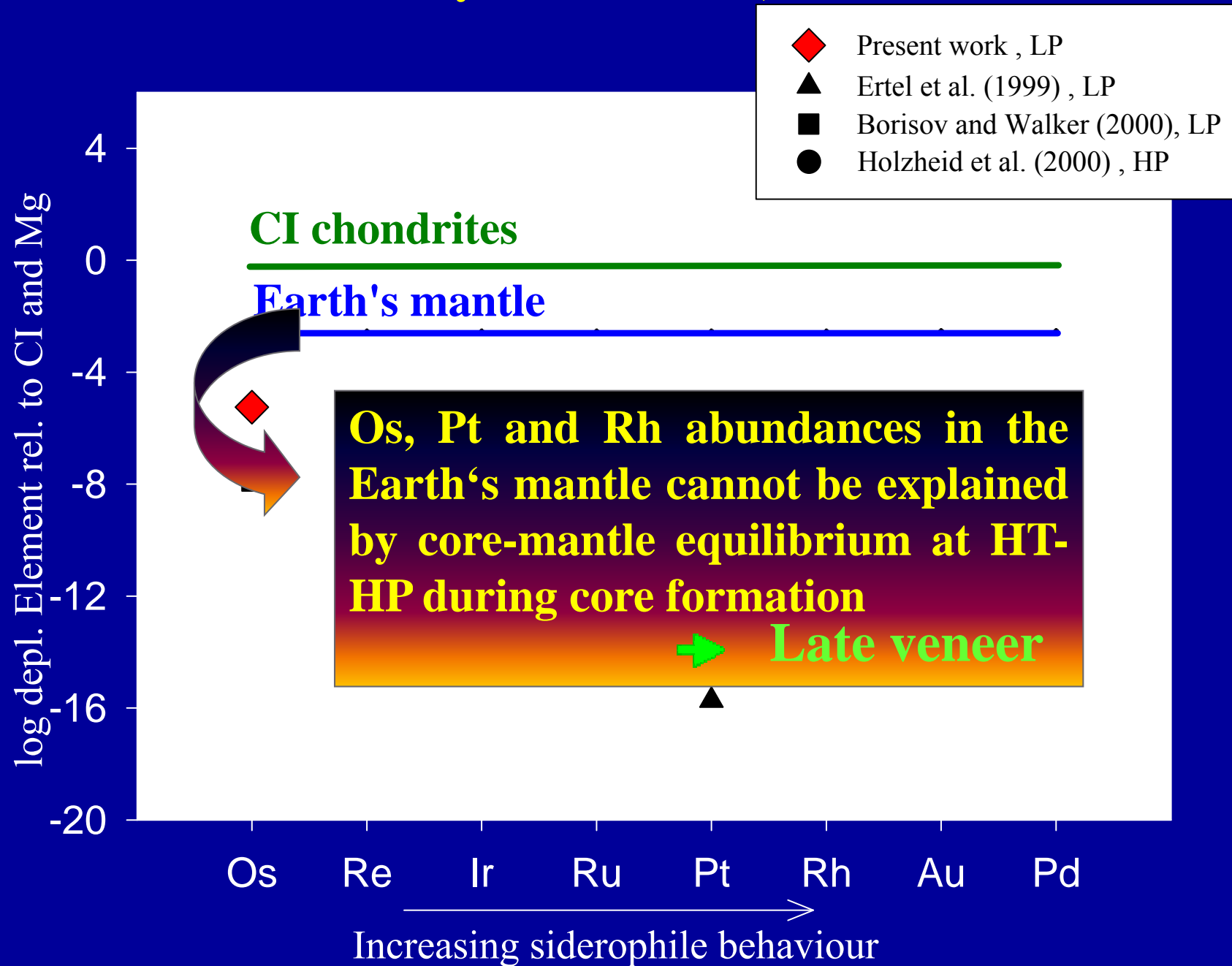
$$D_M^{\text{core/mantle}} = \frac{1}{A \times [\text{M}]_{T, f\text{O}_2} \times \gamma_M^{\text{Fe liq}, \infty}}$$

Solubility of M at T and fO₂ conditions relevant to core formation (3000K, 10^{-4.5} bars)

Thermodynamic data of the Fe-M system

$$[\text{M}]_{\text{mantle}} = \frac{[\text{M}]_{\text{CI}}}{(0.32D_M^{\text{core/mantle}} + 0.68)}$$

1-bar solubility results for Os, Pt and Rh

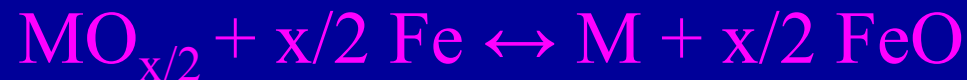


HP-HT liquid metal-magnesiowüstite partitioning of Re and Os

- During a HP-HT multi-anvil experiment, the oxygen fugacity is not directly measured.
- To interpret the data on the partitioning of an element M between liquid metal and oxide at high pressure and high temperature, it is common to use the two-component distribution coefficient KD_M :

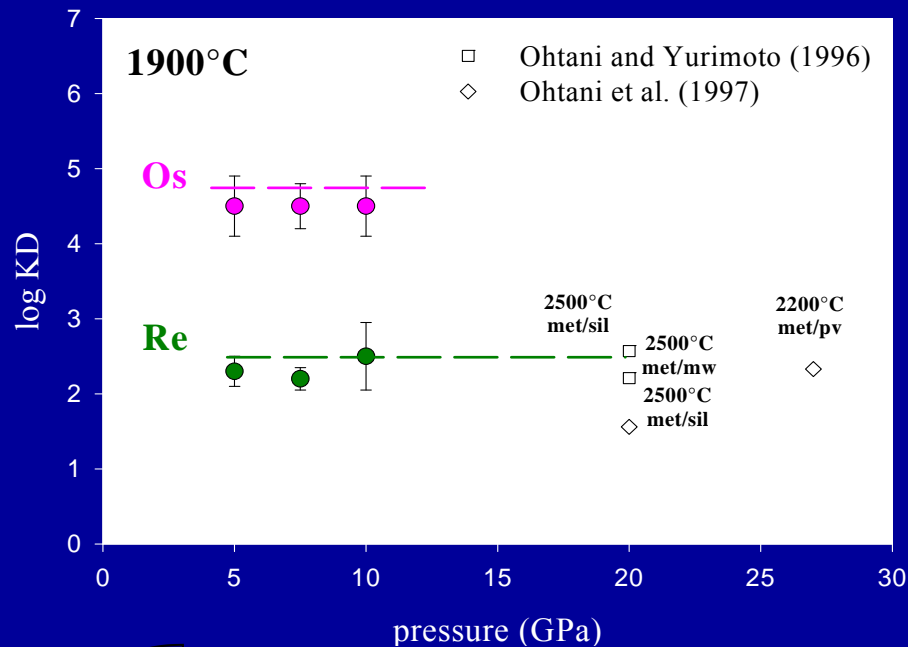
$$KD_M = \frac{D_M^{\text{met/ox}}}{\left(D_{\text{Fe}}^{\text{met/ox}}\right)^{x/2}} = \frac{[M]}{[MO_{x/2}]} \times \frac{[FeO]^{x/2}}{[Fe]^{x/2}}$$

which describes the distribution reaction



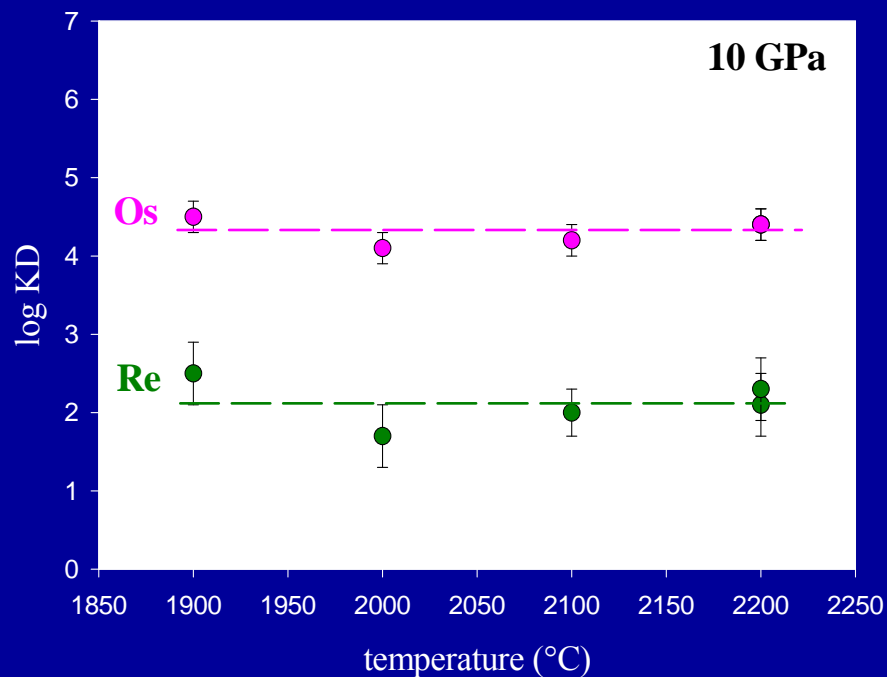
 KD_M is independent of fO_2

HP-HT liquid metal-magnesiowüstite partitioning of Re and Os



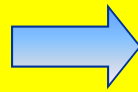
No temperature dependence

No pressure dependence



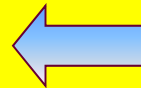
HP-HT liquid metal-magnesiowüstite partitioning of Re and Os

Experimental
results

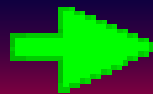


$$\begin{aligned} \log KD_{Os}^{\text{met/mw}} &\approx 4.1 - 4.5 \\ \log KD_{Re}^{\text{met/mw}} &\approx 1.8 - 2.5 \end{aligned}$$

$$\begin{aligned} \log KD_{Os}^{\text{met/mw}} &= 2 \\ \log KD_{Re}^{\text{met/mw}} &= 1.6 \end{aligned}$$





Values calculated
assuming core-mantle
equilibrium



~~core-mantle equilibrium~~

General Conclusions

-  MSE: based on the results from the 1 bar study on the compositional effect on the solubilities of Ni, Mo and W in melts, **the abundances of these MSE in the Earth's mantle could be the result of core-mantle equilibrium at HP-HT in a deep magma ocean of peridotitic composition.**
-  HSE: based, on the one hand, on the solubility results for Os, Pt and Rh obtained at 1 bar and on the HP/HT results on the partitioning of Re and Os between liquid metal and magnesiowüstite on the other hand, **equilibrium between core and mantle during core formation cannot explain the abundances of Re, Os, Pt and Rh in the Earth's mantle. The elevated and chondritic abundances of the highly siderophile elements observed in the Earth's mantle remain better explained by the addition of a late veneer of chondritic material after core segregation.**

Current research

Polysulfide as intermediate sulfur species are important for electron transfer processes in anoxic aquifers

1. Development of an analytical method for polysulfide identification and quantification
2. Determination of the importance of polysulfides as electron shuttles during microbial reduction of ferrihydrite by *Sulfurospirillum deleyianum*
3. Determination of the importance of polysulfides as electron donors and acceptors in different anoxic sulfur redox processes