



Science, the 5 Questions and Exploration

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Mineral Systems Work Flow

The Why Question

Why is the orebody there?

5 Questions

1. Geodynamics
2. Architecture
3. Fluid reservoirs
4. Flow drivers & pathways
5. Deposition

Inputs from:
Data Compilation
Data Collection
Modelling

The Where Question

Where is the next orebody?

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The 5 Questions

- 1. What are the geodynamic and P-T histories?**
- 2. What is the architecture of the system?**
- 3. What are the fluid reservoirs?**
- 4. What are the fluid flow drivers and pathways?**
- 5. What are the metal and sulphur transport and depositional processes?**

*Developed from inspiration provided by Tom Loutit
by (amongst others) John Walshe, Alison Ord,
Bruce Hobbs & Greg Hall in the AGCRC 1997-98*

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How do the 5Q help?

- **Mineral System focus**
- **Away from Source - Transport –Trap**
 - Ignores geodynamic and geological context, mixes fluid, metal (and sulphur) sources, implicitly links fluid flow drivers and pathways and finishes with erroneous concept of traps
- **Integrated framework for understanding ore deposits**
- **Concentrate on processes (not deposits)**
- **Provide the framework for a systematic approach to analysing prospectivity and exploration risk**



Talk Outline

- **Why do giant ore deposits form?**
 - Phenomenological approach
- **A set of 5 necessary, sufficient and independent factors**
- **Fluids in Mineral Systems**
- **The 5 factors**
- **Implications:**
 - Exploration risk analysis
 - Research and numerical simulation



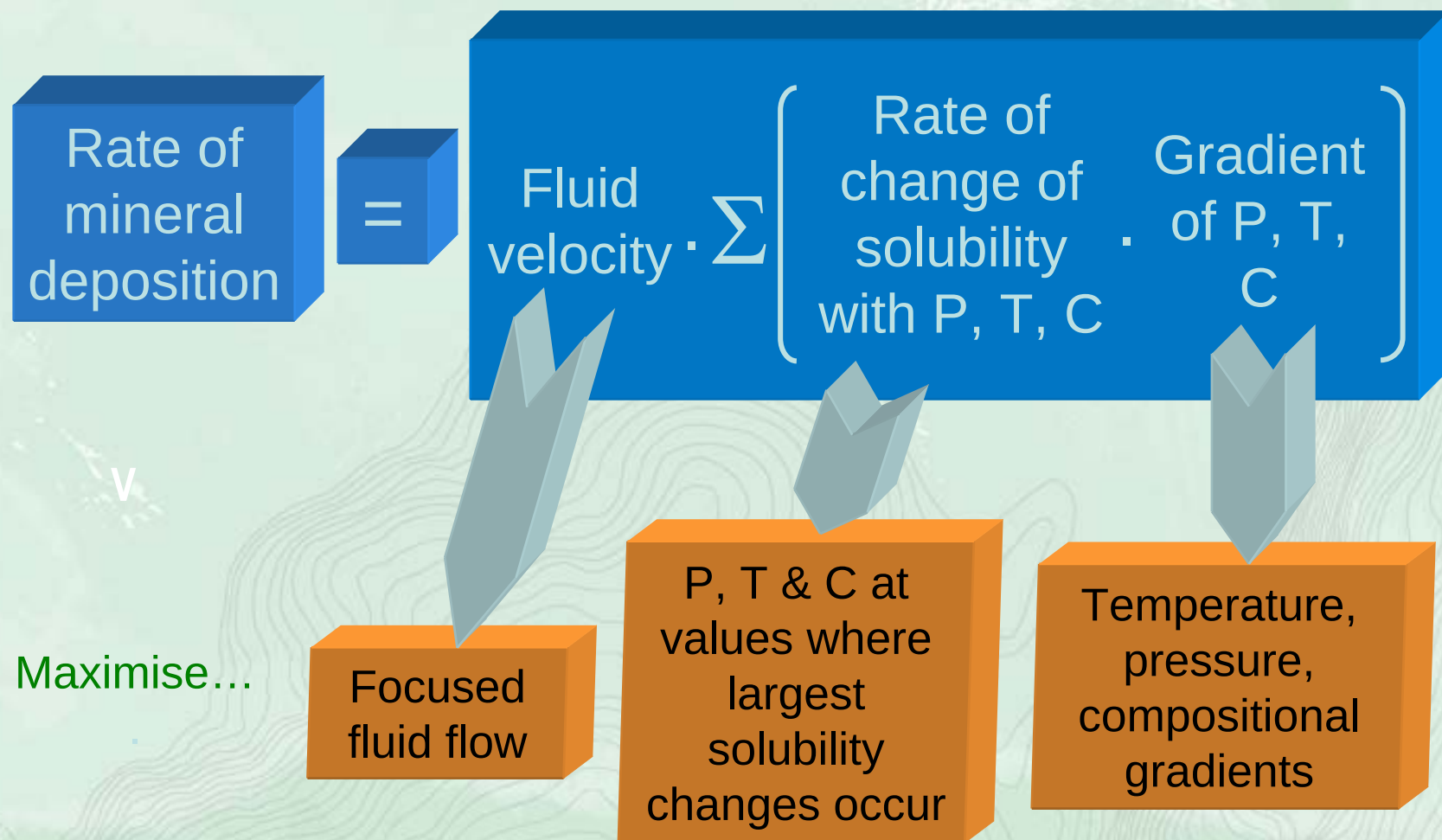
Why Do Giant Ore Deposits Form?

....because a lot of the appropriate mineral(s) precipitate

So what controls mineral precipitation?



Fundamental Relationship



After Phillips, 1990, 1991

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Two Additional Relationships

Fluid velocity = $\frac{k\rho g}{\mu}$ · Gradient in hydraulic potential

This diagram illustrates the relationship between fluid velocity and the gradient in hydraulic potential. It features a blue box on the left containing the text 'Fluid velocity', followed by a small blue box with an equals sign. To the right is a larger blue box containing the mathematical expression $\frac{k\rho g}{\mu}$ multiplied by 'Gradient in hydraulic potential'.

Total quantity of mineral deposited = $\int$ Rate of Mineral Deposition · dt

This diagram illustrates the relationship between the total quantity of mineral deposited and the rate of mineral deposition. It features a green box on the left containing the text 'Total quantity of mineral deposited', followed by a small green box with an equals sign. To the right is a larger green box containing the mathematical expression $\int$ multiplied by 'Rate of Mineral Deposition' and 'dt'.

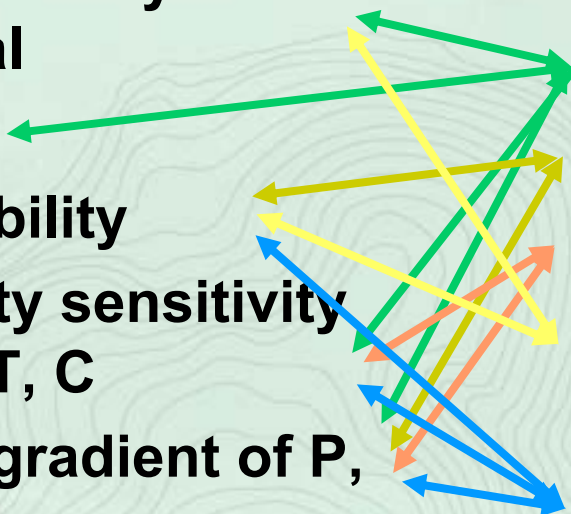
So.....

Quantity of Ore is a function of:

- A. Gradient in hydraulic potential
- B. Time
- C. Permeability
- D. Solubility sensitivity with P, T, C
- E. Spatial gradient of P, T, C

5 Questions

- Geodynamics
- Architecture
- Reservoirs
- Flow paths & drivers
- Deposition



The Largest Ore Deposits Form...

When and where

- **Gradient in hydraulic potential**
- **Permeability**
- **Solubility sensitivity to P, T, C**
- **Spatial gradient of P, T, C**

**have all been maximised
over prolonged period of
time**

Drivers; Geodynamics

Pathways; Architecture

Deposition;
Architecture;
Geodynamics

Architecture,
Drivers,
Deposition

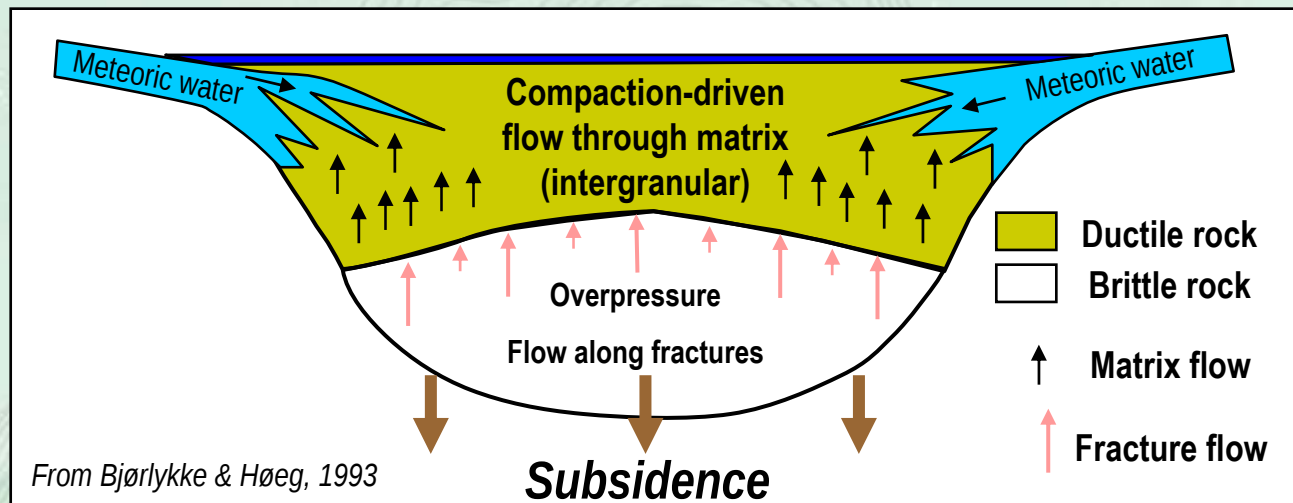
Geodynamics



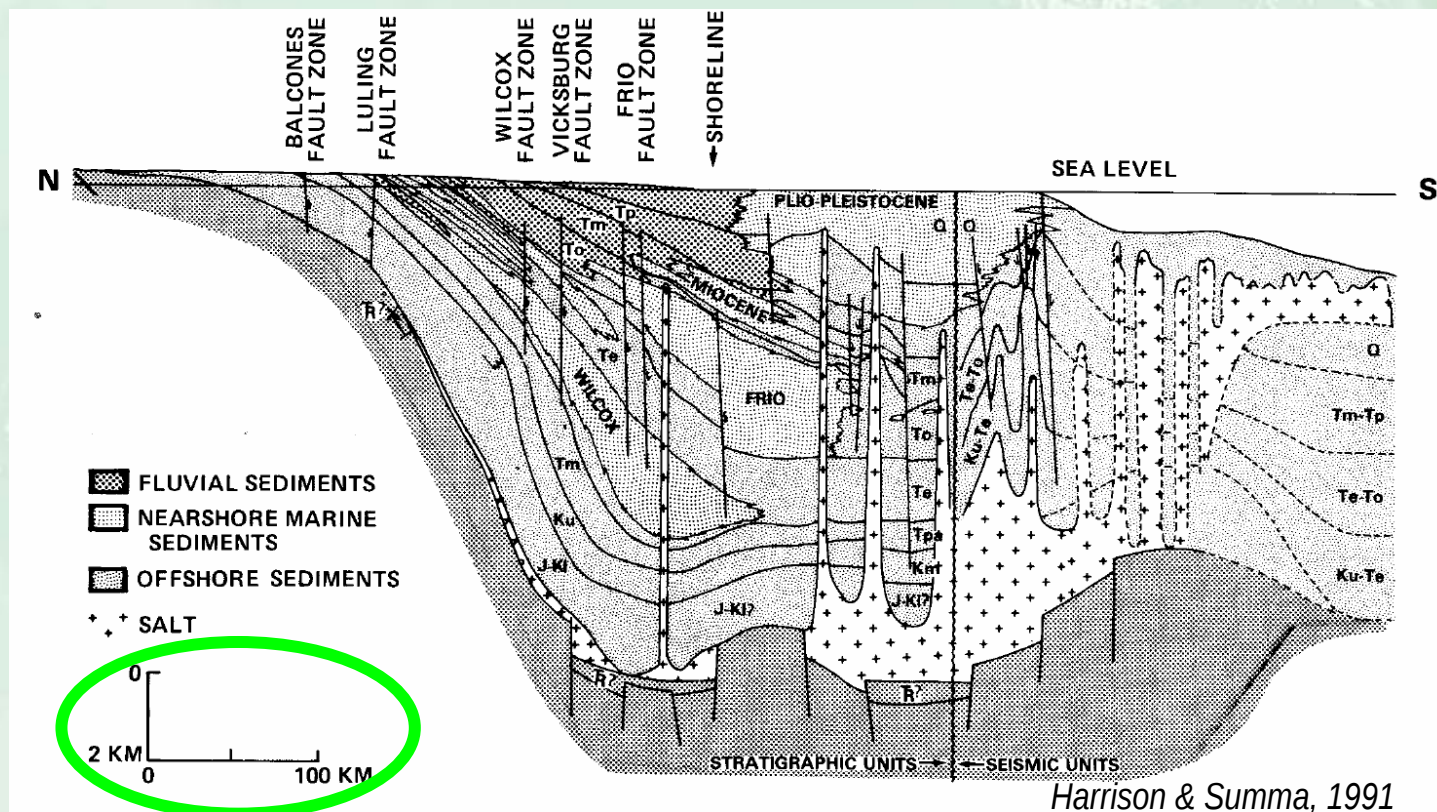
Fluid Sources: 1

Hydrosphere and Upper Crust:

- Meteoric, including bitterns
- Basinal, including compactional
- Fluids liberated during diagenesis



Fluids in Basins: Gulf of Mexico

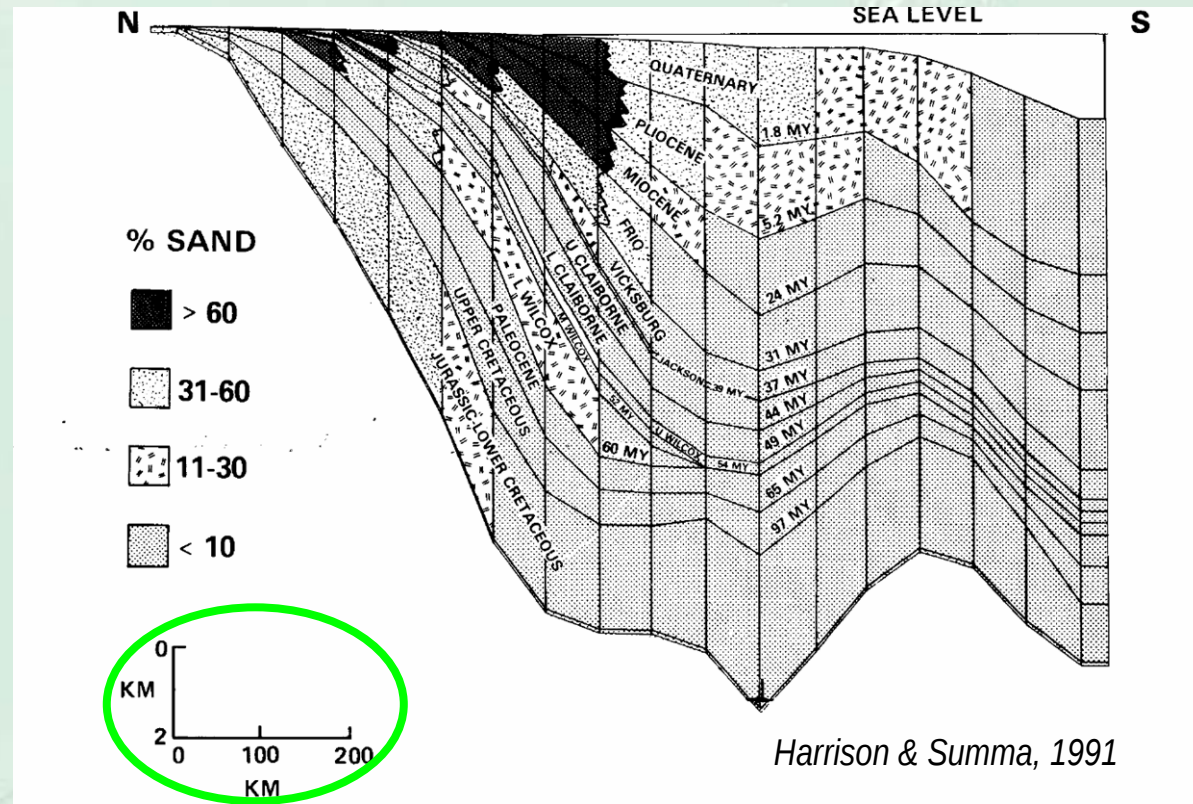


- Mesozoic-Recent sedimentation
- Up to 10 km thick – rapid sedimentation

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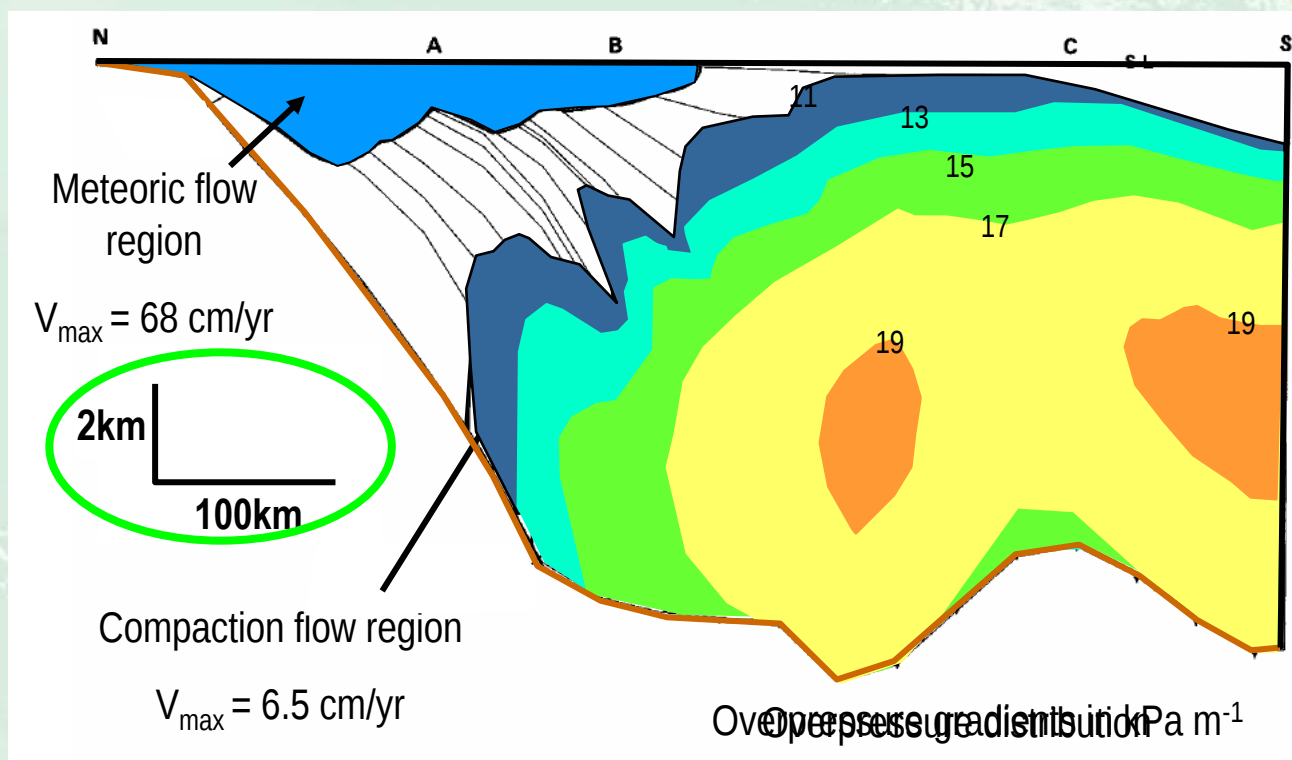
Fluids in Basins: Gulf of Mexico 2



- Modelled distribution of lithologies
- Based on drilling and seismic data

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Fluids in Basins: Gulf of Mexico 3



Harrison & Summa, 1991

- **Good match with present day observations**
- **Overpressures dynamic – not related to over seals**

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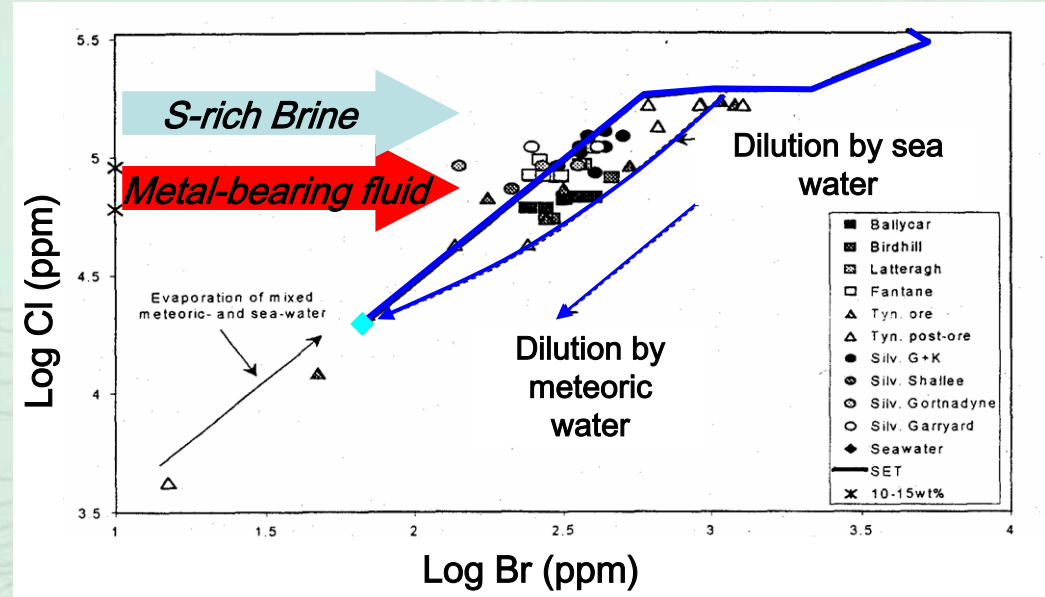


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Tracking Upper Crustal Fluids using Chemistry

- Irish deposits and prospects
- Two fluids derived from evaporated seawater present; one has reacted with basement



Gleeson et al., 1999

- **Complex patterns**
 - Most saline fluids represent sea water evaporated to beyond halite saturation at surface
 - Others represent diluted bitterns

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Fluid Sources: 2

Mid Crustal and deeper:

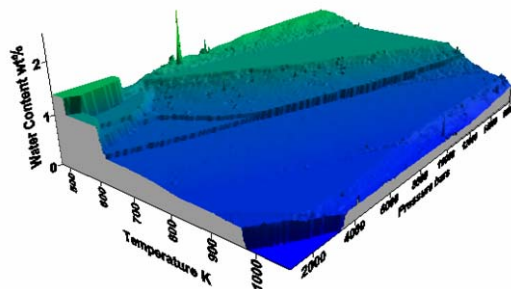
- **Metamorphism**
- **Magmatism**
- **Mantle and other sub-crustal sources**
- ***Timing and fluid budgets***

Granite in the Karakoram Mountains

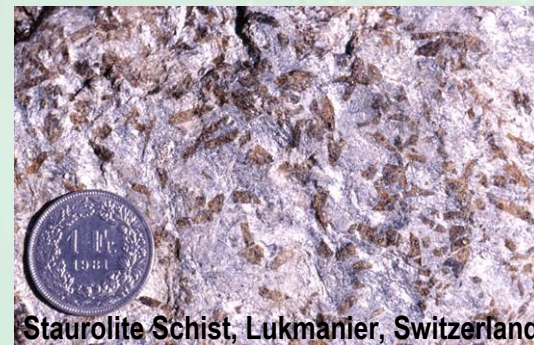


Metamorphic Fluxes

Granite maximum bound water content

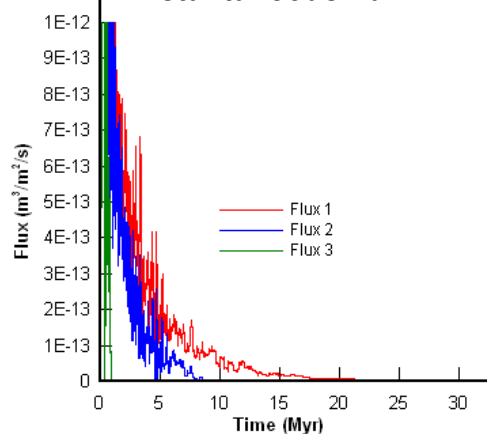


Garbenschiefer, Lukmanier, Switzerland

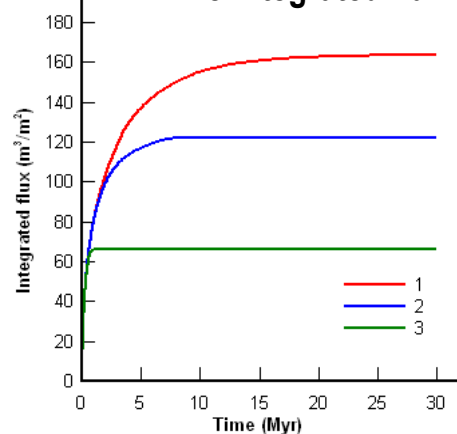


Staurolite Schist, Lukmanier, Switzerland

Instantaneous flux



Time-integrated flux



- Heat granitic crust so base at 40km reaches 1200°C
- Monitor flux at different depths in homogeneous crust
- Fluid flux decreases with time as steady state re-established and water lost

— Modelling by Heather Sheldon

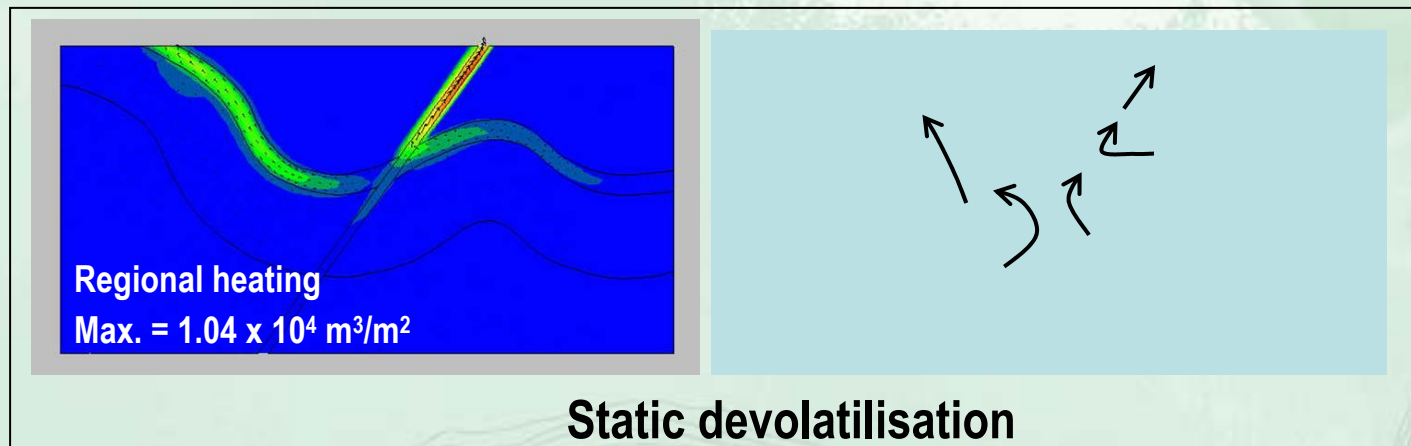
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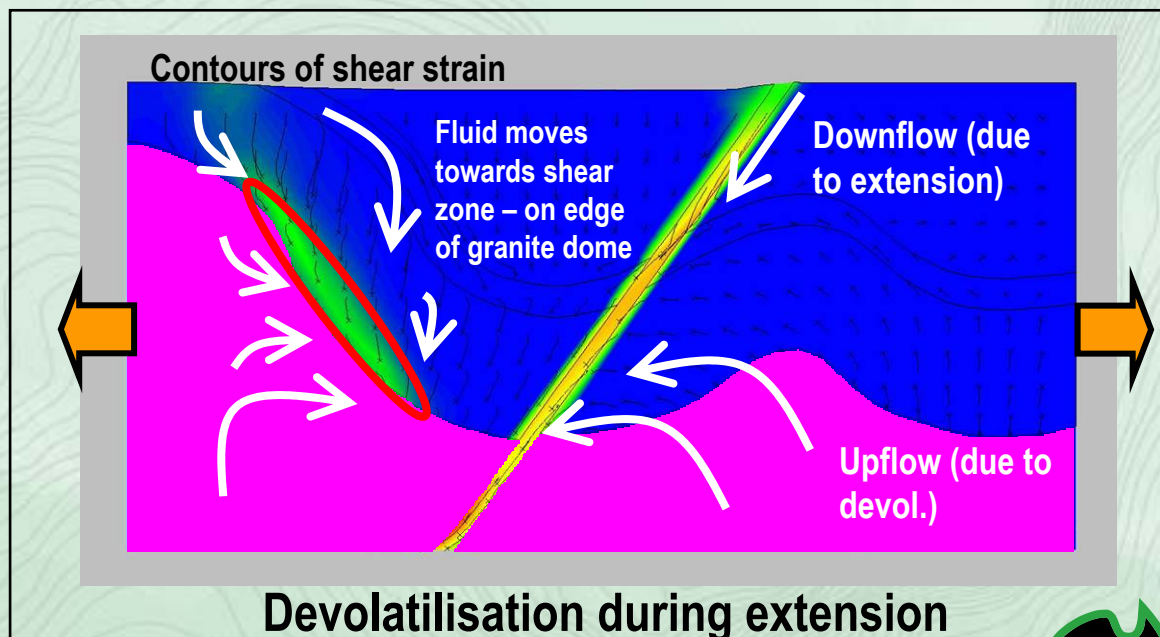
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Metamorphic Fluxes



- Upper crustal devolatilisation fluxes focussed – possibly enough
- Extension leads to downflow but facilitates mixing



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Modelling of Heather Sheldon & Yanhua Zhang

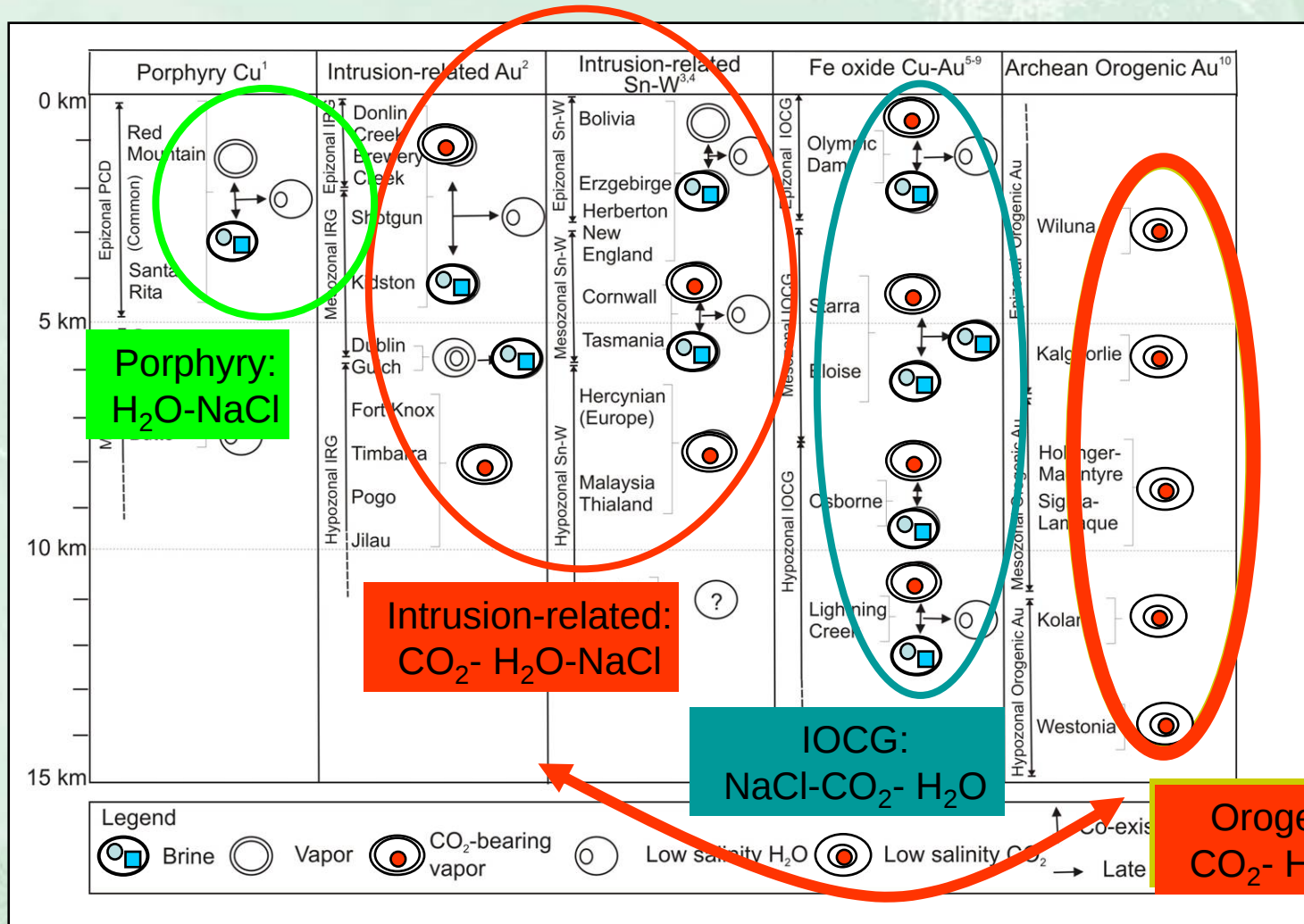


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pmd*CR

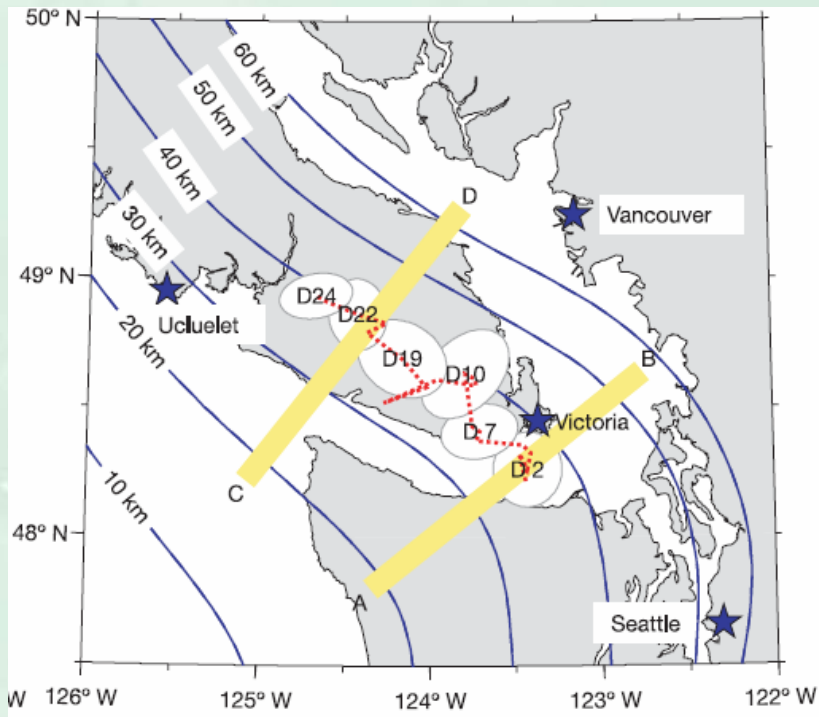
Crustal Fluid Sources



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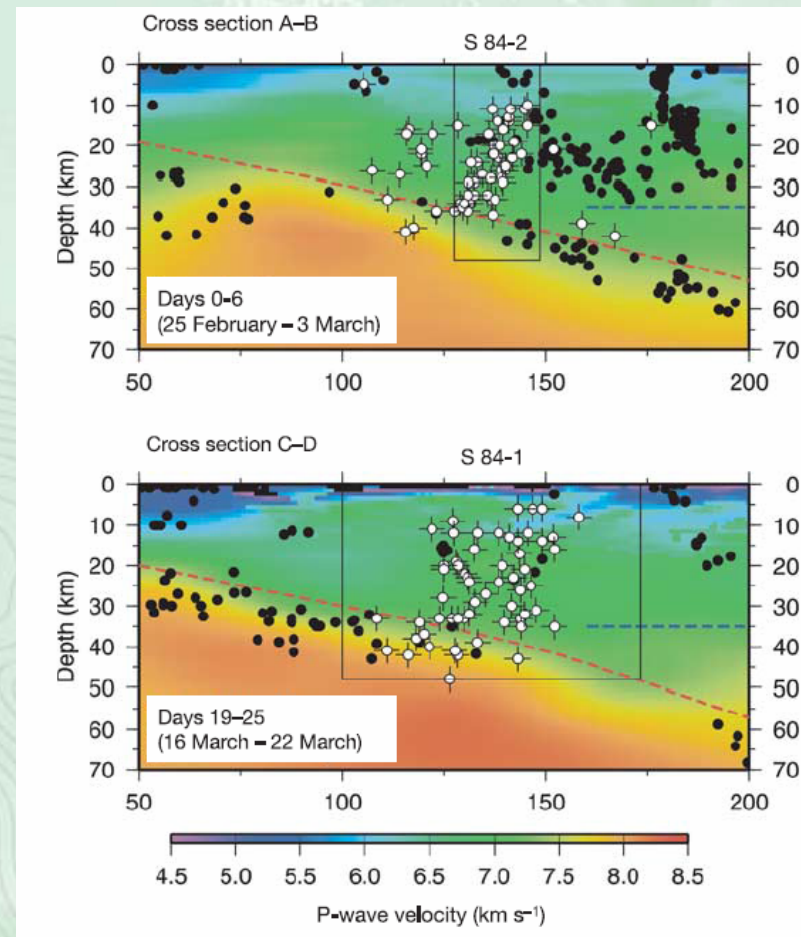
Stolen from a Tim Baker paper

Sub-Crustal Fluid Reservoirs



- **Cascadia Margin**
- **Tremor swarms indicative of fluid movement**
- **Fluid sourced from dewatering slab?**

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Kao et al, 2005

Fluid Sources: 3



- **Fluids are only available at certain times and places in the geological evolution of an area**
- **‘Surface’ fluids (basinal; diagenetic) transient unless a pathway is created to sealed compartments or the surface**
- **Fluxes?**



A. Creating Hydraulic Potential Gradients

- **Topographic relief**
- **Fluid pressure changes induced by deformation**
 - **Compaction is a subset of this**
- **Convection induced by thermal or chemical buoyancy**
- **Fluid pressure changes generated by chemical reactions**
 - **Diagenesis, hydrocarbon generation, metamorphism**
- **Pressure gradients generated by fluids released from magmas**

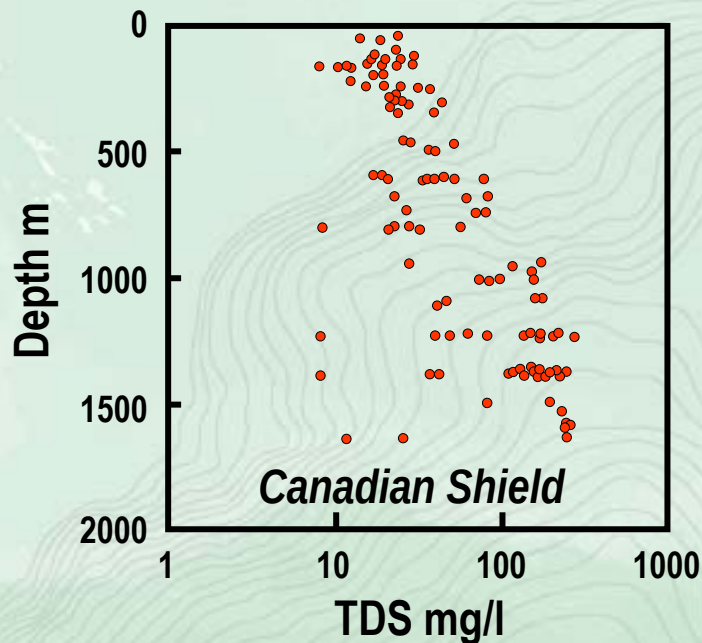
After Bruce Hobbs

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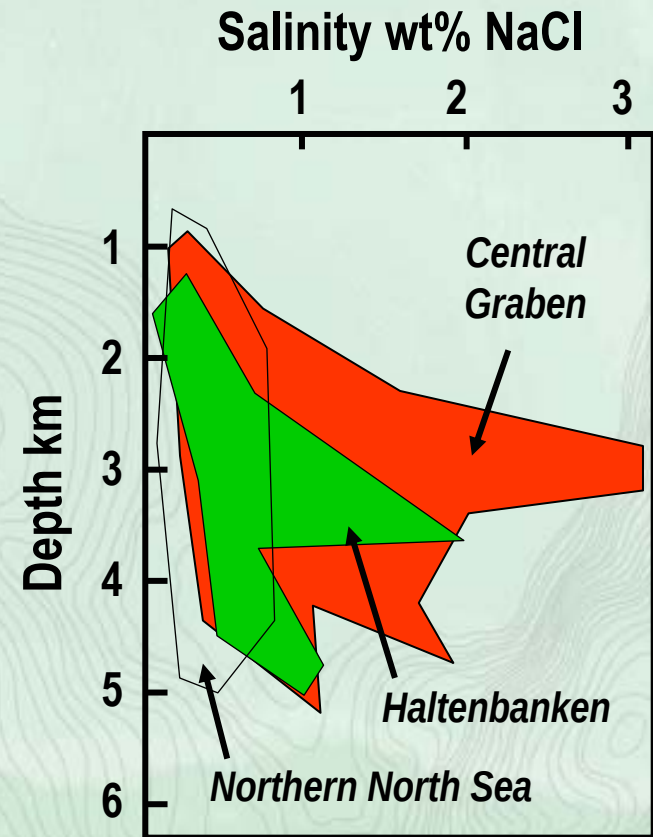
Upper Crustal Fluid Migration

- Salinity distributions imply a lack of convection

– And real difficulty in establishing it



Deming, 1994



Bjorlykke & Gran, 1994

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B, C. Pathways

Permeability maximised *over time*
where:

- **deep structures can be repeatedly reactivated**
- **structures intersect**
- **damage zones develop**
- **mineral dissolution occurs**



C. Permeability Generation and Destruction

- **Competition between**

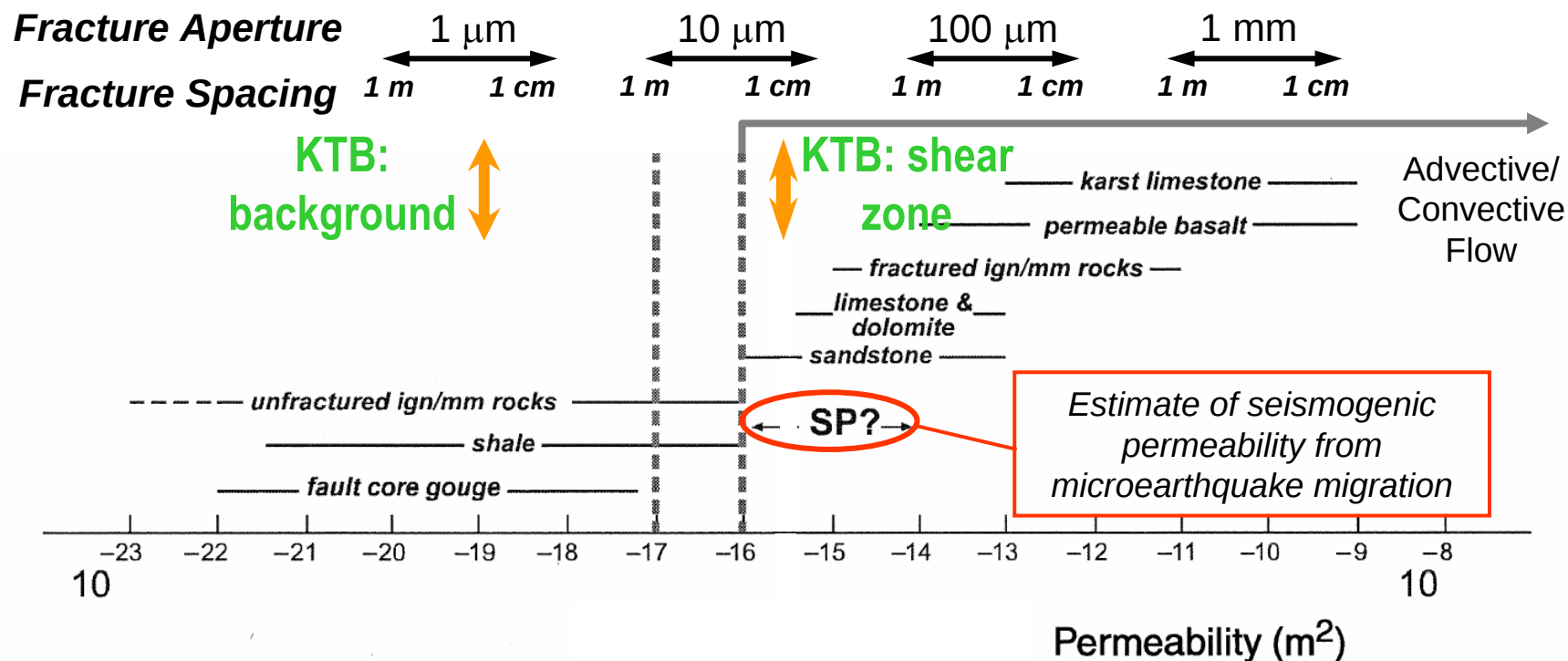
- 1. Permeability Creation**

- **Deformation**
 - Extensional and compressional
- **Increase in porosity due to reaction**

- 2. Permeability Destruction**

- **Compaction (not just in basins)**
- **Decrease in porosity due to mineral precipitation**

Permeability Values

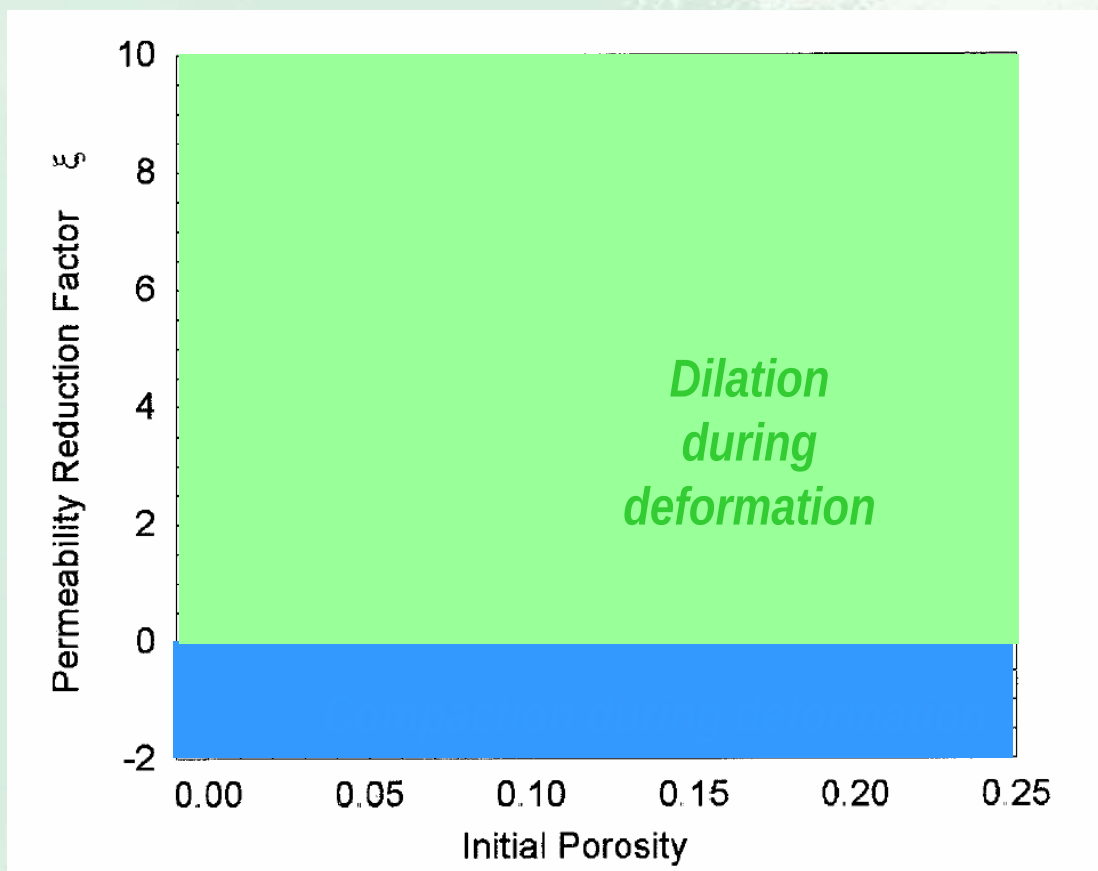


Based on Rowland & Sibson, 2004

Data from McDermott et al., 2006

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Fault Permeability and Rock Type

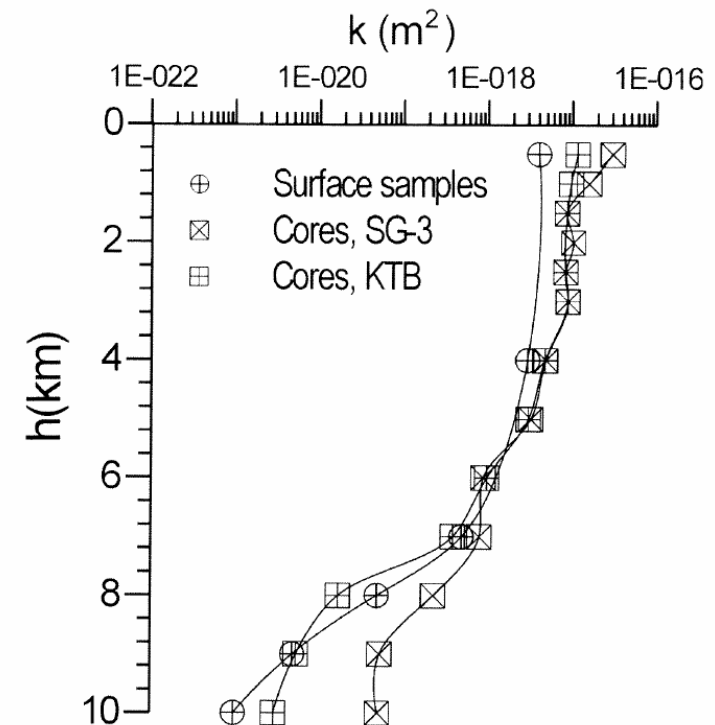
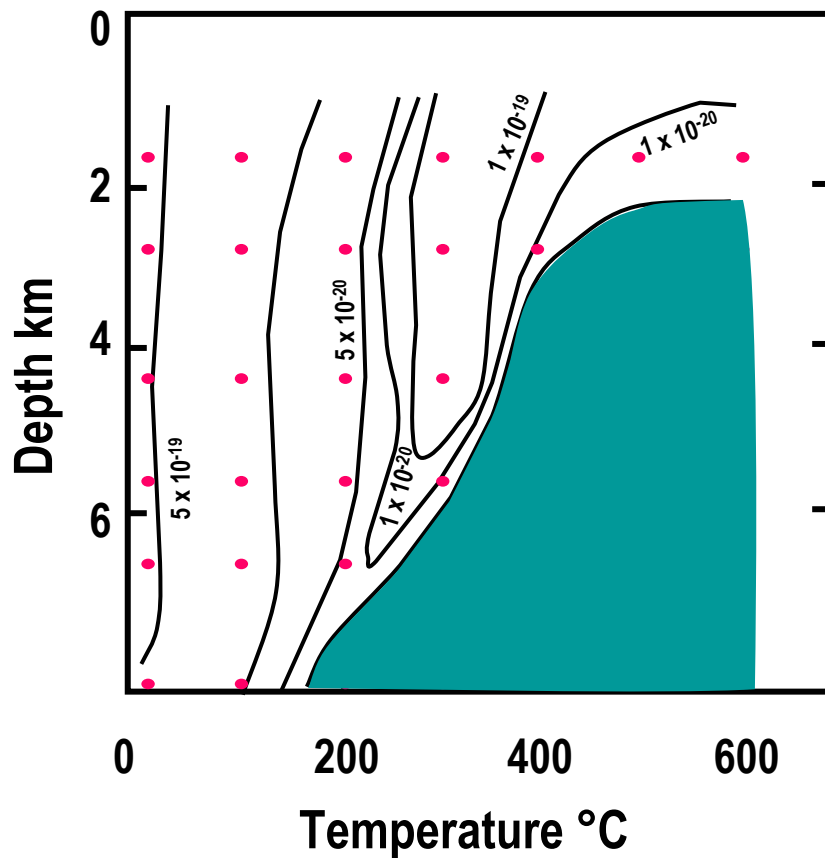


- **Ductile regime – variation in permeability with lithology**
- **Brittle regime – variation in permeability with porosity**
- **When do fault permeabilities depend on host lithologies?**

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Permeability Variation

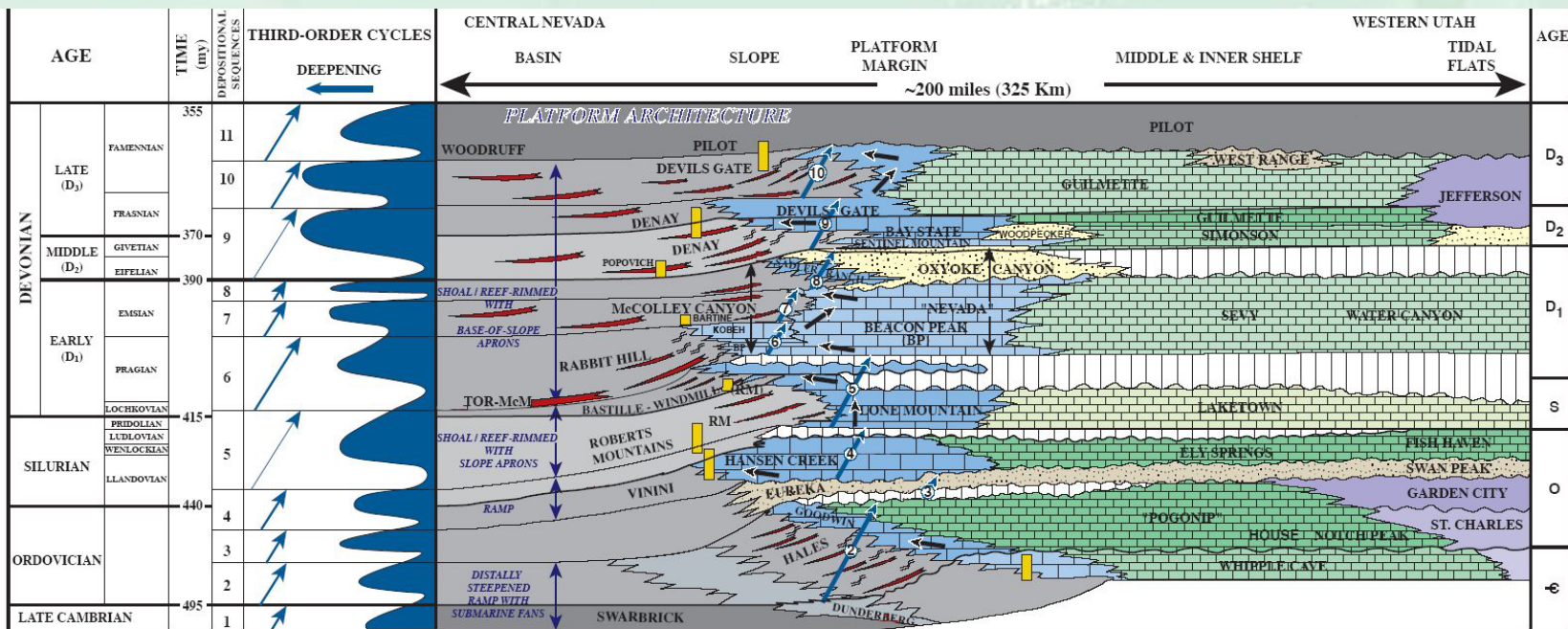


Zharikov et al., 2005

- **Permeability of plugs from Kola borehole**
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Great Basin USA: Lower Palaeozoic Architecture



STRATIGRAPHIC OCCURRENCES OF GOLD



SLOPE AND BASIN



TURBIDITE/DEBRIS FLOWS



PLATFORM MARGIN



MIDDLE AND INNER SHELF



TIDAL FLATS



SILICICLASTIC SANDSTONE

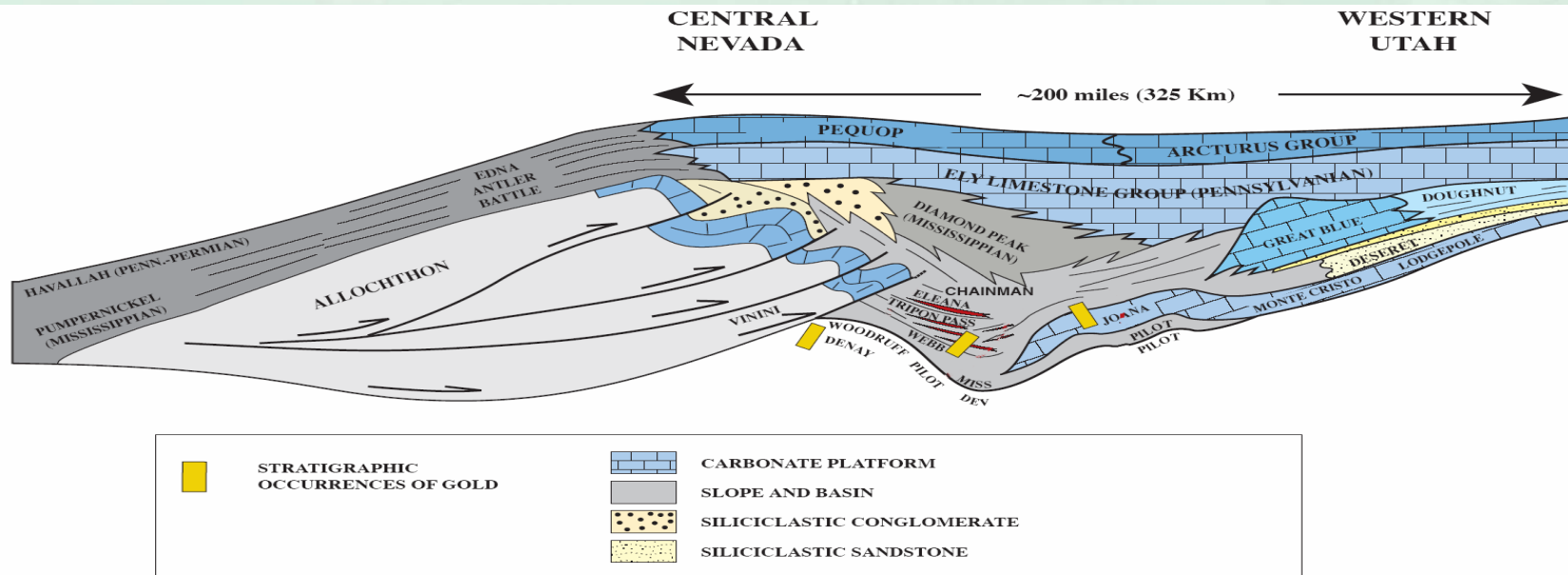
Cook & Corboy, 2004

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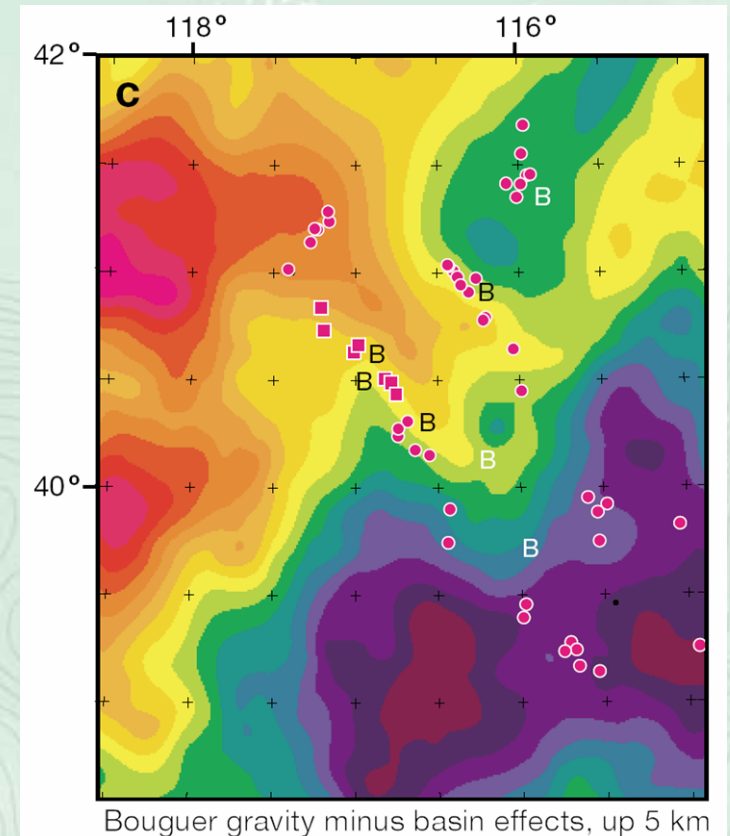
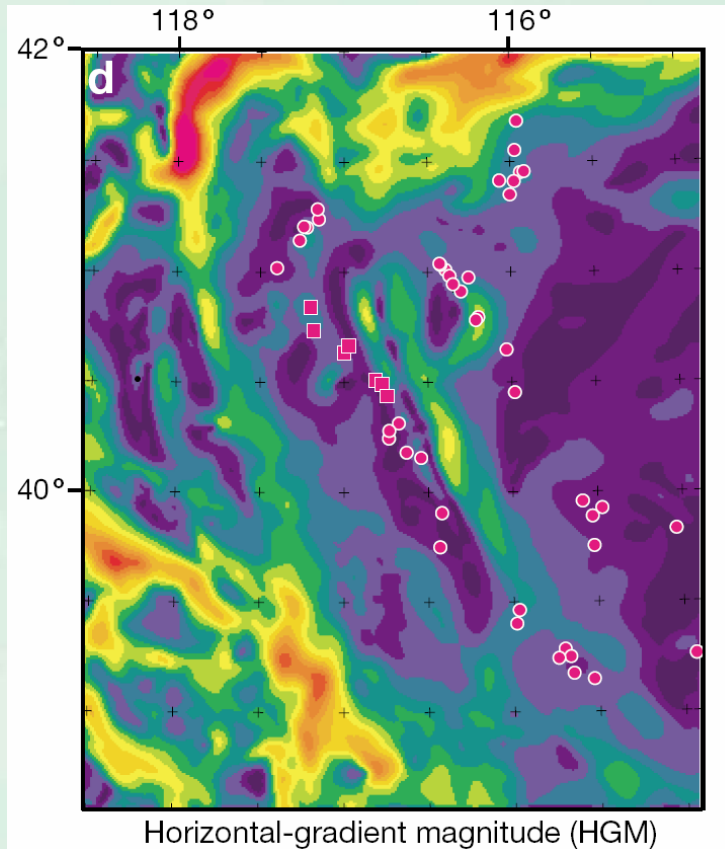
Great Basin: Controls on Architecture



Deep Structures
seen in Geophysics
as well as isotope
data etc.

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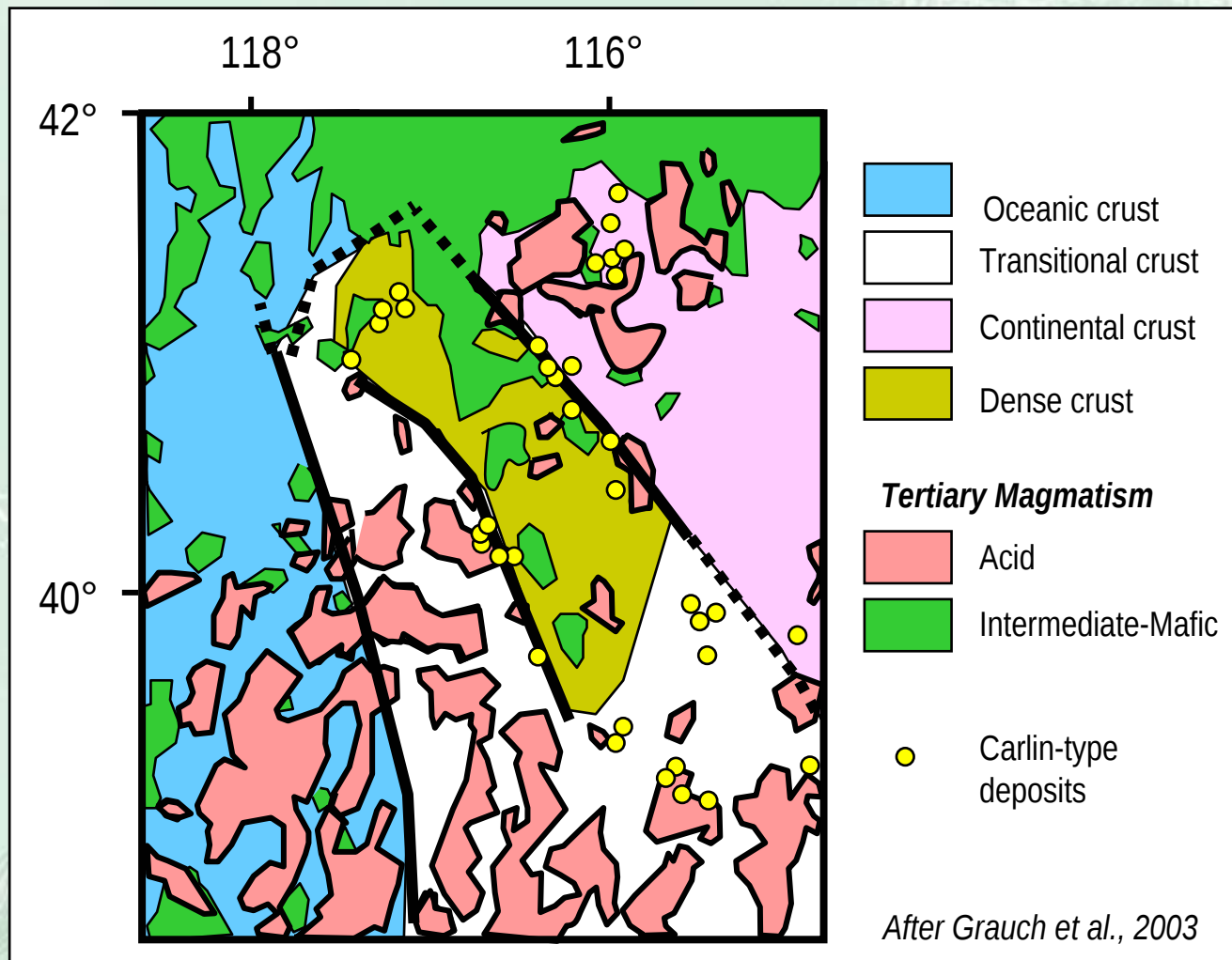
Great Basin: Architecture



From Grauch et al., 2003

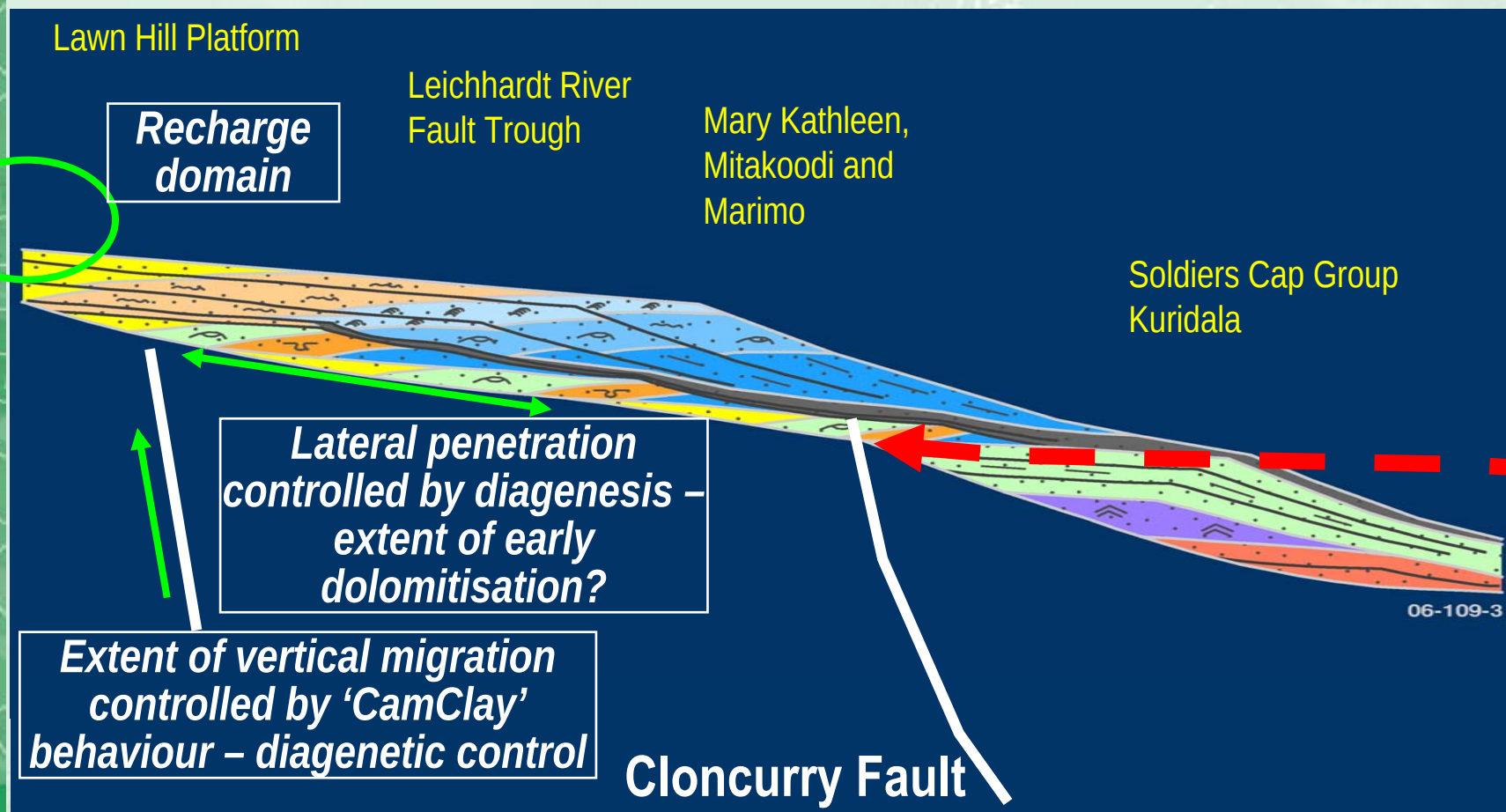
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Great Basin: Geodynamics



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Mt Isa Sediment Architecture 1690 – 1650 Ma



From Peter Southgate

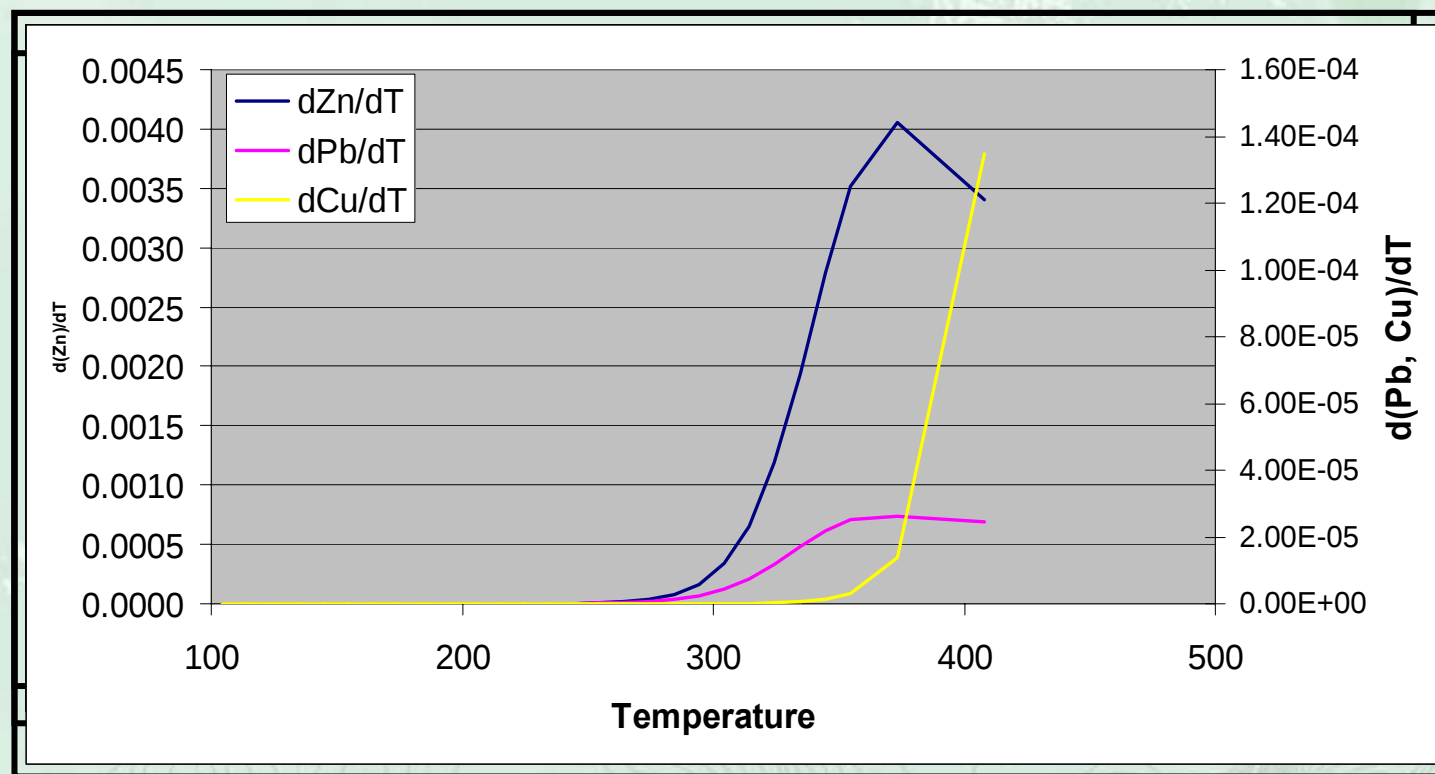
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D. Solubility Sensitivity - Temperature

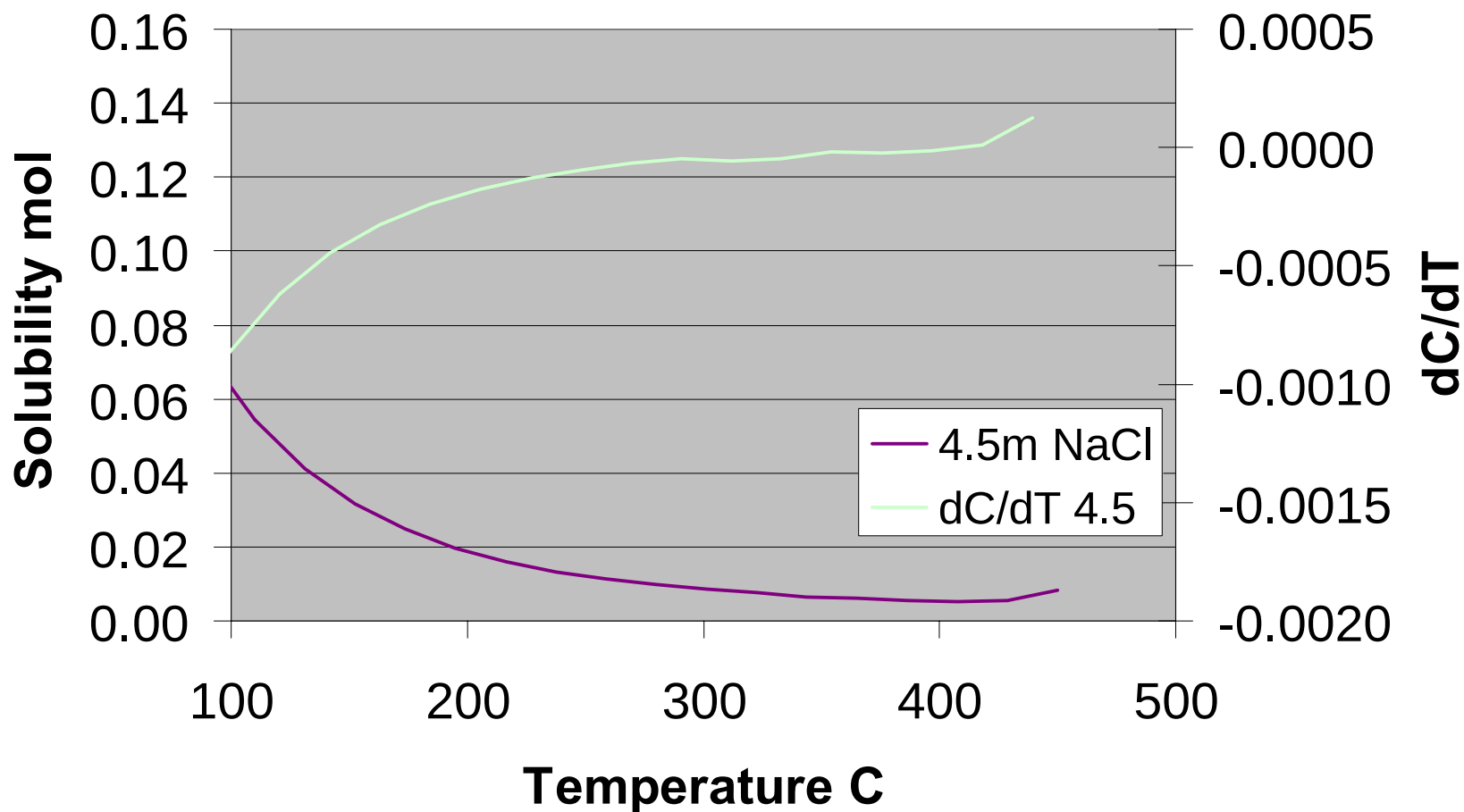


- Solubility of Zn (sphalerite), Pb (galena) and Cu (chalcopyrite) at 1 kbar
 - pH controlled by K-feldspar-muscovite-quartz
 - $\Sigma S = 0.01m$
 - Redox controlled by haematite-magnetite

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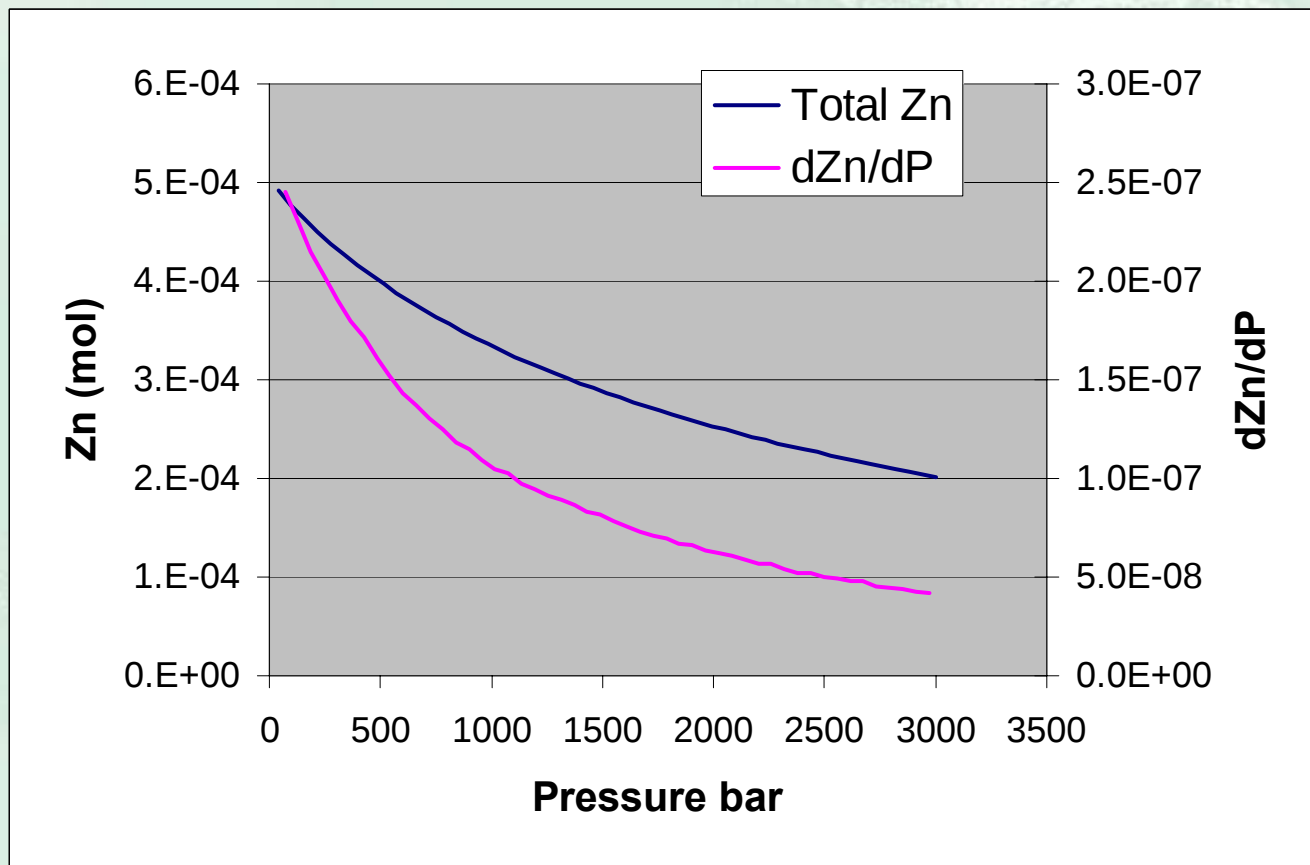
Solubility Sensitivity - Temperature



- **Ca in solution controlled by Anhydrite at 500 bars**
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Solubility Sensitivity - Pressure

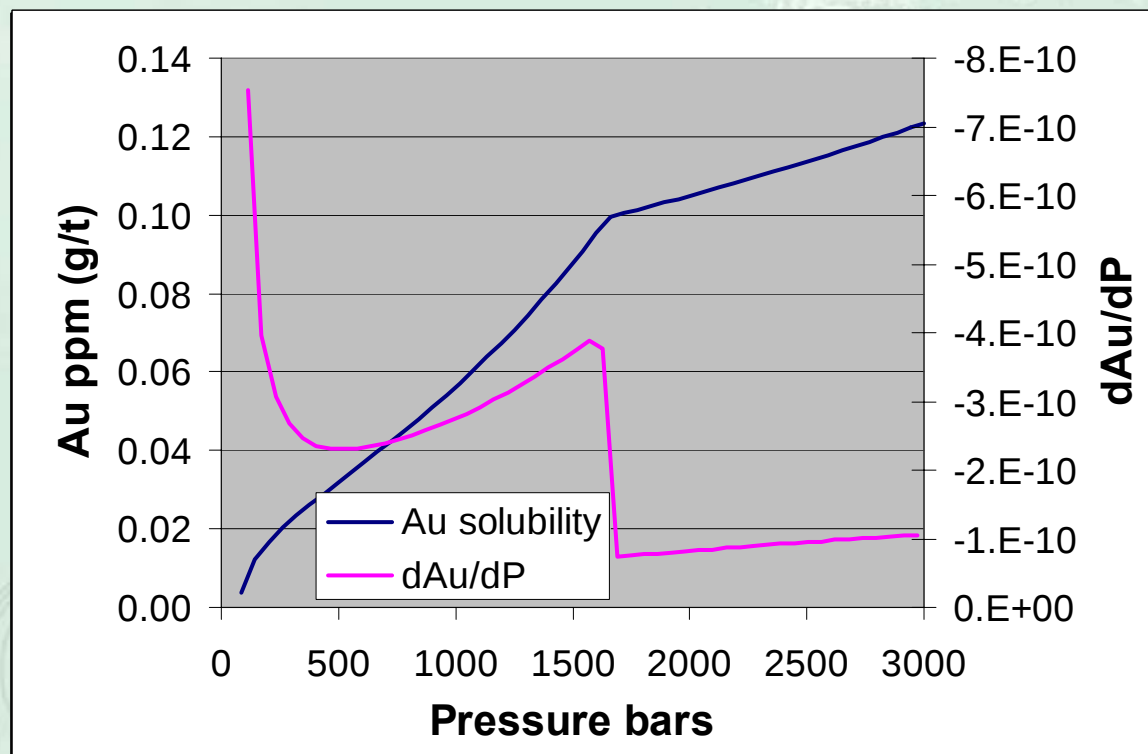


- **Solubility of Zn (sphalerite) at 250°C**
 - pH controlled by K-feldspar-muscovite-quartz
 - $\Sigma S = 0.01m$
 - Redox controlled by haematite-magnetite

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Solubility Sensitivity - Pressure

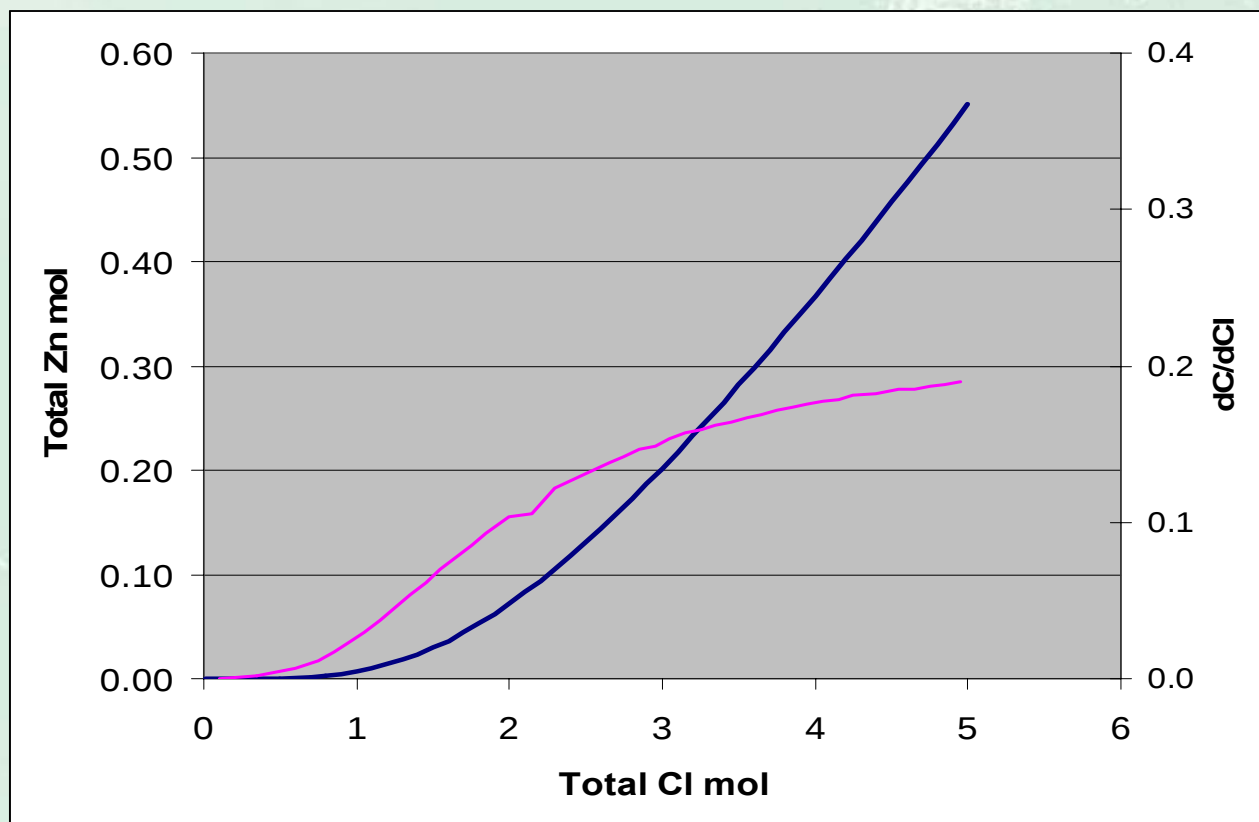


- **Solubility of Gold at 300°C**
 - pH controlled by K-feldspar-muscovite-quartz
 - SS = 0.01
 - Redox below Haematite-magnetite
 - Gas allowed to escape once formed

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Solubility Sensitivity - Chemistry



- **Solubility of Zn (Sphalerite) at 1kbar**
 - pH controlled by K-feldspar-muscovite-quartz
 - $\Sigma S = 0.01m$
 - Redox controlled by haematite-magnetite

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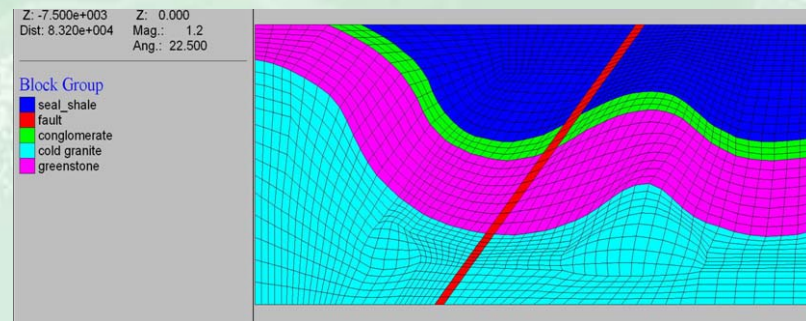
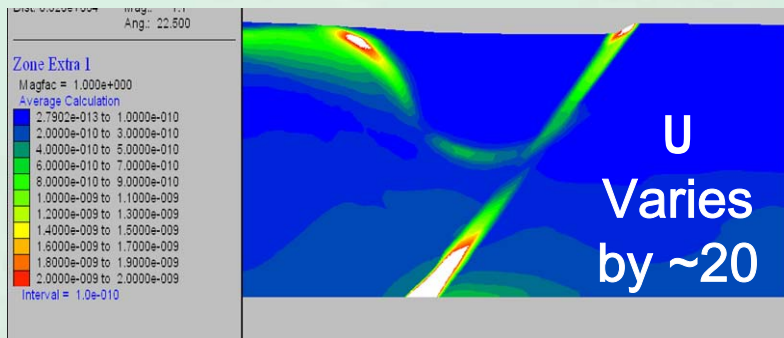


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P, T, C Gradients and Deposition

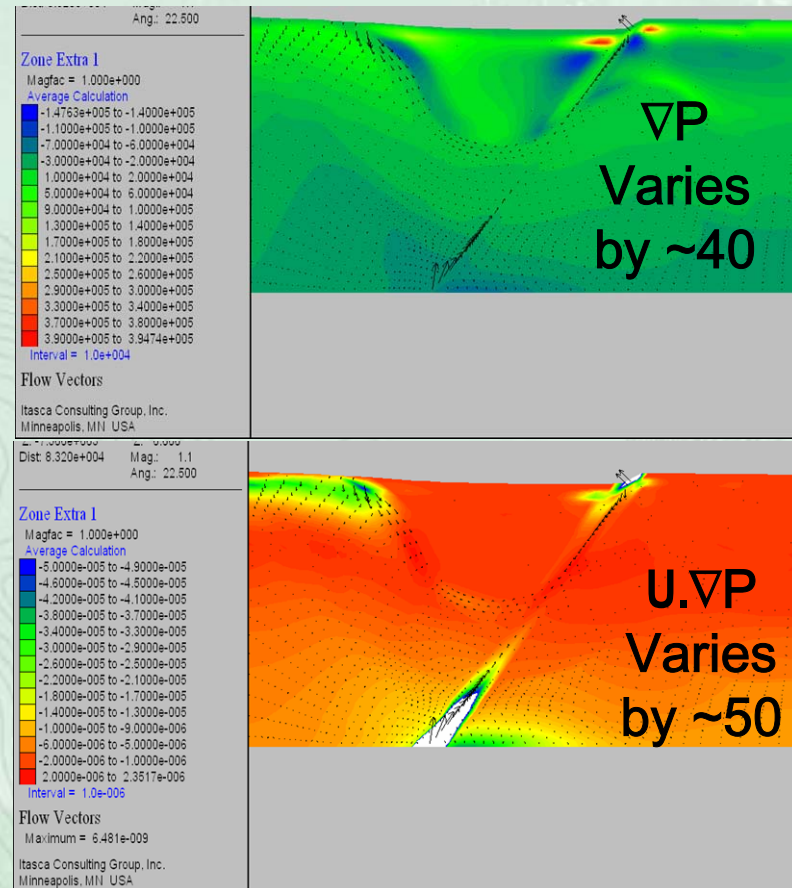
- **Temperature gradients**
 - Steepest near surface or around igneous centres
- **Pressure gradients**
 - Greatest (transiently) in dilation sites
- **Can induce unmixing**
- **Chemical gradients**
 - **Fluid-rock disequilibrium**
 - Due to both temperature and compositional effects
 - **Fluid mixing**
 - Efficacy demonstrated e.g. in Ireland

E. Spatial Gradients in T, P & C



- Yilgarn late basin models of Yanhua Zhang

— Plots generated specially by Heather Sheldon



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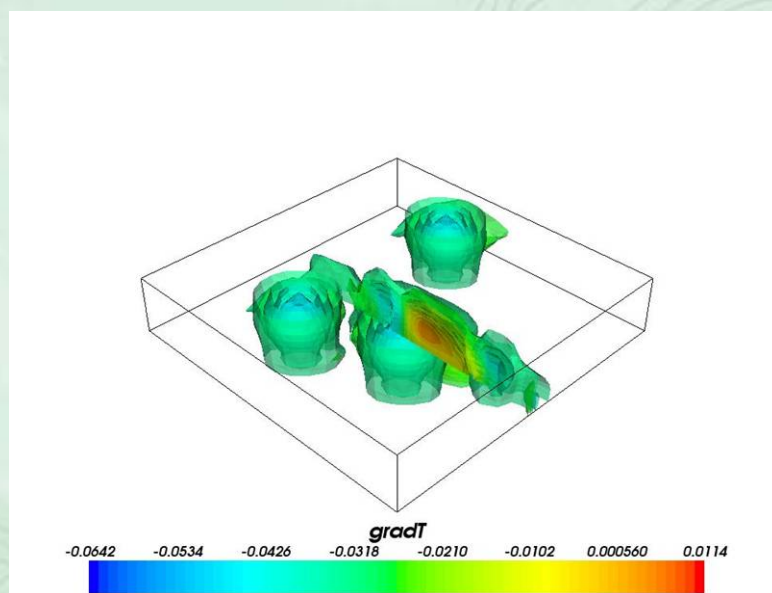
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Spatial Gradients in T , P & C

- Yilgarn granite convection models run by Julia Blockeel

— Plots generated specially by Heather Sheldon



- Velocity varies by a factor of $\sim 10^6$
- ∇P and ∇T change by ~ 10

	Spatial Gradient	Solubility Sensitivity	Spatial Sensitivity mol m ⁻¹
P (Zn)	$\sim 10^3$ Pa m ⁻¹	10^{-9} mol Pa ⁻¹	10^{-6}
P (Au)	$\sim 10^3$ Pa m ⁻¹	10^{-15} mol Pa ⁻¹	10^{-12}
T (Zn)	$\sim 10^{-2}$ K m ⁻¹	2×10^{-3} mol K ⁻¹	2×10^{-5}
T(Au)	$\sim 10^{-2}$ K m ⁻¹	3×10^{-9} mol K ⁻¹	3×10^{-11}

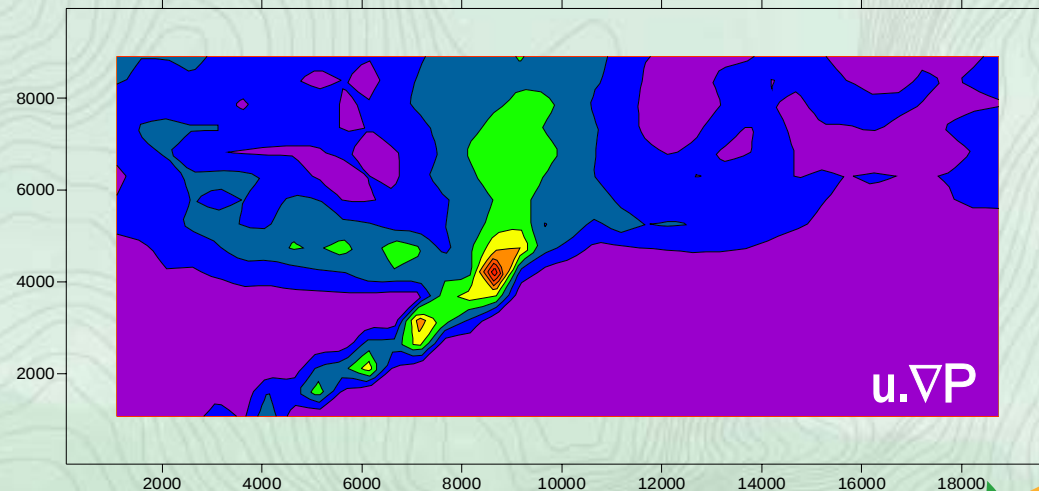
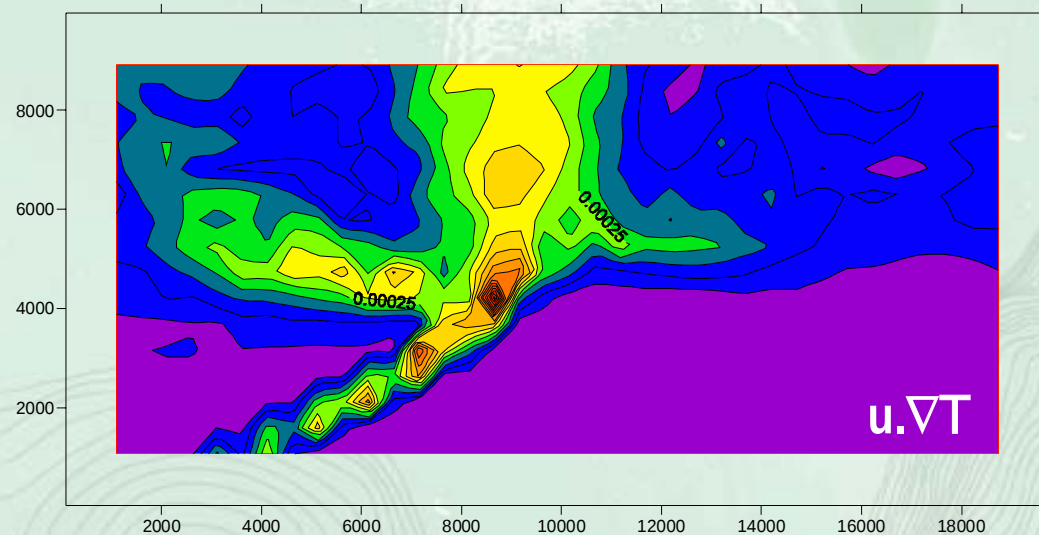
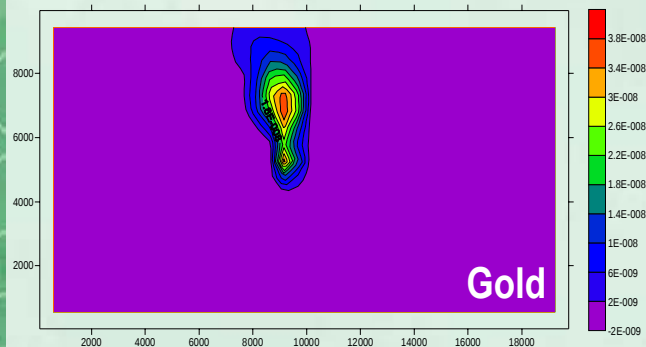
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Spatial Gradients in T , P & C



- Upwelling plume has elevated ∇P and ∇T and correlates with gold

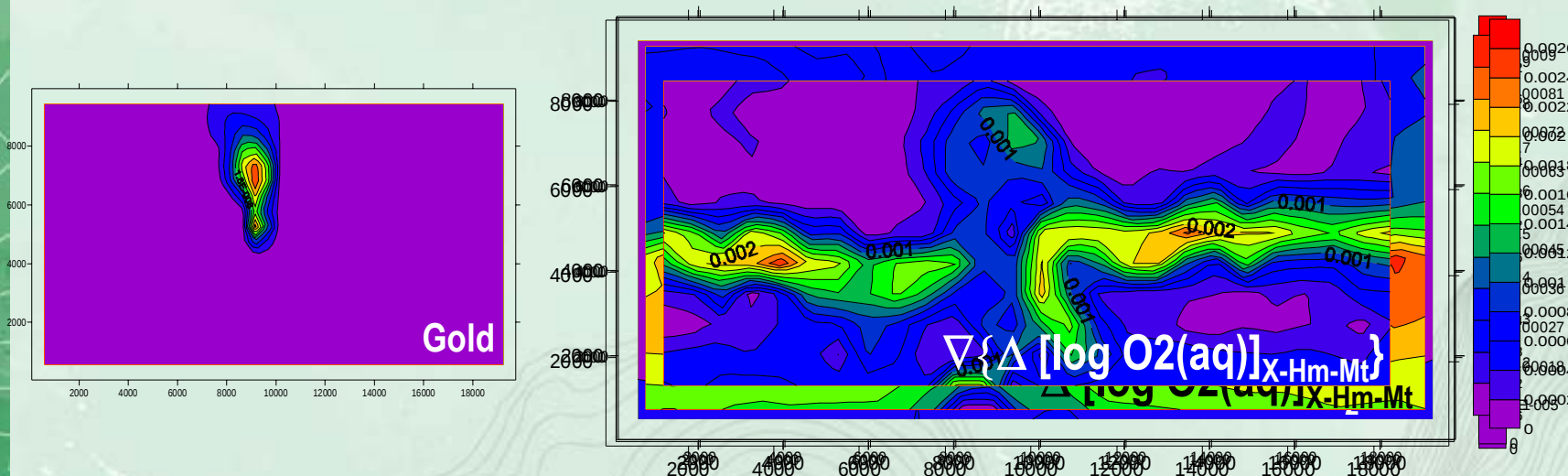
–Which is more important?

• Consider $\partial Au / \partial P$ and $\partial Au / \partial T$

–What about chemistry?



Spatial Gradients in T , P & C



- Gold solubility in model dominated by $AuHS_{(aq)}$:



Spatial Gradients in T, P & C

Parameter	Spatial Gradient	Solubility Sensitivity	Spatial Sensitivity mol m⁻¹
Pressure	0.008 bar m ⁻¹	7 × 10 ⁻¹² mol bar ⁻¹	5.6 × 10⁻¹⁴
Temperature	0.026 °C m ⁻¹	1 × 10 ⁻⁹ mol °C ⁻¹	2.6 × 10⁻¹¹
H₂S	2.5 × 10 ⁻⁶ mol m ⁻¹	2 × 10 ⁻⁴ mol mol ⁻¹	5 × 10⁻¹⁰
O₂	3 × 10 ⁻³⁵ mol m ⁻¹	1 × 10 ²⁷ mol mol ⁻¹	3 × 10⁻⁸



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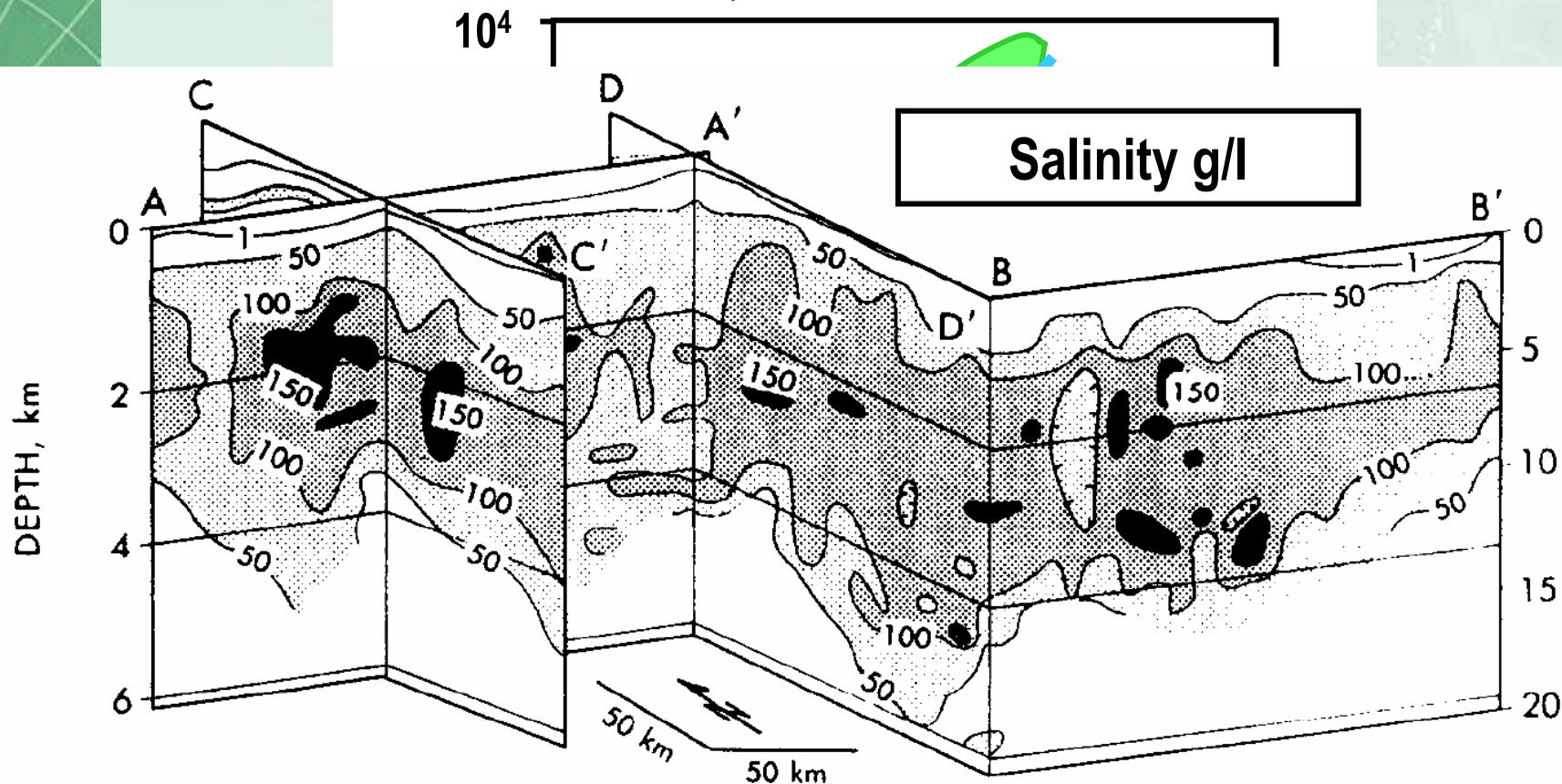


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Real World Spatial Gradients

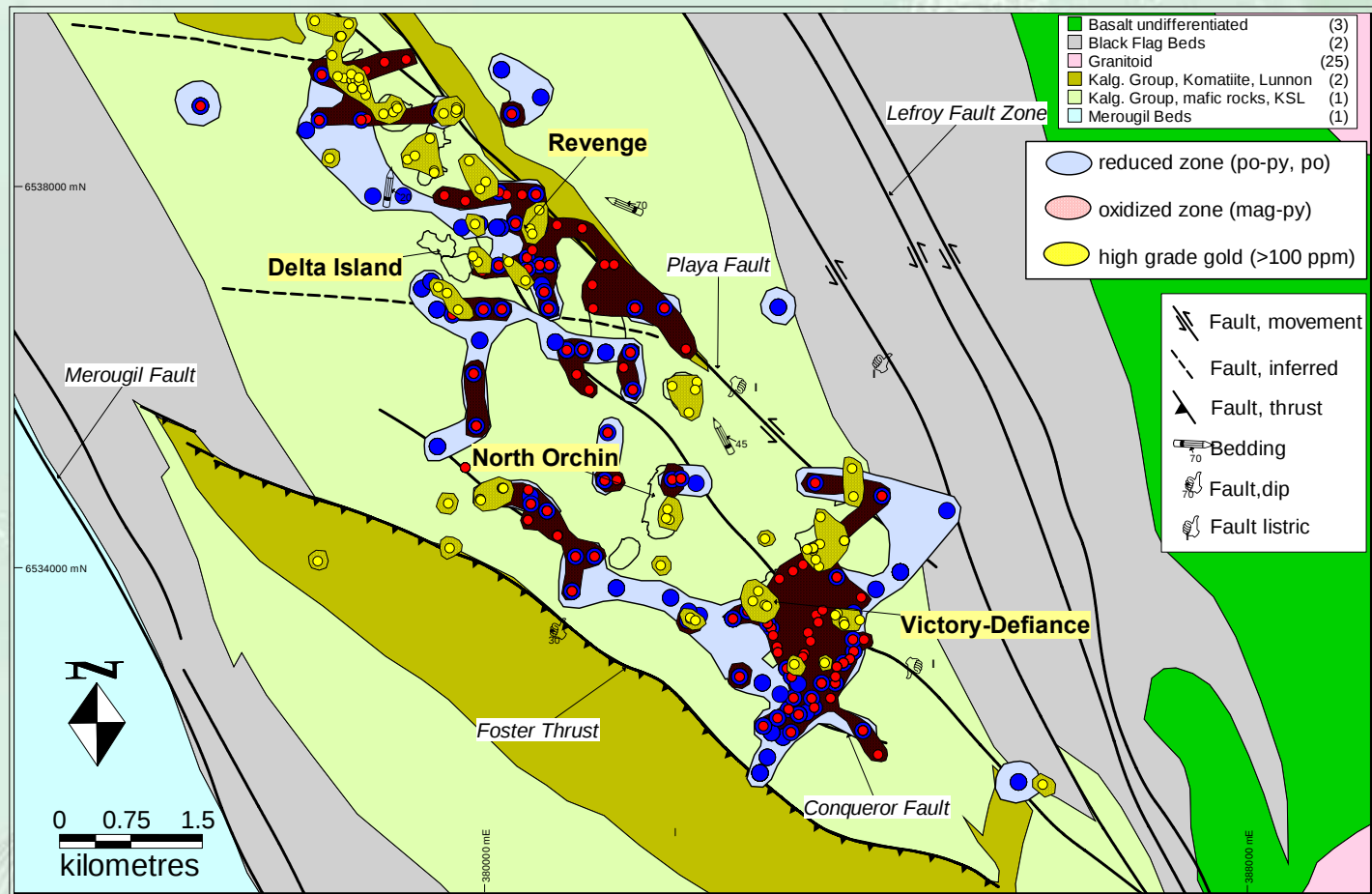
- Gulf of Mexico, onshore Louisiana



Hanor, 1994

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Real World Spatial Gradients

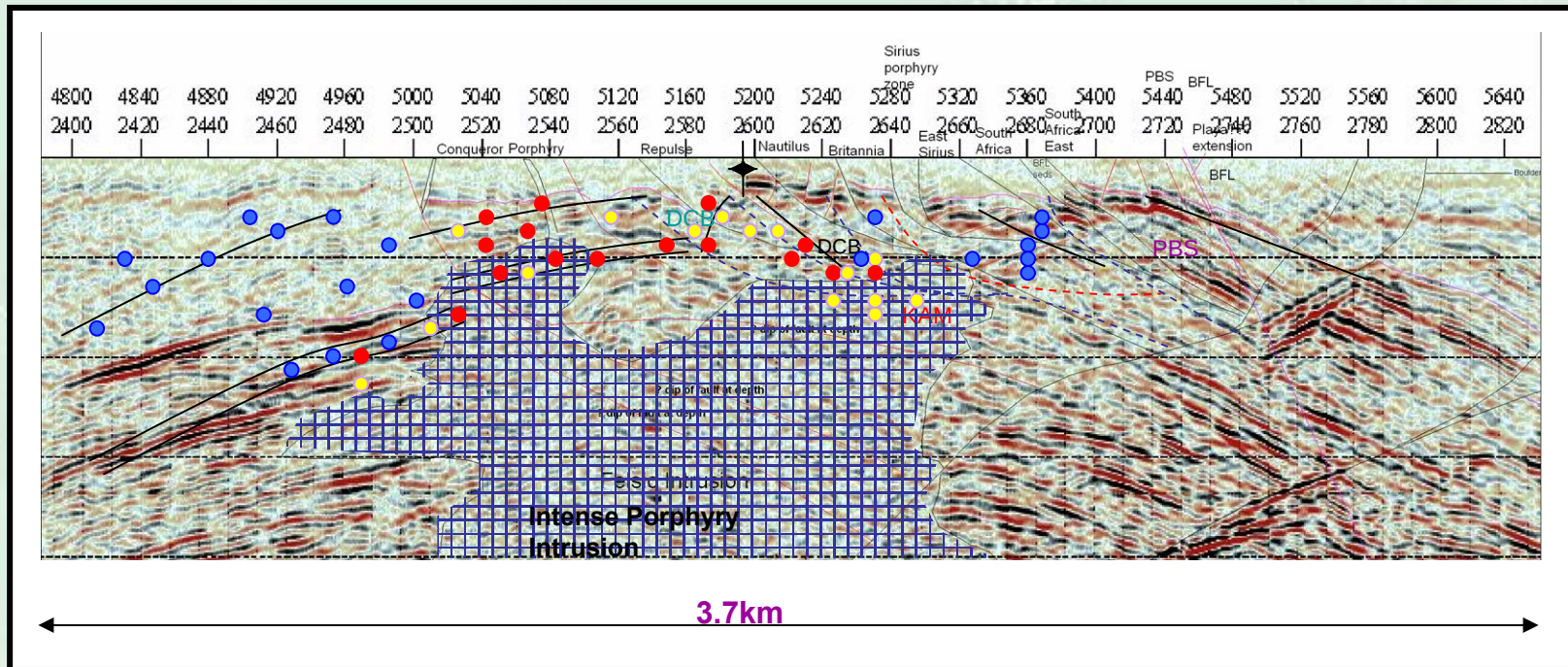


From Y3/Y4/MERIWA team

- St Ives, Western Australia

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Real World Spatial Gradients



From Y3/Y4/MERIWA team

- Redox state (coloured dots) varies with proximity to zone of intense porphyry intrusion

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B, D. Maximising Time....

Steep gradients in P, T (& X) are typically transient

Unless the location is...

- near surface (*T, P?, X?*)
- subject to repeated igneous activity (*T, X, P?*)
- experiences repeated breaching and healing of seal (*P*)
- undergoes repeated deformation-induced dilation effects (*P*)
- possesses a stable hydrology (*X, P?, T?*)



How is This Relevant to Exploration?

- **Provides an alternative view of critical parameters for ore formation**
 - **Gradient in hydraulic potential**
 - **Permeability**
 - **Solubility sensitivity to P, T, C**
 - **Spatial gradient of P, T, C**
 - **Timing & duration**



Fluid Sources

- How many are significant?

- Perhaps only 2:

- Meteoric

- which may be extensively modified

- Magmatic

- which may be H_2O , CO_2 , NaCl or perhaps CH_4 rich

- Meteoric fluids, driven by sub-aerial topography with deep penetration for fluids by faults

- These will only be active at certain times

- Karakoram Mountains
- Anthracite emissions – 20 Mt/d
- Isa Inlier Cu-D3 timing indicated

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Risking

- This analysis yields explicitly *necessary and sufficient* independent parameters suited to use in Bayesian risk analysis

$$P = P_1 \times P_2 \times P_3 \times P_4 \dots$$

(c.f. Lord et al., 2001)

- Not based on specific processes for any one system

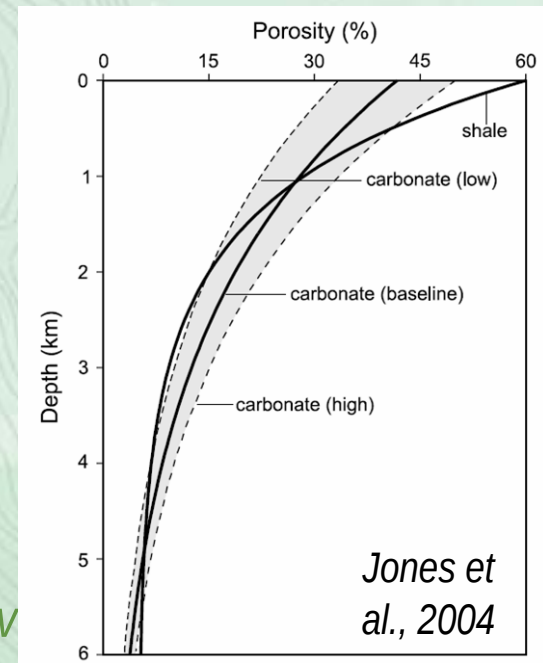
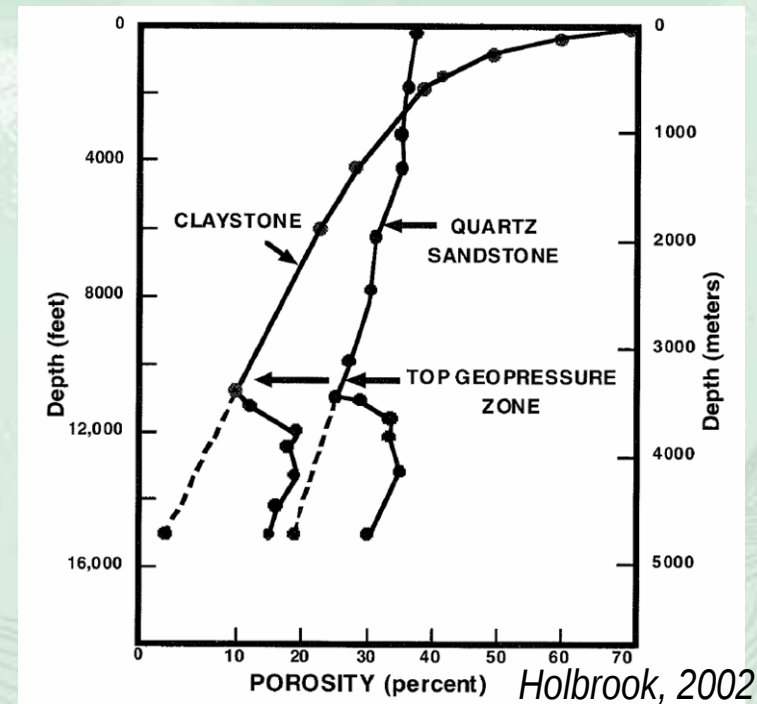
Some Lessons

- **Role of reactivation and other inherited architectures (facies & permeability potential)**
- **There are less variables than at first glance – fluids available only at certain times**



Further Lessons

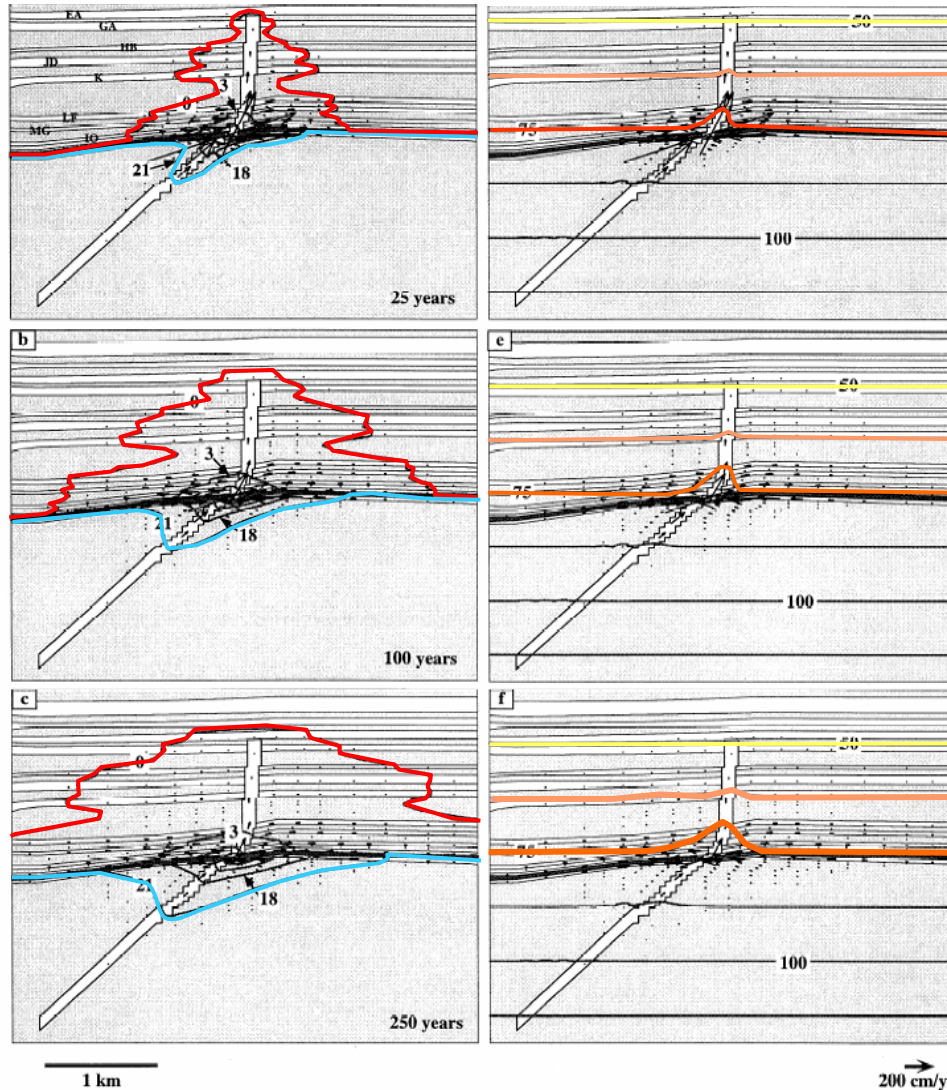
- Much information already exists on the behaviour of basinal systems
- We must use it!



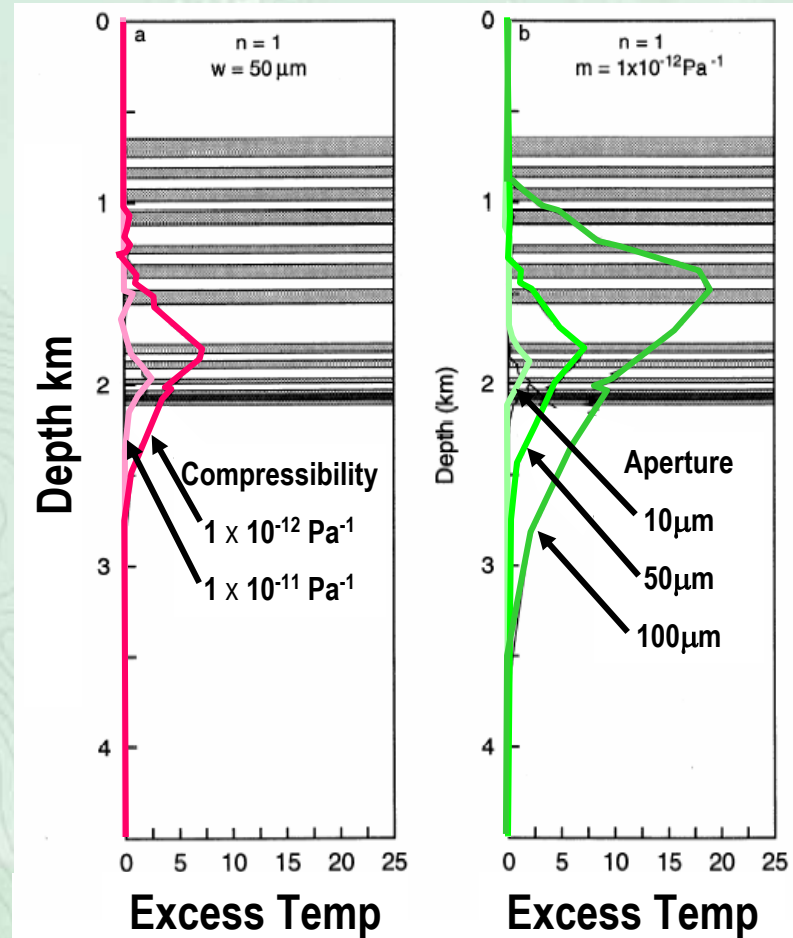
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Excess fluid pressure MPa

Temperature °C



Permeability Evolution



- Fault permeability varies with fluid pressure

predictive mineral discovery Cooperative Research Centre

Roberts et al., 1996 *pmd*^{*}CRC



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In Summary.....

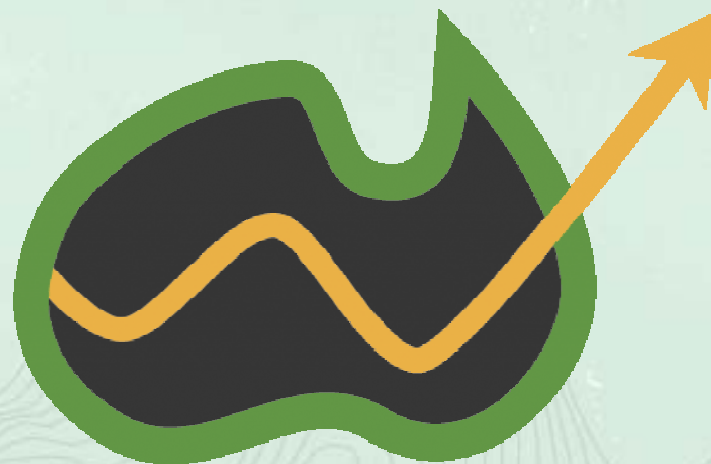
$$\text{Amount of mineral} = \int \frac{k \rho g}{\mu} \cdot \text{Hydraulic Gradient} \cdot \sum \left(\frac{\text{Solubility} \cdot \text{Spatial}}{\text{sensitivities Gradients}} \right) \cdot dt$$

• Critical factors for forming a (giant) ore system:

- Gradient in hydraulic potential **DRIVER**
- Permeability **ARCHITECTURE; PATHWAY**
- Solubility Sensitivity * 
- Spatial gradient of P, T, X **ARCHITECTURE; DEPOSITION**
- Maximum duration **GEODYNAMICS**

**This will be zero if an appropriate fluid is not present!*





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Another view of the 5 Questions

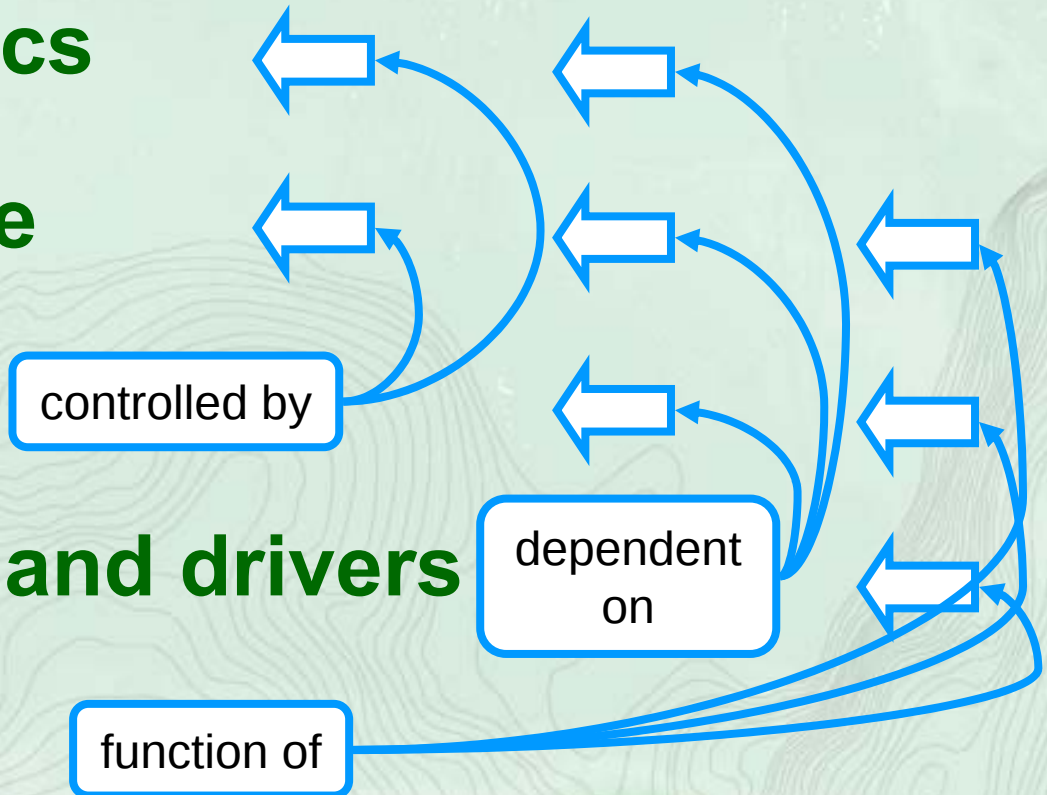
- **Geodynamics**

- **Architecture**

- **Reservoirs**

- **Flow paths and drivers**

- **Deposition**



This Means....

- **The processes of ore deposition are a culmination of normal earth processes – it is their coincidence that leads to mineralisation**
- **Effective understanding must be based on the *geology of economic deposits*, and *not economic geology***

