

# Course of Geothermics

**Dr. Magdala Tesauro**

## **Course Outline:**

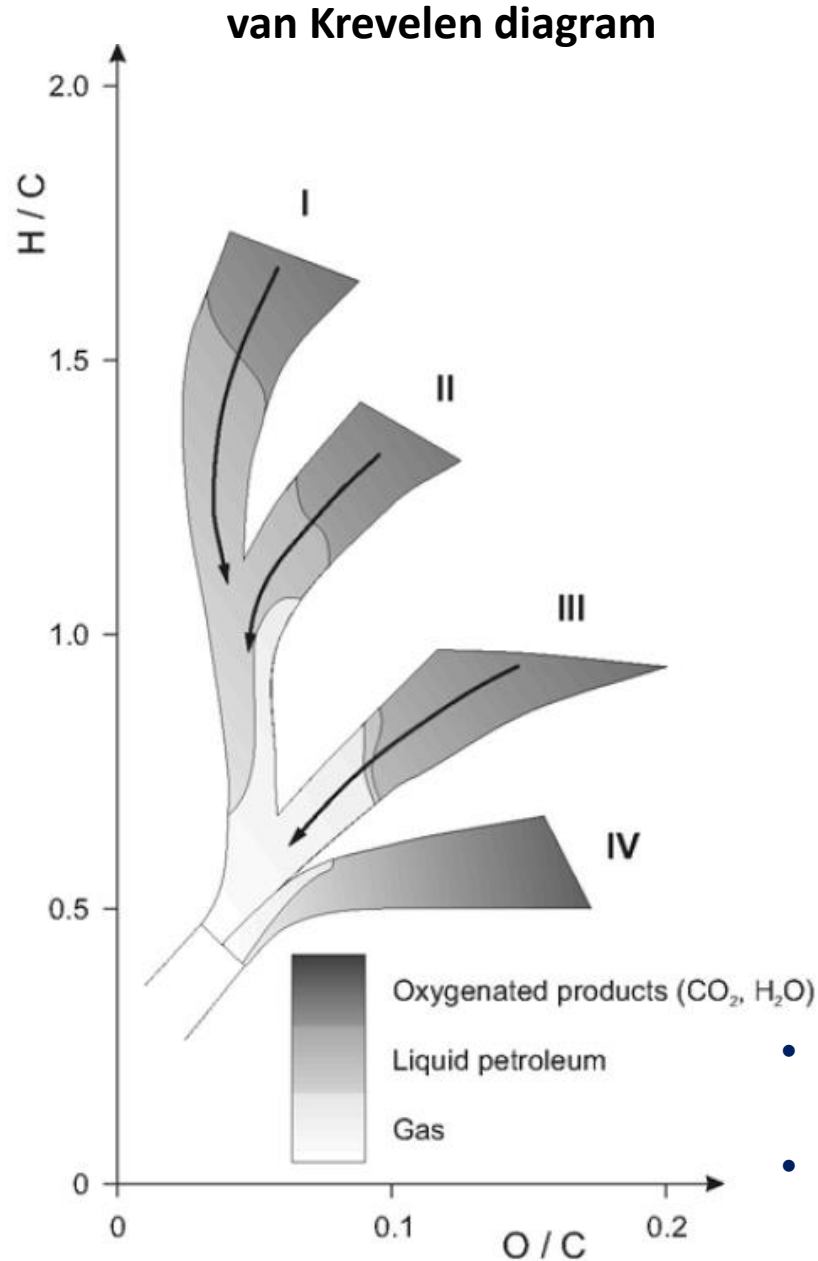
1. Thermal conditions of the early Earth and present-day Earth's structure
2. Thermal parameters of the rocks
3. Thermal structure of the lithospheric continental areas (steady state)
4. Thermal structure of the lithospheric oceanic areas
5. Thermal structure of the lithosphere for transient conditions in various tectonic settings
6. Heat balance of the Earth
7. Thermal structure of the sedimentary basins
- 8. Thermal maturity of sediments**
9. Mantle convection and hot spots
10. Magmatic processes and volcanoes
11. Heat transfer in hydrogeological settings
12. Geothermal Systems

# Thermal evolution of sedimentary basins

- **Hydrocarbons** are derived from the metamorphism of organic material, OM (everything from woody cell walls, spore cases, leaf cuticles, pollen, resins, algae, bacteria and plankton), a process involving the expulsion of volatiles, gas, liquids, and oil.
- The **thermal maturity** of a rock is a measure of the degree to which this metamorphism has progressed (or degree of thermal degradation of the OM).
- Sedimentary OM can be divided into **bitumen**: the fraction of sedimentary OM soluble in typical organic solvents (e.g., chloroform, benzene) and **kerogen**: the fraction of sedimentary OM insoluble, left in the rocks after washing with solvents.
- The type and maturity of kerogen depends on the content of carbon (*C*), hydrogen (*H*), and oxygen (*O*).
- The temperature limits of the **oil window** are within the range of about 50-150 °C (323–423 K) and they depend on the age of source rocks with significant decrease in *T* with increase in age.
- For deposits of ~300 Myr, the formation of oil starts at  $T < 50^{\circ}\text{C}$  and continues up to  $T \geq 100^{\circ}\text{C}$  and gas is formed above this limit up to  $T \sim 215^{\circ}\text{C}$ . Above that *T* both oil and gas are destroyed. For rocks ~10 Myr, the oil generation starts at *T* of  $\sim 83^{\circ}\text{C}$  and continues to about  $170^{\circ}\text{C}$ , above which gas is usually formed to *T* of  $\sim 300^{\circ}\text{C}$ .
- Within the oil window increasingly more oil is generated with increase in *T* until maximal production is reached, at which point the quantity of oil decreases and increasingly more gas is formed.

# Thermal evolution of sedimentary basins

## van Krevelen diagram



**I Hydrogen-rich, oxygen-poor:** they derive from the deposition in lakes as bacterial and vegetable waxes or algae.

**II Intermediate H:C and O:C ratios:** they derive from bacterially altered marine organic matter. With some terrigenous material (e.g., leaf cuticles and spores).

**III Hydrogen-poor, oxygen rich:** they derive from terrigenous woody OM and cuticular waxes, associated with near-shore deltaic environments.

**IV very low H:C, high O:C:** they consist of highly reworked OM with no hydrocarbon generative potential.

- The kerogens trace characteristic path through the van Krevelen diagram during the gradual maturation (alteration of atomic ratios).
- The amount of petroleum generated and expelled increases with the increase of H/C.

# Thermal evolution of sedimentary basins (chemical kinetics)

- Temperature (exponential dependence) and time (linear dependence) are the main factors influencing OM maturation.
- Arrhenius equation, expressing the relationship between temperature and rate of chemical reactions, suggests that reaction rates ( $k$ ) increases exponentially with  $T$ , but the rate of increase in reaction rate slows down with increasing  $T$ :

$$k(T) = A \times \exp(-E/RT)$$

$A$ =frequency factor ( $10^{13}$ - $10^{16}$  s<sup>-1</sup>), it is possibly related to the vibrational frequency of the reacting molecules

$E$ =Activation energy, 160-340 kJ mol<sup>-1</sup> (energy barrier that must be overcome before the reaction can proceed)

$R$ =universal gas constant (8.314 JK<sup>-1</sup> mol<sup>-1</sup>)

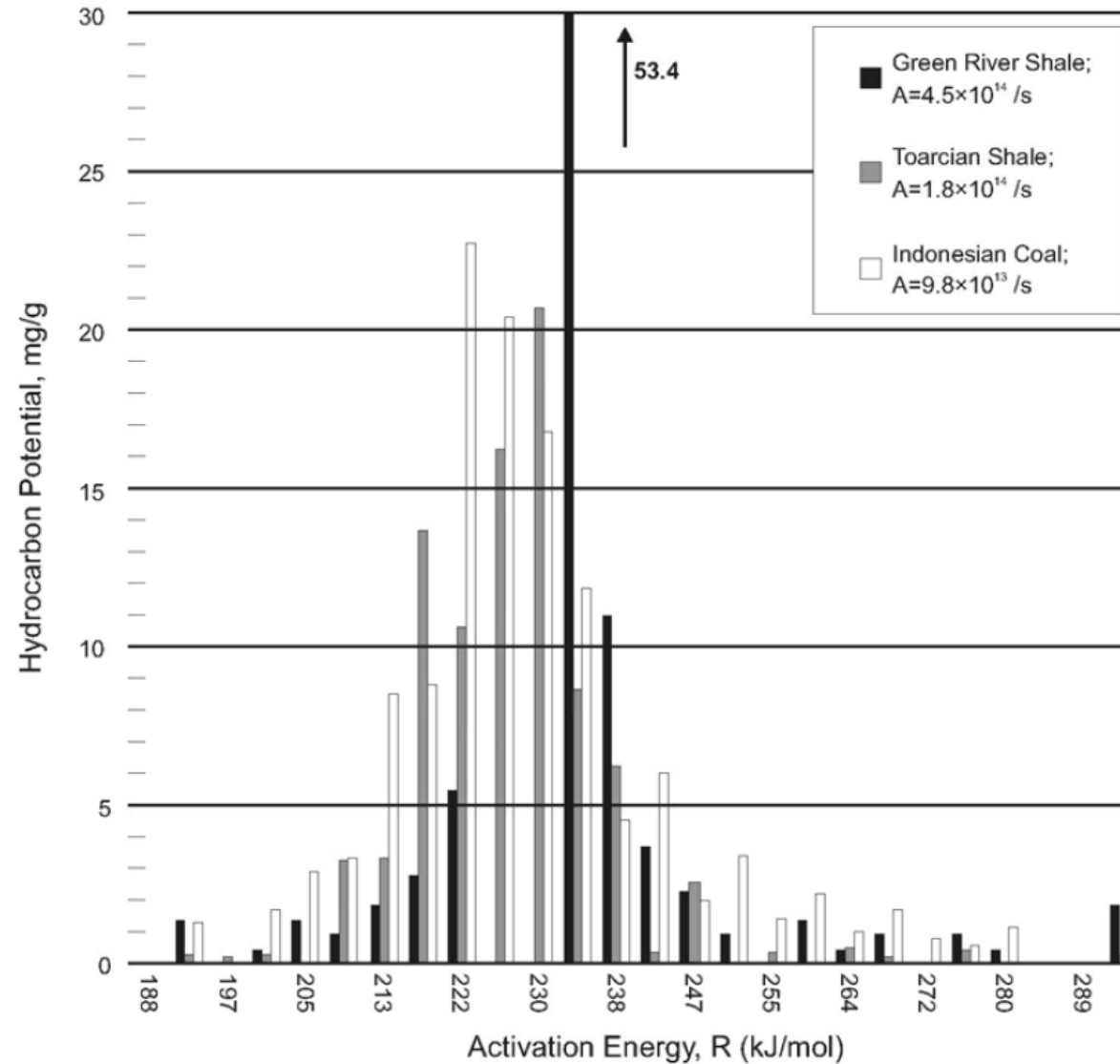
$E$  has been determined using **pyrolysis**, which artificially forces a source rock to release all of its potential hydrocarbon product, by increasing  $T$  at constant heating rate,  $r$ .

**Kinetic Parameters for the Generation of Oil from Type II Source Rocks**

Source	$E$ (kJ mol <sup>-1</sup> )	$A$ (s <sup>-1</sup> )	Sulphur Content (wt% kerogen)
Monterey shale	143.4	$2.224 \times 10^7$	11.0
Phosphoria shale	178.7	$1.338 \times 10^{10}$	9.0
Alum shale	201.3	$4.899 \times 10^{11}$	7.4
Woodford shale	218.3	$1.792 \times 10^{13}$	5.4

Source: Hunt et al. (1991)

# Thermal evolution of sedimentary basins (chemical kinetics)



Kinetic data derived for the generation of hydrocarbon for Type I and Type II kerogens

# Thermal evolution of sedimentary basins (chemical kinetics)

We can represent the kerogen as initially having a certain potential to produce hydrocarbons,  $\mathbf{M}_0$ , which decreases as the hydrocarbon yield,  $\mathbf{m}$ , increases:

$$-(\partial \mathbf{M} / \partial t) = (\partial \mathbf{m} / \partial t) = k \times (\mathbf{M}_0 - \mathbf{m}) \quad k=\text{reaction rate}$$

The rate at which hydrocarbons are released from samples is  $T$  dependent:

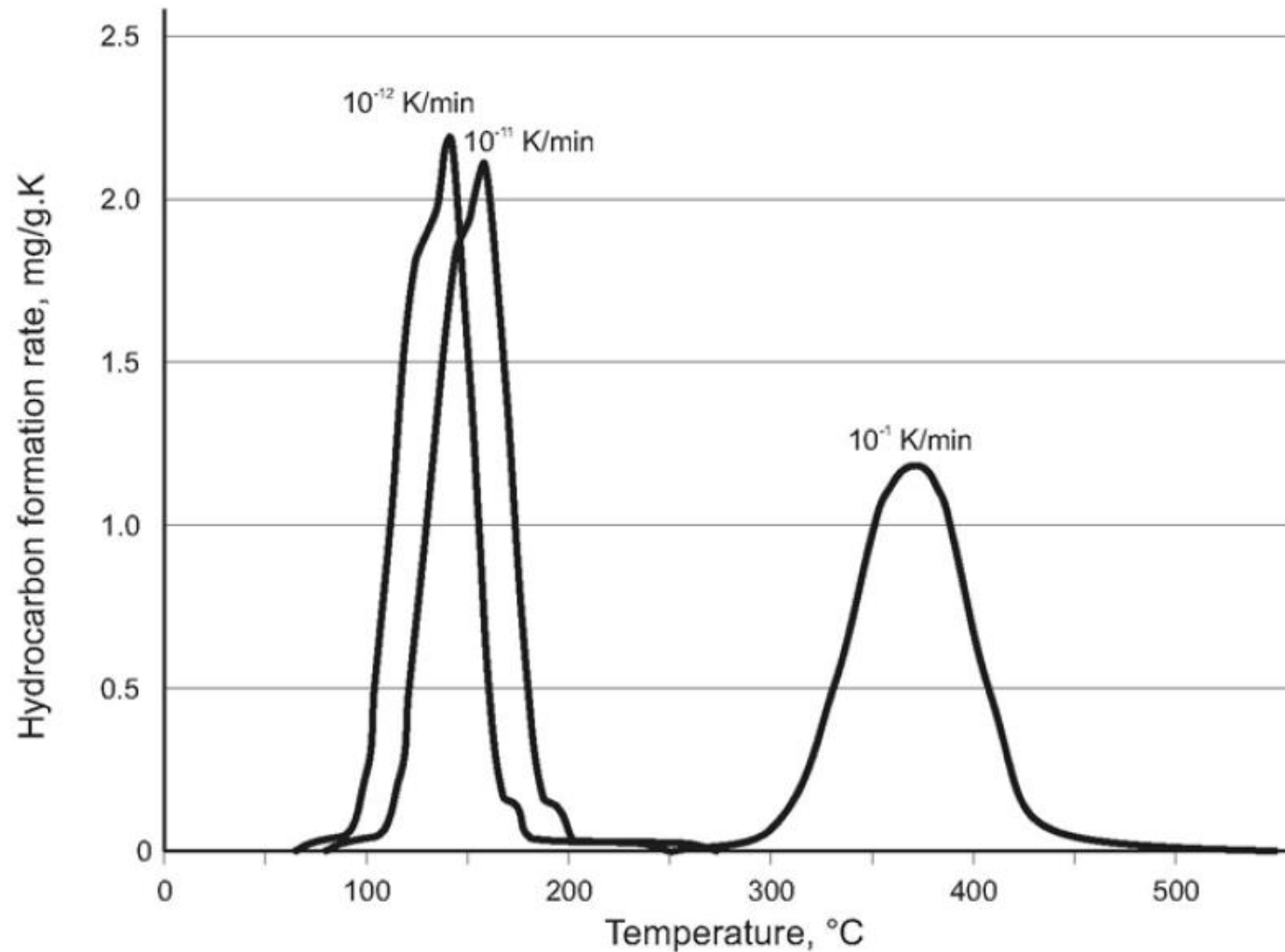
$$\frac{\partial \mathbf{m}}{\partial T} = \sum_{i=1}^n \mathbf{M}_{i0} \frac{A}{r} \left( -\frac{E_i}{RT} - \frac{A}{r} \mathbf{J}_i \right) \quad \mathbf{J}_i = \int_0^T \exp\left(-\frac{E_i}{RT}\right) dT \quad r=\text{heating rate}$$

- According to Connan (1974), the threshold of the principal zone of oil generation is related to the logarithm of the age of the formation.
- The cumulative effect of increasing  $T$  can be evaluated from the maturation integral (the reaction rate integrated over time), estimated if decompacted burial history, heat flow through time, and thermal conductivity of sedimentary basin is known:

$$C = A \int_0^t \exp(-E_a/RT) + C_0$$

$C_0$ =original level of maturation of the organic material at  $t=0$  (depositional time)

## Thermal evolution of sedimentary basins (chemical kinetics)



- Petroleum forms at a lower and narrower  $T$  range at geological heating rates than during pyrolysis.
- $T$  of the sediments should be in the range 70-120°C for significant period of time, at cooler  $T$  the reaction rate is too slow to produce hydrocarbons, while at higher  $T$ , the rate is too rapid to allow expulsion before the oil degrades.

# Paleotemperature Indicators

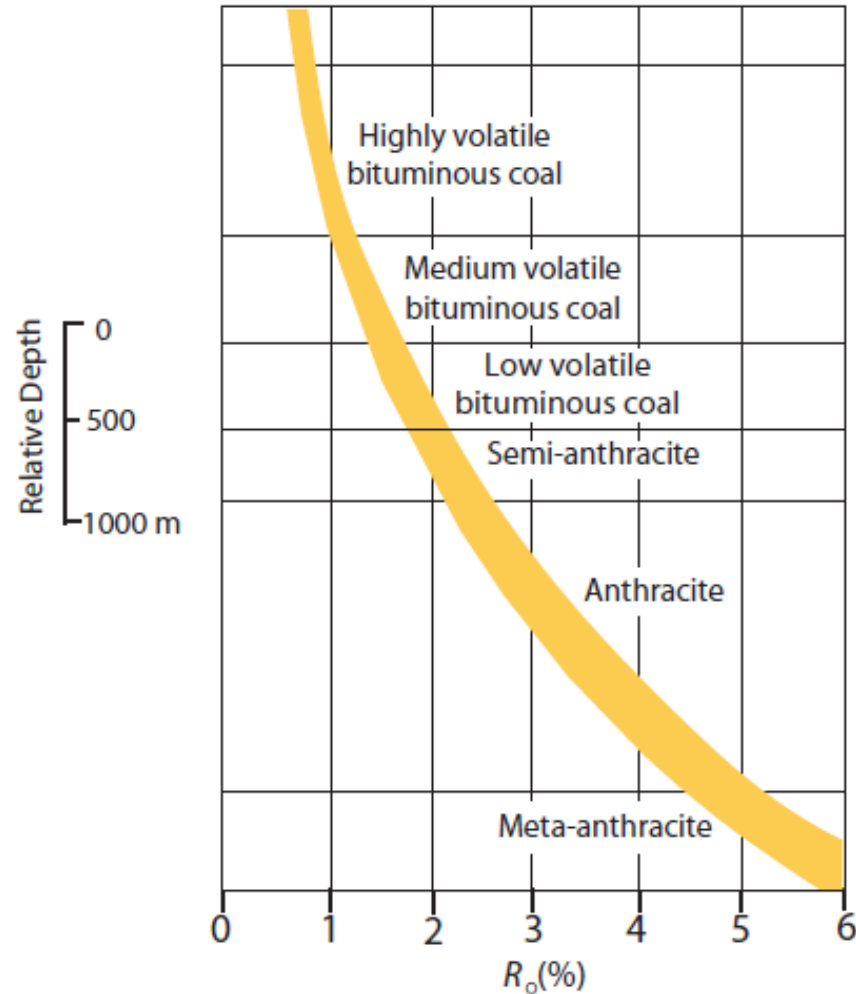
## Vitrinite Reflectance (VR)

- **Vitrinite is a detrital plant material**, transported with other sediments to the final deposition zone. It is a class of macerals, microscopic organic components of coal.
- Macerals can be broadly divided into three classes on the basis of morphology, reflectance in white light, and fluorescence:
- **Liptinite**: It is derived from the waxy, lipid-rich and resinous part of plants (e.g., leaf cuticles, algae). Liptinite is the least reflective among the macerals and fluoresces under ultraviolet light.
- **Vitrinite**: It is derived from plant cell walls, cell contents or precipitated gels. The reflectance of vitrinite lies somewhere between that of liptinite and inertinite, and increases with the rank of coal. It may show a weak brownish fluorescence.
- **Inertinite**: It has the same derivation of vitrinite, but has been oxidised, altered or degraded in early stage of coal formation. Inertinites are the most reflective, but do not fluoresce.
- The percentage of carbon increases with increase of burial time and  $T$ . A higher **reflectance** implies that the coal is of a higher rank (percentage) and has been raised to a higher  $T$  or buried for long time.
- **Reflectance ( $R_0$ ) or Vitrinite Reflectance (VR)** is a measure of the percentage of an incident beam of polarized white light that is reflected off a polished surface (optical parameter) and gives an indication of the geothermal gradients in the history of the basins.
- Estimates of paleo- $T$  based on the analysis of VR relies on the **property of vitrinite to undergo increasing reflectance with increasing  $T$** . The amount of light reflected by vitrinite particles ( $R_0$ ) is affected by time and  $T$  of burial and irreversibly changes with increases in  $T$ .
- **Vitrinite is an anisotropic, birefractant substance**: There are two identical reflectance maxima at  $180^\circ$  separation, with minima mid-way between the maxima. The magnitude of anisotropy is related to sediment compaction and increases with increasing rank.

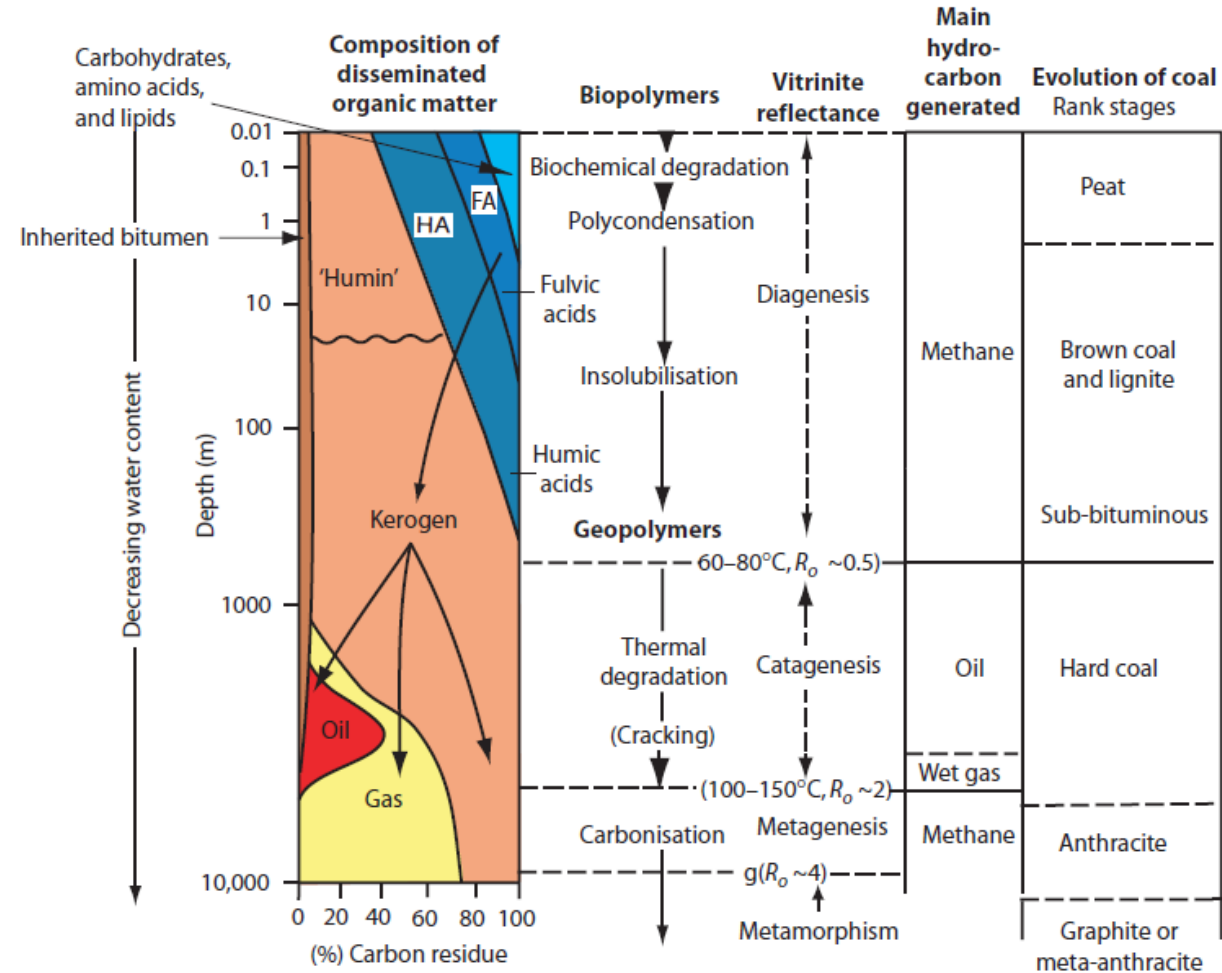


# Paleotemperature Indicators

## Vitrinite Reflectance (VR)



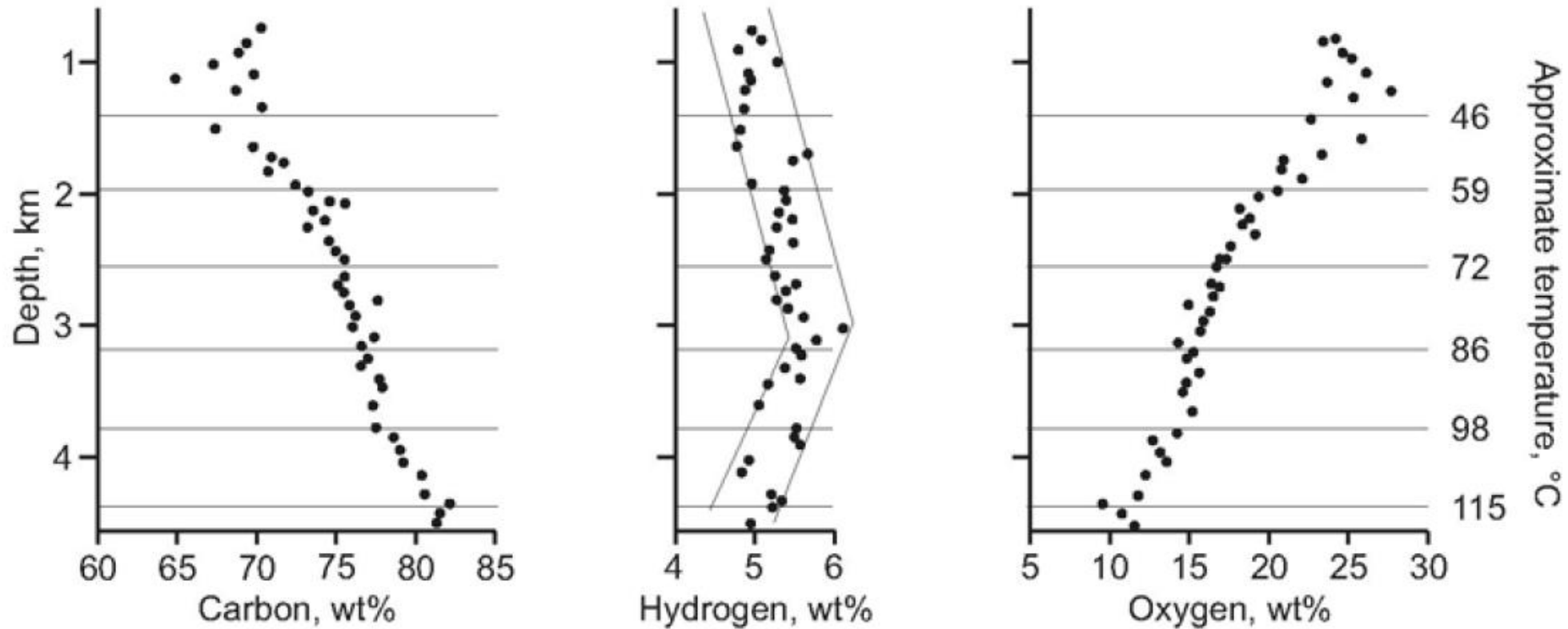
### Evolution of organic matter and coal



- VR is the most widely used indicator of maturity of OM.  $R_o < 0.5-0.7\%$  indicates that the source rocks are immature.
- Progressive maturation of OM causes a coalification process, which changes peat to anthracite through brown coal and bituminous coal.

# Paleotemperature Indicators

## Vitrinite Reflectance (VR)



Changing atomic composition of coal with  $P$  and  $T$

# Paleotemperature Indicators

## Vitrinite Reflectance (VR)

Since vitrinite is an anisotropic substance, there are two methods to estimate the reflectance:

- (1) It records the maximum reflectance ( $R_{vMAX}$ ) for each vitrinite grain. It is more precise, but it requires rotation of the microscope through  $360^\circ$  to detect the maxima.
- (2) It records only a single randomly oriented reflectance for each grain ( $R_o, R_o\%, R_v\%$ ). It is faster, but less precise.

$$R_{vmax}(\%) = R_o(\%) \times 1.066$$

The thermal history of a rock can be broken down into a number of discrete  $T$  intervals (Lopatin, 1976), so that the thermal maturity or time-temperature index (TTI) of a kerogen is:

$$TTI = \sum(\Delta t_i)(2^{n_i})$$

$\Delta t_i$ =time(My) spent by the kerogen in the  $i$ th  $T$  interval

$n_i$ =integer weighting factor for the  $i$ th interval

Each  $T$  interval covers  $10^\circ\text{C}$  and is assigned a weighting factor  $n_i$  (which changes by 1 for each  $10^\circ\text{C}$ ). The reference interval for which  $n=0$  corresponds to  $100\text{-}110^\circ\text{C}$ .

According to McKenzie (1981), the TTI function is a continuous integral:  $TTI = \int_{t_s}^{t_f} 2^{[(T_t - 105)/10]} dt$

$T_t$ =time-dependent temperature ( $^\circ\text{C}$ ),  $t_s$  and  $t_f$ =start and finish times (Myr)

## Paleotemperature Indicators

### Vitrinite Reflectance (VR)

**Question:** Organic matter is deposited within sediment at a surface temperature of 20°C. It is buried to a temperature of 100°C at a constant rate over a period of 40 Myr. Sedimentation then slows, and after another 30 Myr the temperature has increased to 130°C. How mature is the sediment, expressed in terms of TTI?

$$(1) \quad \text{TTI} = \Sigma(\Delta t_i)(2^{n_i})$$

$$(2) \quad \text{TTI} = \int_{t_s}^{t_f} 2^{[(T_t - 105)/10]} dt$$

**1<sup>st</sup> Answer:** In the first period of burial, the sediment passes through eight temperature intervals in 40 Ma, spending 5 Ma in each. The initial interval (20-30°C) has a weighting factor  $n^{-8}$ .

$$\text{TTI} = \Sigma(\Delta t_i)(2^{n_i}) = (\Delta t_i)\Sigma(2^{n_i}) = 5 \times (2^{-8} + 2^{-7} + 2^{-6} + 2^{-5} + 2^{-4} + 2^{-3} + 2^{-2} + 2^{-1}) = 4.98$$

The sediment then spends 10 Myr in each of the next three temperature intervals,  $n = 0, 1$  and  $2$ . The total maturity is then:

$$\text{TTI} = 4.98 + 10 \times (2^0 + 2^1 + 2^2) = 74.98$$

## Paleotemperature Indicators

### Vitrinite Reflectance (VR)

**2<sup>nd</sup> Answer:** The thermal history should be broken into two segments:

The first:

$$T_t = 20 + 2t, \quad 0 < t < 40 \text{ Ma}$$

the second:

$$T_t = 60 + t, \quad 40 < t < 70 \text{ Ma}$$

$$\text{TTI} = \int_{t_s}^{t_f} 2^{[(T_t - 105)/10]} dt$$

$$\text{TTI} = \int [2^{(-8.5+0.2t)}] dt = 2^{-8.5} \int [2^{0.2t}] dt = 2^{-8.5} \times [1/(0.2 \times \ln 2)] \times [2^8 - 2^0] = 5.08$$

$$\text{TTI} = \int [2^{(-4.5+0.1t)}] dt = 2^{-4.5} \int [2^{0.1t}] dt = 2^{-4.5} \times [1/(0.1 \times \ln 2)] \times [2^7 - 2^4] = 71.41$$

$$\text{TTI} = 5.08 + 71.41 = 76.49$$

# Paleotemperature Indicators

## Vitrinite Reflectance (VR)

It is possible to estimate the max paleo  $T$  ( $T_{max}$ ) experienced by a sample of vitrinite from its reflectance (from a compilation of > 600  $R_o$ - $T_{MAX}$  ( $^{\circ}\text{C}$ ) data pairs):

$$\ln(R_o) = 0.0078 \times T_{max} - 1.2$$

**Waples (1980)** presented a calibration between  $TTI$  (using discrete  $T$  intervals) and the reflectance of vitrinite:

$$\ln(TTI) = A \times \ln(R_o) + B \quad \begin{matrix} A=4.144 & B=4.168 \\ \text{(over the range } 0.5 \leq R \leq 5.0\%) \end{matrix}$$

The estimates of vitrinite reflectance are used to estimate the thermal maturity of a kerogen in terms of  $TTI$  (forward modelling).

According to **Issler (1984)**:

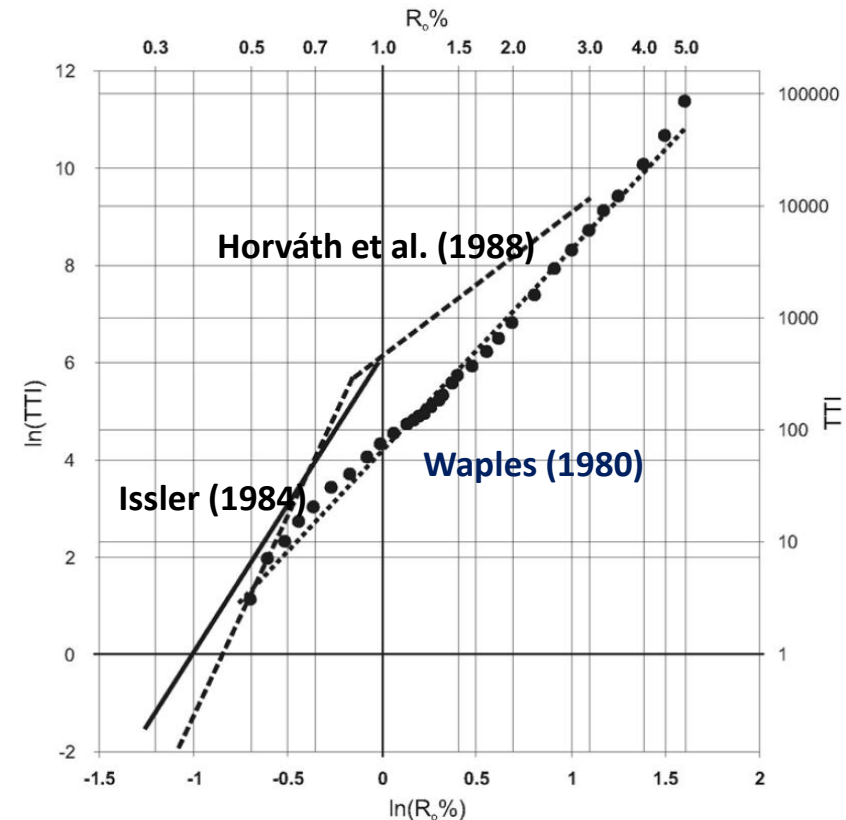
$A = 6.184$  and  $B = 6.115$

(over the range  $0.29 \leq R_o \leq 0.98\%$ ).

According to **Horváth et al. (1988)**:

$A = 8.278$ ,  $B = 6.992$  for  $TTI < 292$  (for  $R_o < 0.853\%$ )

$A = 2.936$ ,  $B = 6.143$  for  $TTI > 292$  (for  $R_o > 0.853\%$ )



## Paleotemperature Indicators

### Vitrinite Reflectance (*VR*)

what VR value should we expect?

According to Waples' (1980) calibration:

$$\ln(\text{TTI}) = A \times \ln(R_o) + B$$

$$\ln(74.98) = 4.144 \times \ln(R_o) + 4.168$$

$$\ln(R_o) = [\ln(74.98) - 4.168] / 4.144 = 0.036$$

$$R_o = 1.04\%$$

According to Issler (1984) calibration:

$$A = 6.184 \text{ and } B = 6.115: \quad \ln(76.49) = 6.184 \times \ln(R_o) + 6.115 \quad R_o = 0.75\%$$

According to Horvath et al. (1988) calibration:

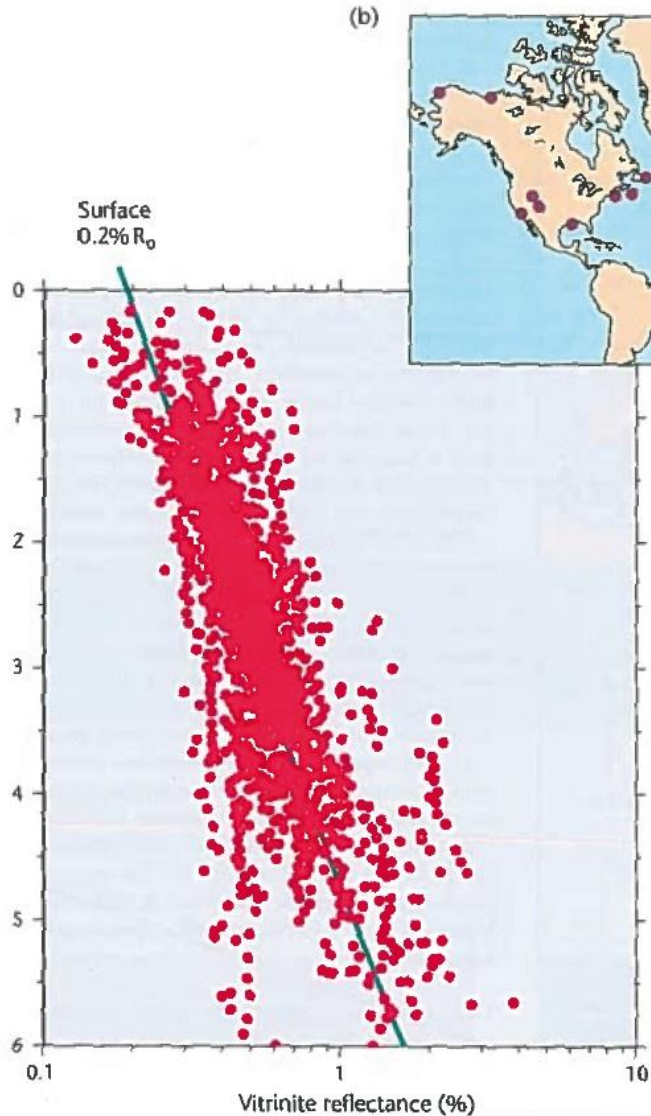
$$A = 8.278 \text{ and } B = 6.992: \quad \ln(76.49) = 8.278 \times \ln(R_o) + 6.992 \quad R_o = 0.73\%$$



# Paleotemperature Indicators

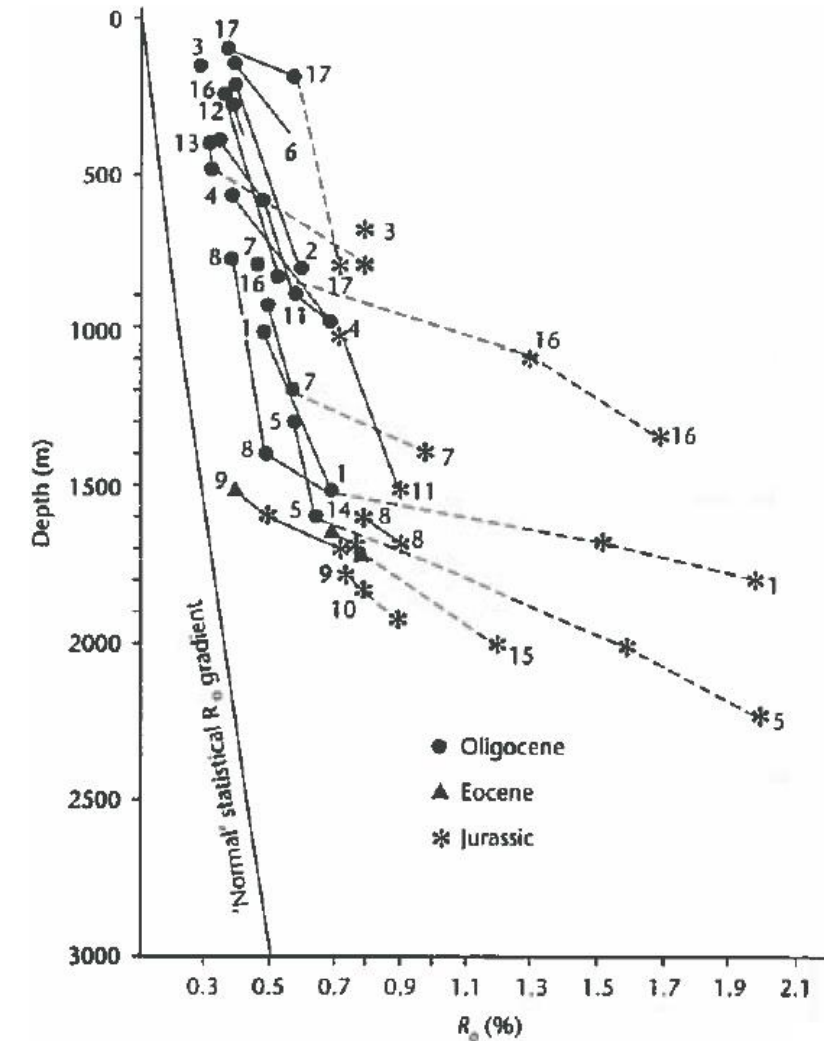
## Vitrinite Reflectance (VR)

### 28 World extensional basins



- There is a linear relationship between depth and  $\log R_0$  for the sedimentary basins, characterized by no variation of the geothermal gradients with time (due to the exponential evolution of  $OM$  with  $T$ ).
- A break in the slope of these curves identify the occurrence of periods of different geothermal gradient: In the Upper Rhine Graben, the rifting occurred in the late Eocene caused higher than normal maturity (sedimentary rocks have high  $R_0$  in relation to their burial).

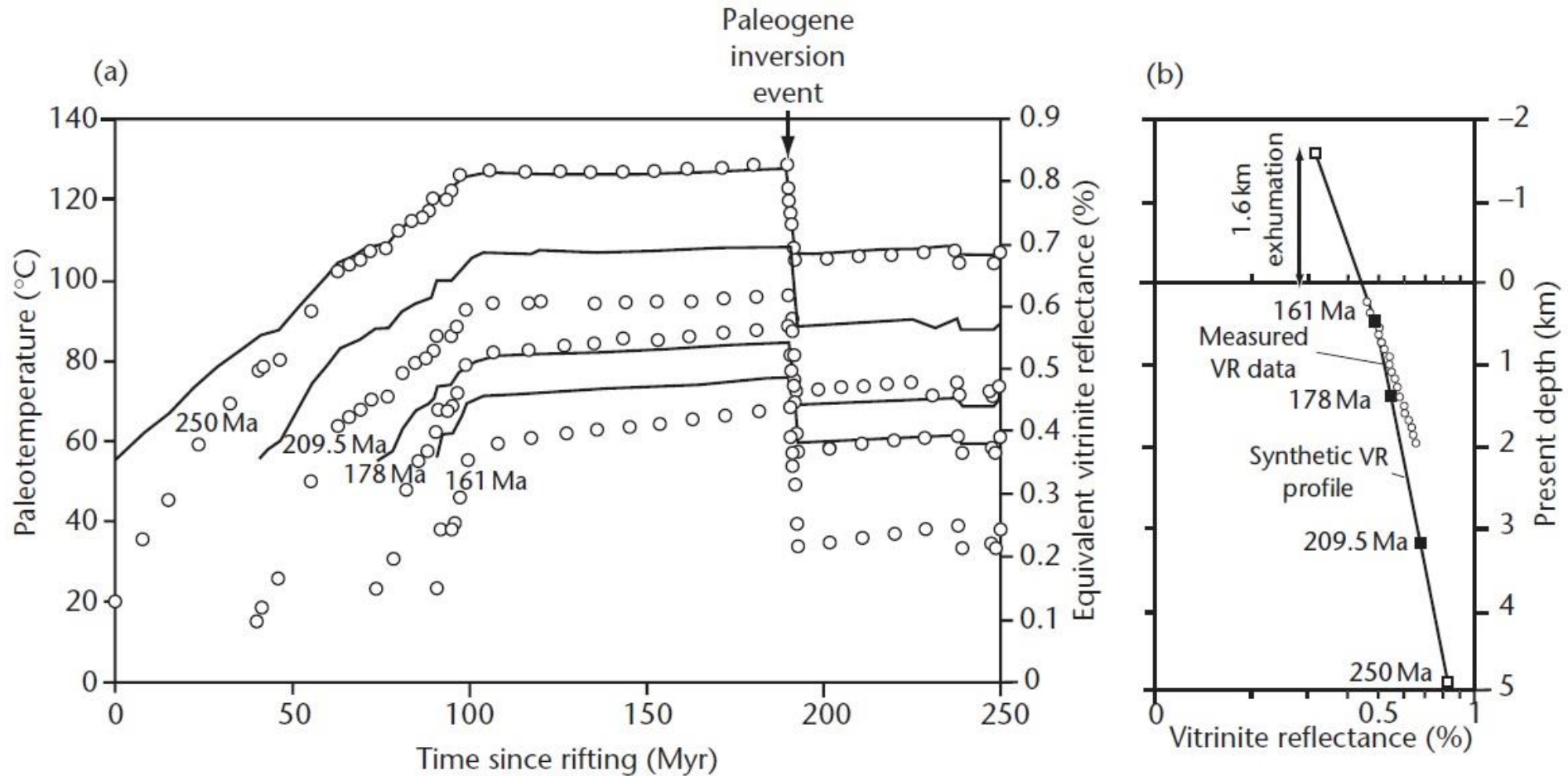
### Upper Rhine Graben





# Paleotemperature Indicators

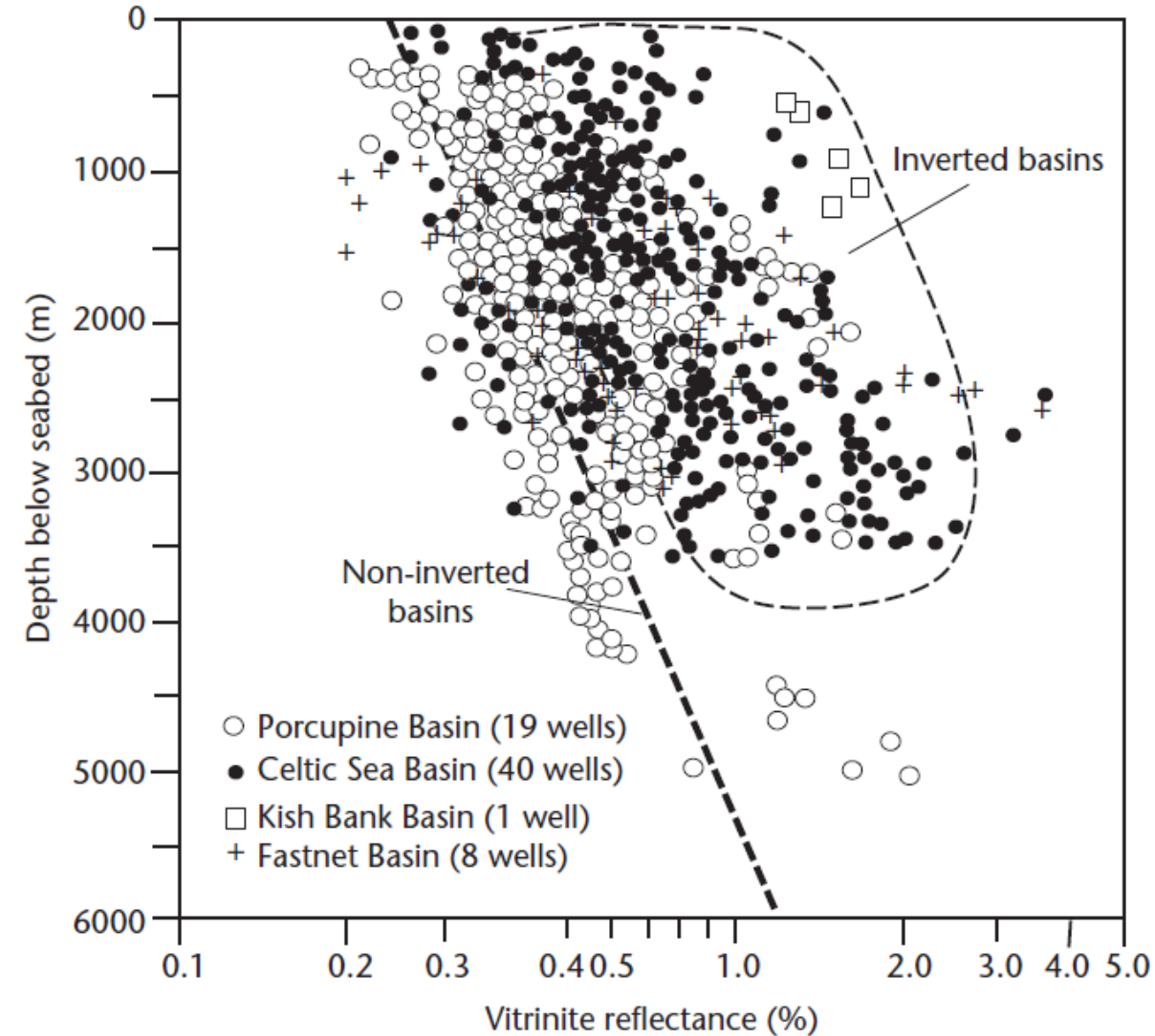
## Vitrinite Reflectance (VR)



Paleotemperature and equivalent vitrinite reflectance versus time for four different horizons in a well in the extensional Irish Sea Basin ( $\beta = 1.5$  at 250 Ma) with internal heat generation ( $A = 9.6 \times 10^{-10} \text{ W kg}^{-1}$ ;  $h_r = 10 \text{ km}$ ) and variable thermal conductivity.

# Paleotemperature Indicators

## Vitrinite Reflectance (VR)



- Basins that experienced tectonic uplift and exhumation show  $R_0$  profiles indicating higher thermal maturity than expected from their depth of burial.
- Maximum depths of burial can be estimated from  $R_0$  values:

$$\ln R_0 = 0.00096T - 1.4 \quad (\text{Barker and Pawlewicz, 1986})$$

$T$  must be converted to depth using geothermal gradient (high uncertainty)

# Paleotemperature Indicators

## Vitrinite Reflectance (*VR*)

**Drawbacks of VR measurements arise from:**

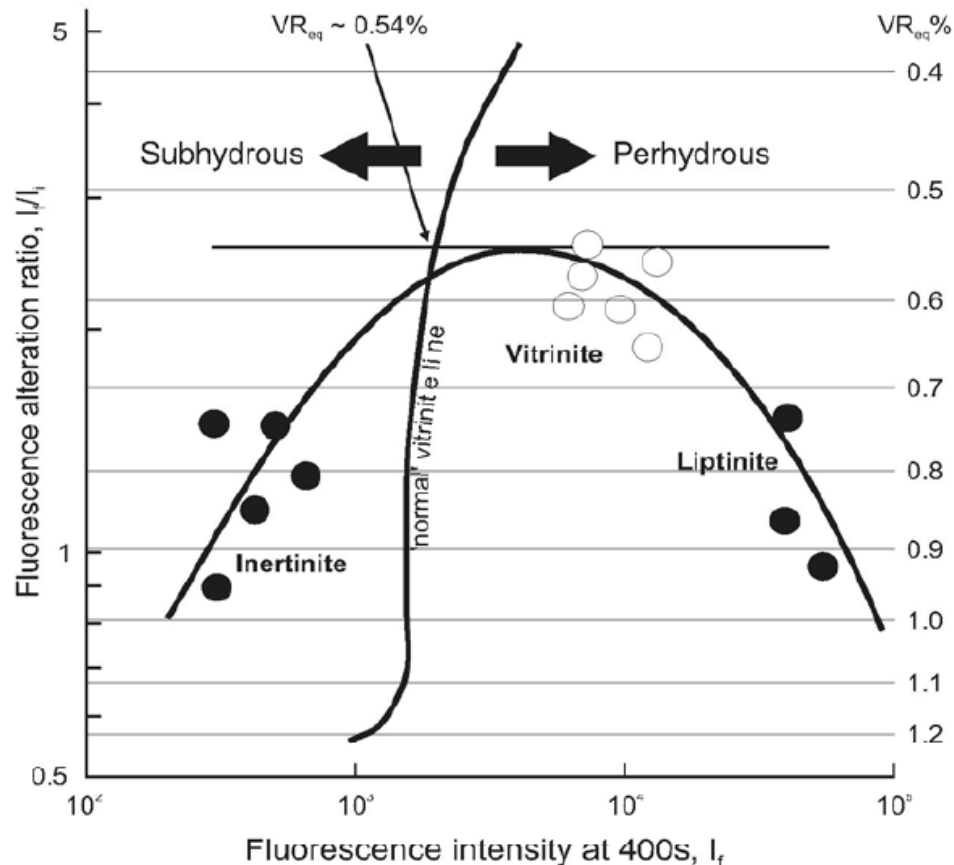
- $R_o$  measurements from different maceral types (within or outside the vitrinite group) may be significantly different.
- *OM* may have been reworked.
- Lack of plants having vitrinite before Devonian age.
- $R_o$  tend to be unreliable at low level of maturity ( $R_o < 0.7-0.8$ ).
- At high  $T$  corresponding to a depth  $> 4$  km the vitrinite maceral is highly anisotropic (accurate measurements are difficult).
- Uncertainty in the knowledge of parameters in Arrhenius equation:  
( $E_a = 200-300 \text{ kJmol}^{-1}$   $A = 2.5 \times 10^{10} - 7.48 \times 10^{18} \text{ s}^{-1}$ )

# Paleotemperature Indicators

## Fluorescence Alteration of Multiple Macerals (FAMM)

- This method involves measuring the fluorescence emission from polished maceral samples in response to a laser beam focused to  $\sim 1\text{-}2\mu\text{m}$  diameter.
- The intensity of the emitted red (625 nm) fluorescence varies with time and the effect is quantified by measuring the intensity at the beginning ( $I_i$ ) and end ( $I_f$ ) of a specified period (400 s).

Fluorescence alteration diagram



- The abscissa  $I_f$  is controlled by maceral types (Inertinite the lowest values and liptinite the highest).
- The ordinate  $I_f/I_i$  is mainly controlled by the thermal maturity, decreasing as maturity increases.
- To determine the equivalent vitrinite reflectance ( $VR_{eq}$ ) of the sample, a best-fit second-order polynomial curve is placed through the complete data set.
- This technique has been calibrated such that the curve maximum indicates  $VR_{eq}$  for samples with  $VR_{eq} < 0.9$ .

## Paleotemperature Indicators

### Thermal Alteration Index (*TAI*)

- The ***TAI*** index is an arbitrary scale (from 1 to 4) that ranks the gradual change in colour and opacity of liptinite (spore, pollen, and leaf cuticle material) with *T*.
- ***TAI*** 1 is equivalent to the lowest levels of organic maturity and is assigned to liptinite particles showing a greenish-yellow color.
- ***TAI*** and maturity increase as the colour of liptinite progresses through pale yellow, yellow amber, deep red-brown, and opaque black at ***TAI*** 4.
- The entire oil generation window falls within a narrow range of ***TAI*** values (~2.0-2.7). Below ***TAI*** 2 no oil can be expected, while above ***TAI*** 3.5, *T* is too high (gas zone).
- Paleo*T* are measured with a precision of  $\pm 20^{\circ}\text{C}$ .

# Paleotemperature indicators

## Conodont Alteration Index (CAI)

- The **CAI** index refers to the coloration of conodont fossils that reflects the max paleo*T*.
- It has eight levels, from the lowest to the highest *T*, related to the gradual change in colour and opacity of conodonts with *T*.

**Conodont Alteration Index (CAI) and Corresponding Palaeotemperature**

CAI	Colour	Palaeotemperature (°C)	Thermal Maturity
1.0	Pale yellow	50–80	Immature
1.5	Very pale brown	50–90	Early-mature
2.0	Brown to dark brown	60–140	Mature
2.5		85–180	Late-mature
3.0	Very dark grey	110–200	Over-mature
3.5		150–260	Over-mature
4.0	Light black	190–300	Partially carbonised
4.5		230–240	Partially carbonised
5.0	Dense black	300–400	Carbonised
5.5	Dark grey–black	310–420	Early volatilisation of carbon
6.0	Grey	350–435	Volatilisation of carbon
6.5	Grey-white	425–500	Late volatilisation of carbon
7.0	Opaque white	480–610	Carbon fully volatilised
7.5	Semi-translucent	> 530	Partially recrystallised
8.0	Transparent	> 600	Recrystallised

*Source:* Data from Helsen et al. (1995).

# Paleotemperature Indicators

## Conodont Alteration Index (CAI)

- The disadvantages of the **CAI** index are due to: (1) the broad  $T$  range associated to the index, (2) the large uncertainty ( $\pm 20$ -50 °C), since the colour is subjective, (3) the large number of pristine conodonts required for statistical accuracy (30-100).
- The computer analysis observed that the intensity of the RGB components decreases almost linearly with the increase of **CAI** up to 5.0, and increases rapidly in the range **CAI** 5.0-8.0.
- Nestell and Balsam (1996) found that the reflectance of white light off conodonts increases almost linearly with wavelength in the range of 550-800 nm and that the slope of the reflectance vs wavelength varies with CAI:

$$\text{CAI} = -95.818(\text{slope}) + 623.996(\text{slope})^2 + 4.558$$

Slope = %/nm

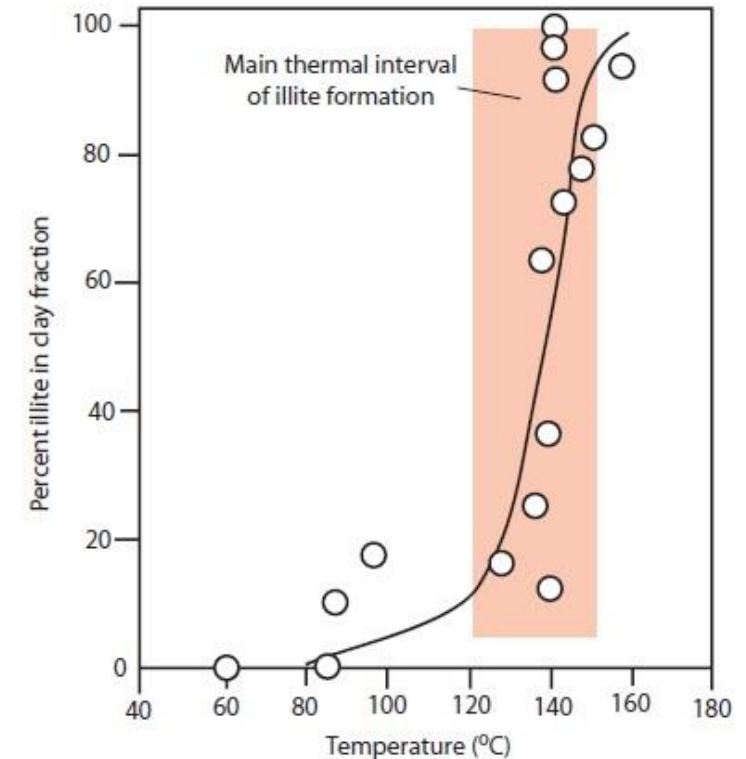
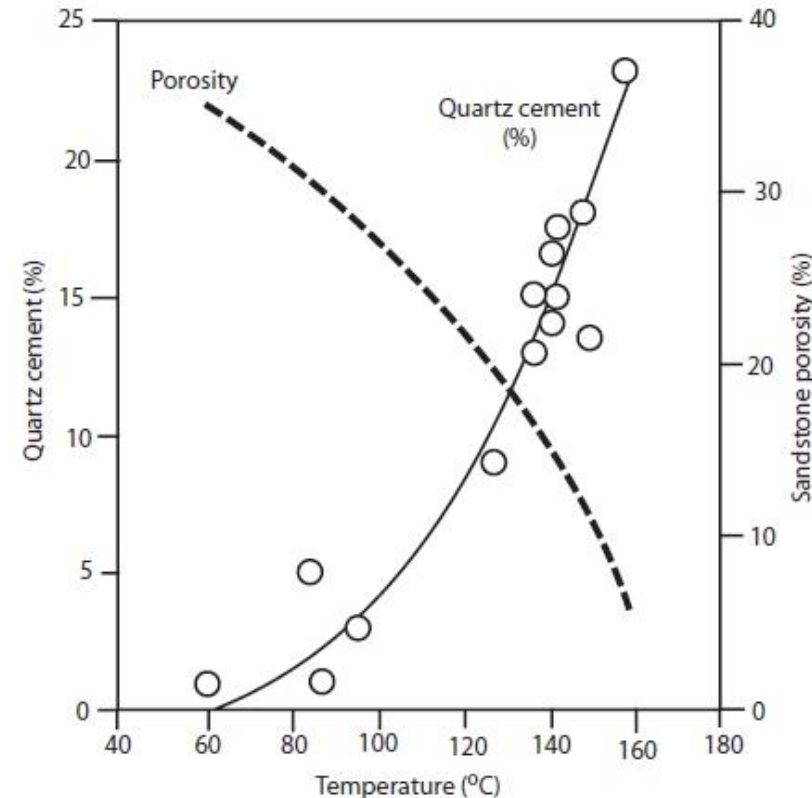
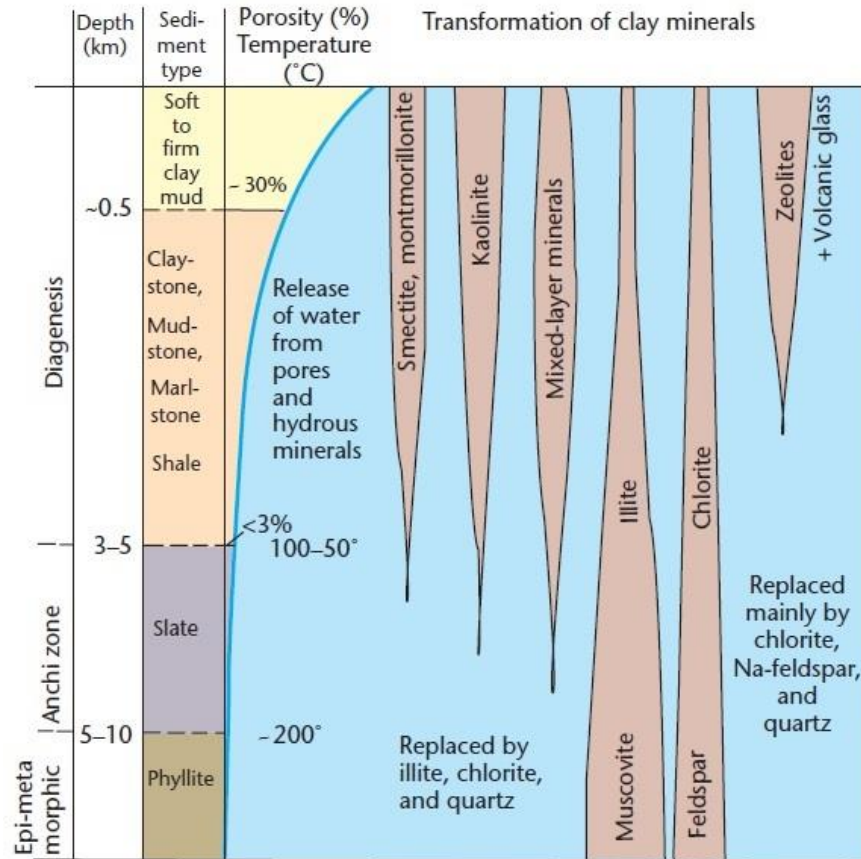
Uncertainty:  $\pm 0.2\%$  with 95% confidence



# Paleotemperature Indicators

## Mineralogical Indices

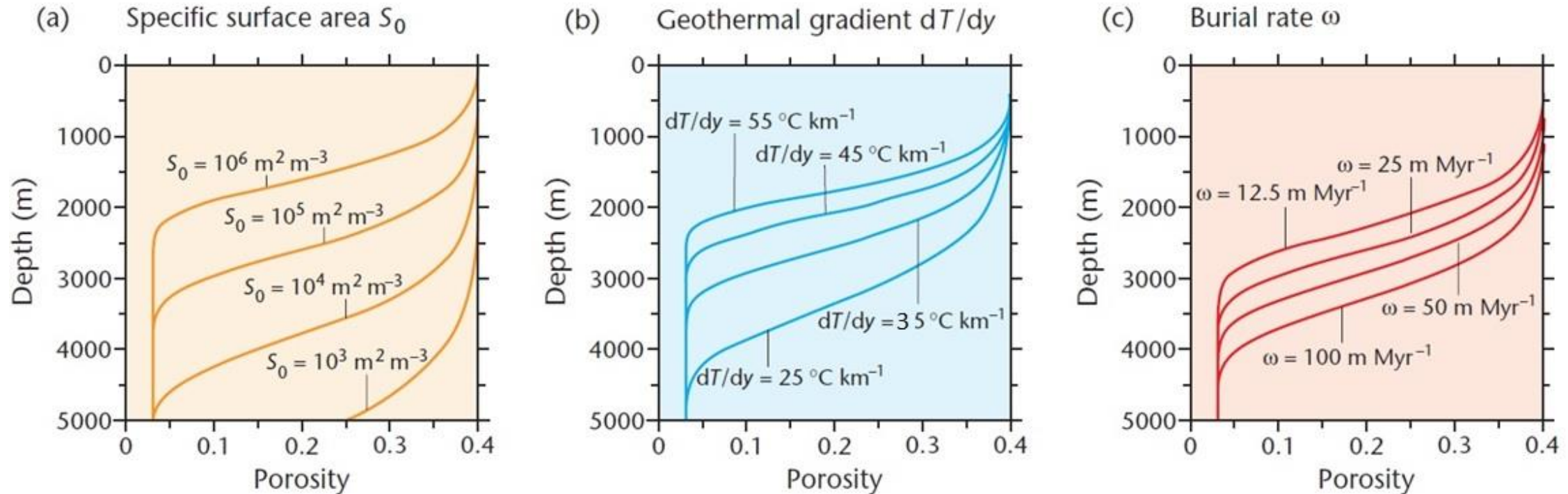
- The beginning of oil generation is correlated with the disappearance of smectite in favour of illite.
- Quartz cementation act in sandstones buried to  $T > 60^\circ\text{C}$  ( $90^\circ\text{--}120^\circ\text{C}$ ) and is mainly due to dissolution in pressure solution seams (stylolites), where silica solubility is enhanced by the presence of clay minerals and micas.





# Paleotemperature Indicators

## Mineralogical Indices



$S_0 = 10^3 \text{ m}^2 \text{ m}^{-3}$  corresponds to grain diameter = 1 mm

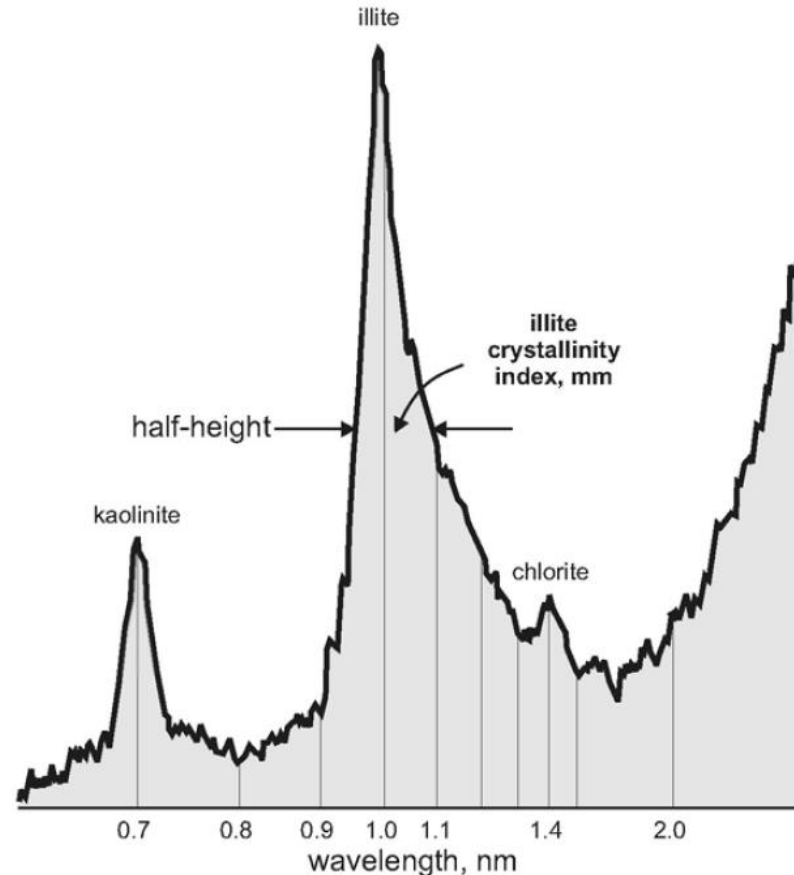
Porosity reduction is only due to cementation (no mechanical compaction is assumed)

- Porosity-depth curve assuming quartz cementation during a constant burial rate depends on (a) the initial specific surface  $S_0$  (a), (b) the geothermal gradient  $dT/dy$ , and (c) the deposition (burial) rate  $\omega$ .

# Paleotemperature Indicators

## Mineralogical Indices

- As diagenetic processes increase clay minerals (kaolinite, smectite, illite and chlorite) enrich in illite and chlorite at the expense of smectite (typically disappears at a depth of 1-2.5 km) and kaolinite.
- The abundance of illite in a material is measured using x-ray diffraction: illite shows on an x-ray spectrum as a characteristic peak at wavelength  $\sim 1$  nm. As  $T$  increases the *index of crystallinity*, defined as the width of the 1 nm illite peak (at half-height) decreases (progressive narrowing of the 1nm illite peak).
- This method works at high maturity level and identifies the transition between diagenesis and metamorphism occurs between indices 4.0-2.5, which is equivalent of vitrinite reflectance of  $R_{vmax}=2.3-3.3\%$ .



# Paleotemperature Indicators

## Mineralogical Indices

Other methods using clay minerals are based on the relative proportion of smectite with respect to illite with increasing time and  $T$ , considering that the ratio of the chemical activities potassium/sodium (K/Na) in the clay exerts some control on the reaction rate.

Pytte and Reynolds, 1989

$$-\partial S/\partial t = S^5 \times (\text{K/Na}) \times A \exp(-E/RT)$$

$S$  = mole fraction of smectite in illite–smectite mixed

$$(\text{K/Na}) = 74.2 \times \exp(-2490/T)$$

$$A = 5.2 \times 10^7 \text{ s}^{-1}$$

$$E = 138 \text{ kJ mol}^{-1}$$

$$R = \text{the gas constant} = 8.314472 \text{ J K}^{-1} \text{ mol}^{-1}$$

$T$  = temperature (K)

$$S^4 = \frac{S_0^4}{1 + (1.543 \times 10^{10}) \times t \times S_0^4 \exp(-19,110/T)}$$

$S_0$ =mole fraction of smectite in the initial mixed-layer clay

$t$ =time (s) spent by the clay at more than 90% of peak  $T$

# Paleotemperature Indicators

## Mineralogical Indices

Bethke and Altaner (1986) found that a smectite layer adjacent to a single illite layer converts to illite at lower energy than smectite layer totally surrounded by either smectite or illite.

They derived a weighted kinetic relationship for the reactions of the three different smectite types:

$$(\partial P_S / \partial t) = -P_S \times (X_0 k_0 + X_1 k_1 + X_2 k_2)$$

$P_S$  = fraction of smectite with respect to total smectite–illite

$X_i = N_i / N$  = fraction of smectite with  $i$  adjacent illite layers/total smectite

$N_i = N_{i0} \exp(-k_i t)$

$N_{i0}$  = initial fraction of Type  $i$  smectite/total smectite

$k_i = A \exp(-E_i / RT)$  = reaction rate for smectite with  $i$  adjacent illite layers

They assumed:  $A = 10^{-3} \text{ s}^{-1}$

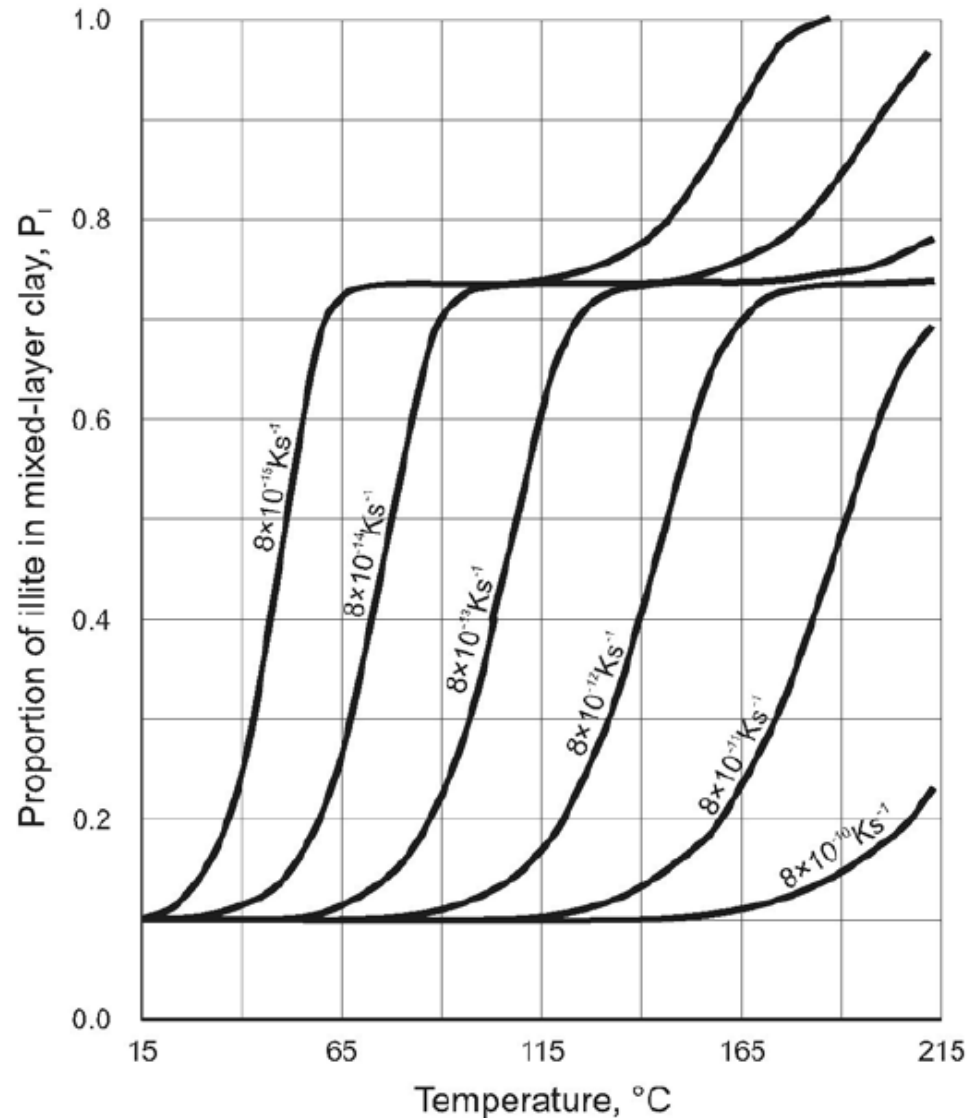
$E_0 = 77.5 \text{ kJmol}^{-1}$  (Type 0)

$E_1 = 75.5 \text{ kJmol}^{-1}$  (Type 1)

$E_2 = 100.5 \text{ kJmol}^{-1}$  (Type II)

# Paleotemperature Indicators

## Mineralogical Indices



**Question:** A particular clay sample is exhumed from a formation that was deposited 25 Myr ago. The present temperature of the formation is 80°C. Assuming an average surface temperature of 15°C, what is the expected illite content in relation to the total smectite-illite?

**Answer:** We need to determine the average heating rate:  
average heating rate = change in temperature / change in time

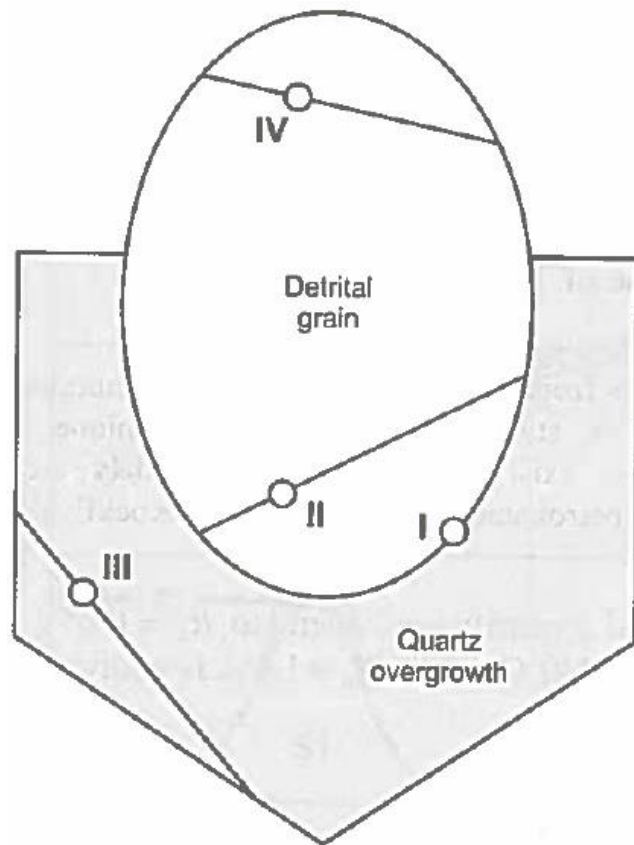
$$dT/dt = (80^\circ\text{C} - 15^\circ\text{C})/25 \text{ Ma} = 65 \text{ K}/(25 \times 3.1557 \times 10^{13} \text{ s}) = 8.24 \times 10^{-14} \text{ K s}^{-1}$$

Assuming a heating rate of  $8.24 \times 10^{-14} \text{ K s}^{-1}$  and a present  $T$  of 80°C:  
 $P_i \sim 0.55$

# Paleotemperature Indicators

## Fluid Inclusion Microthermometry (FIM)

- Fluid inclusions between crystals can be used to constrain the timing of oil migration in relation to mineral diagenesis and give some indications about  $P$ - $T$  conditions at the time of entrapment.
- The fluid (trapped as a single phase) separates into a liquid and vapour components upon reduction to surface conditions. Upon reheating the inclusion, the fluid reverts to a single phase at  $T_h$  (homogenization temperature), the minimum  $T_h$  recorded for a given population of fluid inclusion is likely the actual  $T$  at which the inclusion was originally trapped.



**I: Fluid trapped during crystallization of quartz overgrowth.**

**II: Fluid trapped during healing of fractures prior to quartz overgrowth.**

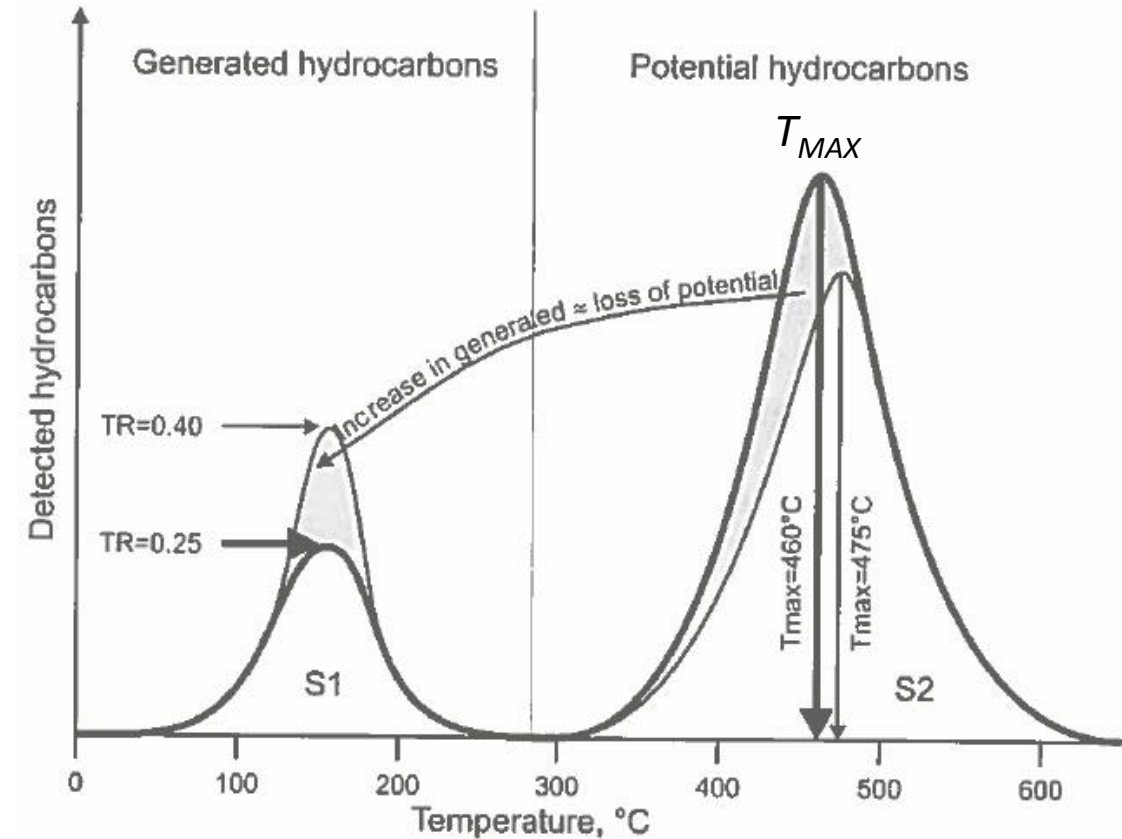
**III: Fluid trapped during healing of fractures during or after quartz overgrowth.**

**IV: It cannot be used for timing in detrital quartz.**

# Paleotemperature indicators

## Pyrolysis (Rock-Eval)

- **Pyrolysis** (annihilation by fire), in the petroleum industry, describes a procedure whereby a rock's sample is heated in an inert gas (e.g., argon).
- Heating rate is kept constant while hydrocarbons boiled from the rock are carried with the inert gas to a chromatograph, which records the volume of hydrocarbons through time.
- The hydrocarbons present in the pores of the rock sample are expelled and detected by the pyrolysis instrument at  $T < 200\text{ }^{\circ}\text{C}$ . Then, a distinct peak of detected hydrocarbons occurs at  $100\text{--}200\text{ }^{\circ}\text{C}$  (S1).
- The kerogen material left in the sample reaches maturation if the  $T$  is raised at sufficient level (reactions generating hydrocarbons occur in a hour) and thus a second distinct peak of detected hydrocarbons occurs at  $400\text{--}500\text{ }^{\circ}\text{C}$  (S2).
- The higher is the thermal maturity, the lower the volume of potential hydrocarbons and the higher is the  $T$  required to mobilize it during pyrolysis.



$$\text{Transformation rate (TR)} = S1 / (S1 + S2)$$

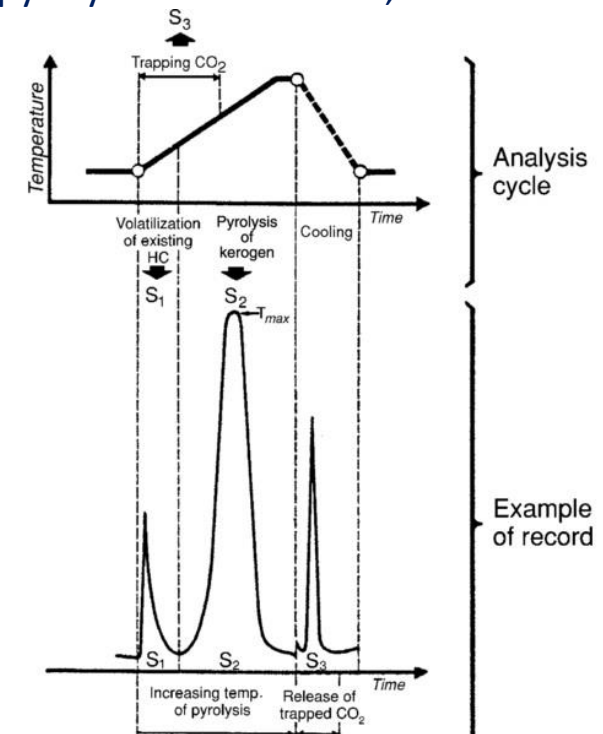
A higher heating rate results in a higher value of  $T_{MAX}$



# Paleotemperature Indicators

## Pyrolysis (Rock-Eval)

- Pyrolysis method is based on determination and analysis of a number of parameters such as the Total Organic Carbon (**TOC**), **S<sub>1</sub>**, **S<sub>2</sub>**, **S<sub>3</sub>**, **T<sub>max</sub>** range, hydrogen index (**HI**), oxygen index (**OI**), production index (**PI**), pyrolyzable carbon (**PC**).
- S<sub>1</sub>** = amount of free hydrocarbons in the sample (in milligrams of hydrocarbon per gram of rock) volatilized at a temperature of <300 °C. **S<sub>1</sub>** normally increases with depth
- S<sub>2</sub>** = amount of hydrocarbons generated through thermal cracking of nonvolatile organic matter by heating in temperature intervals from 300 to 550 °C (~650 °C at the start and upwards to 800 /850 °C). **S<sub>2</sub>** normally decreases with burial depths >1 km.
- S<sub>3</sub>** = amount of CO<sub>2</sub> (in milligrams CO<sub>2</sub> per gram of rock) produced during pyrolysis of kerogen within the 300–400 °C range.
- T<sub>max</sub>** = T at which the maximum release of hydrocarbons from cracking of kerogen occurs during pyrolysis (top of **S<sub>2</sub>** peak). **T<sub>max</sub>** = 400–430 °C represents immature organic matter; **T<sub>max</sub>** = 435–450 °C indicates a mature or oil zone; **T<sub>max</sub>** > 450 °C represents an over-mature zone.
- TOC** (expressed as weight percentage) = weight percentage of carbon in CO<sub>2</sub> formed during pyrolysis below 400 °C, and in CO formed below 500 °C, and in the hydrocarbon peaks, **S<sub>1</sub>** and **S<sub>2</sub>**;
- HI** =  $(100 \times S_2) / \text{TOC}$ ;
- OI** =  $(100 \times S_3) / \text{TOC}$ ;
- PI** =  $S_1 / (S_1 + S_2)$ ;
- PC** =  $0.083 \times (S_1 + S_2)$ .





# Paleotemperature Indicators

## Fission Track Thermochronology (FFT)

- **Fission track analysis** is based on the observation that the crystal lattices of some minerals preserve the scars of spontaneous nuclear fission event ( $^{238}\text{U}$ ). Apatite, zircon and sphene contain uranium and are commonly used for this analysis.
- The initial length of a fission track is relatively constant for a given mineral ( $\sim 16\text{ }\mu\text{m}$ ) for apatite, but gradually anneals starting at each end of the fission track and moving inwards, decreasing the track length at a rate dependent on  $T$ , mineral composition and the angle between the fission track and the c-axis of the crystal.
- Temperature of annealing is different for each mineral (e.g., apatite starts to anneal at  $T > 50^\circ\text{C}$ ). Zircons retain tracks for much longer at higher  $T$  than sphene and apatite.

**Temperature Required to Anneal Fission Tracks in 1 hour at 1 atmosphere Pressure**

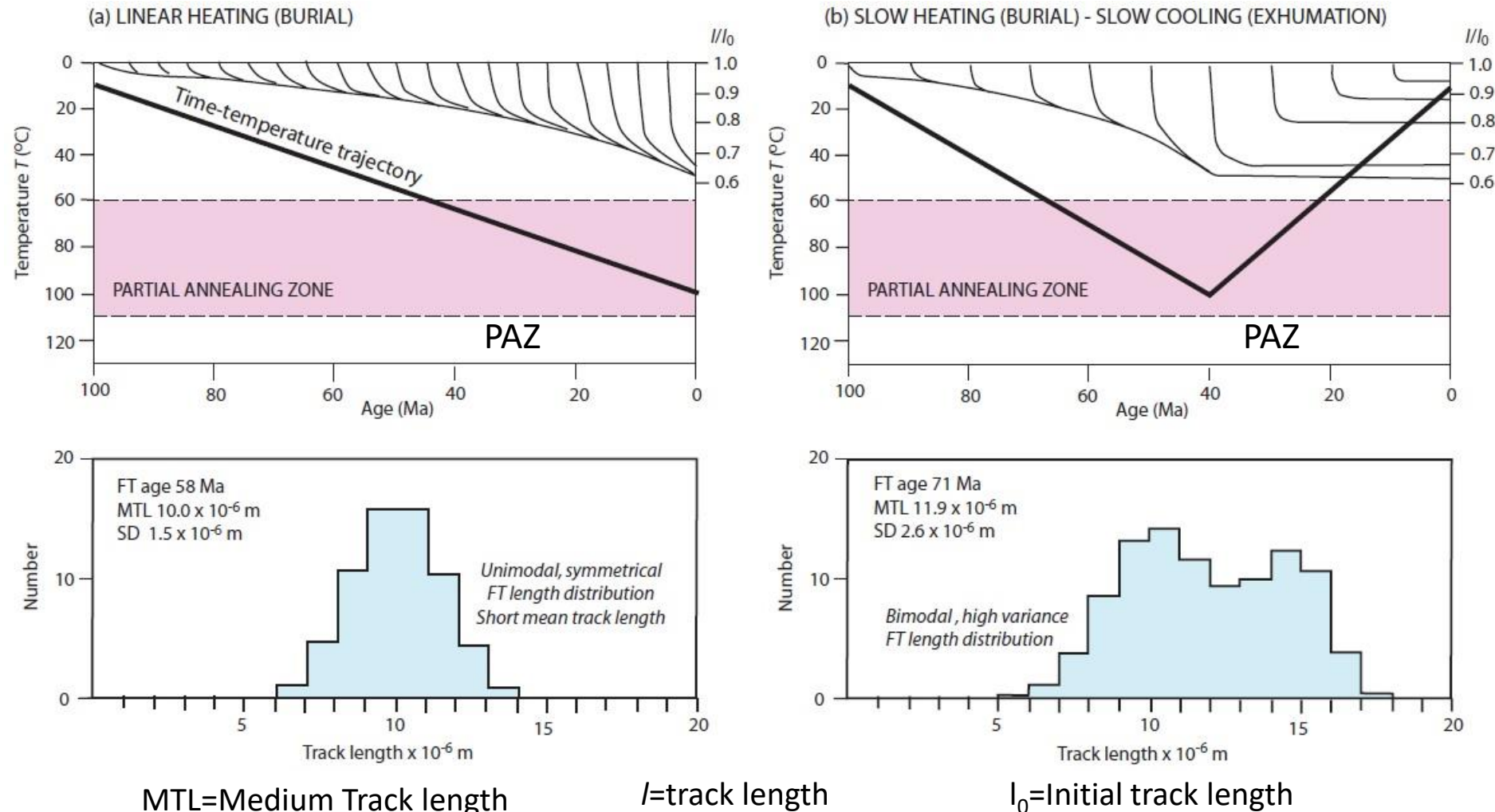
Mineral	Temperature Required to Anneal Indicated Proportion of Tracks		
	0%	50%	100%
Apatite <sup>a</sup>	275–400°C	322°C	375–530°C
Sphene	520°C	620°C	637°C
Zircon		700°C	

<sup>a</sup>Apatite data derived from more than one primary source *Source:* Data from Fleischer et al. (1975).

# Paleotemperature Indicators

## Fission Track Thermochronology (FFT)

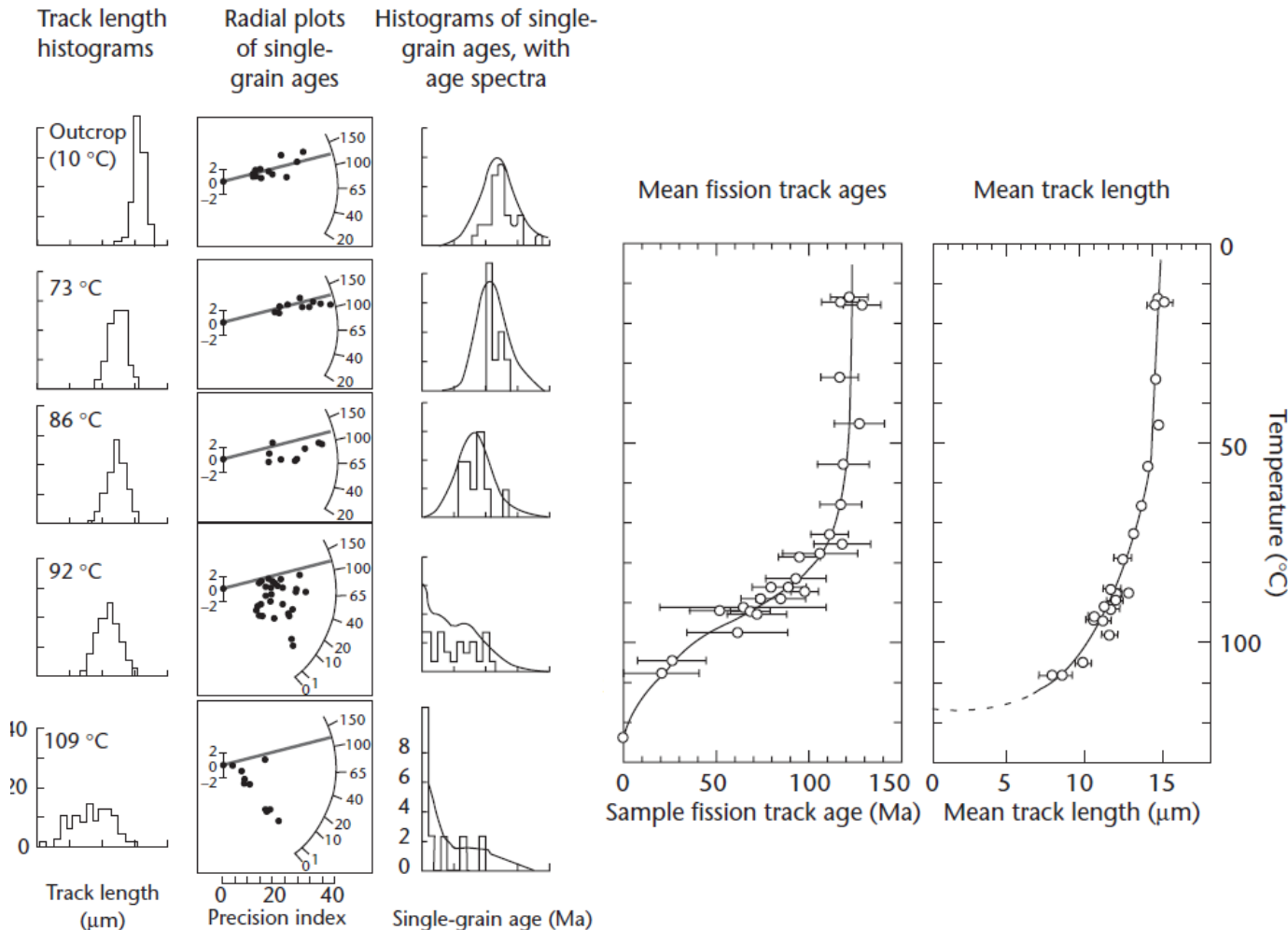
- Above the PAZ, the fission track lengths are tightly clustered and long ( $14\mu\text{m}$ ), indicating minimal annealing.
- Newer tracks experience different maximum  $T$  during the cooling phase (sedimentary basin inversion), causing a bimodal histogram, with a high STD.



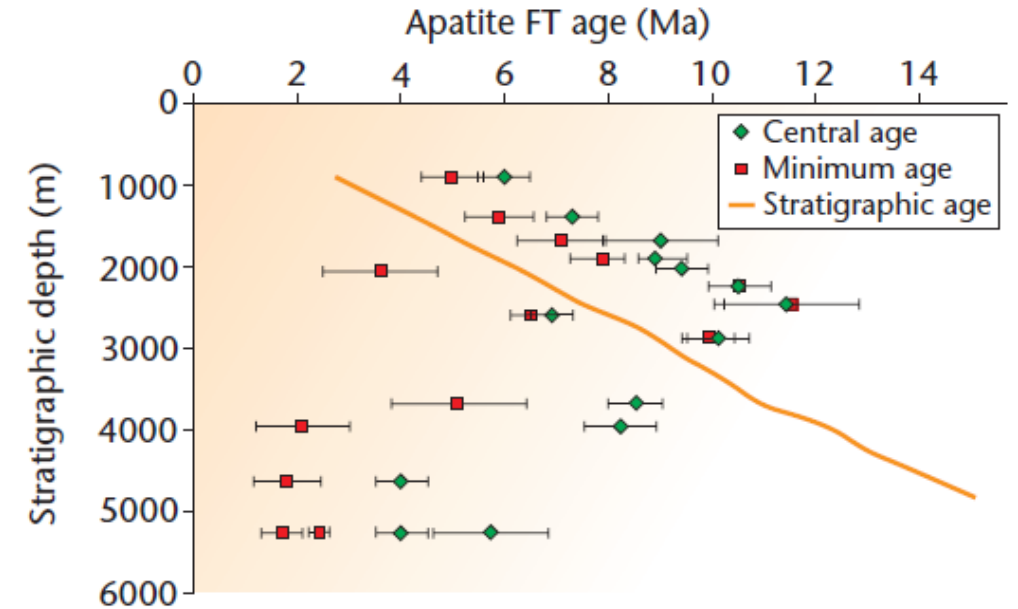
# Paleotemperature Indicators

## Fission Track Thermochronology (FFT)

- With greater depth, fission track length histograms show a wider distribution and a shorter MTL.
- The fission track ages decrease due to the greater amount of annealing at elevated  $T$ , reaching 0 Myr at  $T$  of 120°C.
- Annealing during burial (e.g., at a depth > 2.5 km) causes a decrease of the age of samples (< of the stratigraphic age).
- Samples above the line of stratigraphic age indicate rapid exhumation, transport, and burial without annealing.

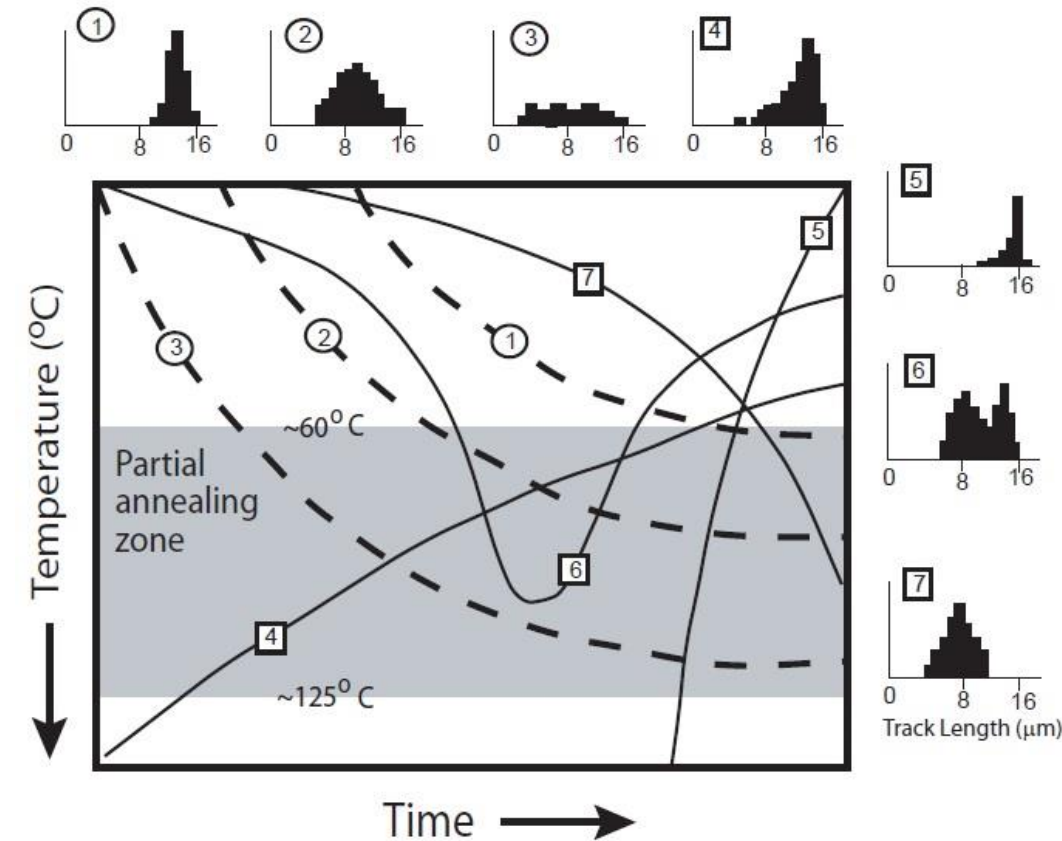


### Western Nepal



# Paleotemperature Indicators

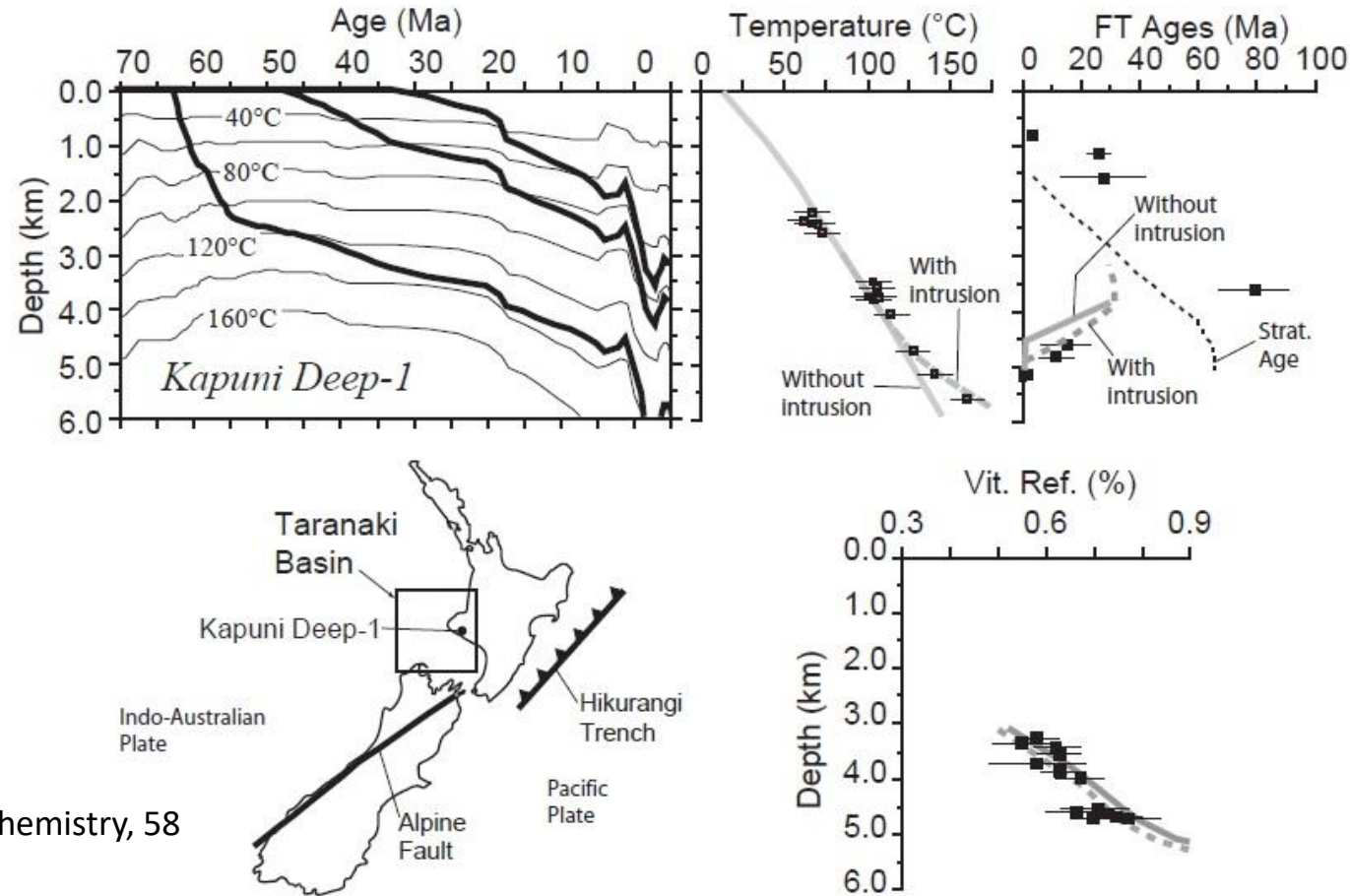
## Fission Track Thermochronology (FFT)



- Paths 1–3 show progressive burial to maximum depths below the top of the PAZ followed by near isothermal conditions: long track lengths with narrow distributions indicative of that sample's grains not being annealed after deposition. With greater depth (and temperature) the average track lengths progressively get shorter and the distribution of tracks spreads out reflecting the partial annealing.
- Paths 4 and 5 show cooling from depths within or below the PAZ. With cooling from temperatures above the base of the PAZ for this apatite species, average track lengths tend to be long but with negative skewness and the distribution gets narrower with more rapid cooling through the PAZ—compare path 4 with 5.
- Path 6 shows heating into the partial annealing zone followed by cooling out of it. For a sample that is heated to PAZ temperatures and then cooled, a bimodal distribution can be expected because tracks that were shortened while the sample was in the PAZ are mixed with longer tracks that formed after cooling out of the PAZ.
- Path 7 shows rapid heating to temperatures in the partial annealing zone. In this case average track lengths will be relatively short but not as spread out as in the case of longer periods of time in the PAZ (paths 2 and 3).

# Paleotemperature Indicators

## Fission Track Thermochronology (FFT)



Armstrong, 2005, Rev. Min. & Geochemistry, 58

Burial/thermal history and output parameters temperature, apatite fission-track, and vitrinite reflectance data for well Kapuni Deep-1 in Taranaki Basin.

Solid curves in the temperature, fission track, and vitrinite reflectance plots are model values for case of tracking thermal history with burial and erosion only. Thick dashed curves are for case of including sill intrusion below the sediments in the last <1 M.y.

- Vitrinite reflectance data show essentially the same fit with both the intrusion and non-intrusion model, indicating that the vitrinite reflectance data are less sensitive to the short-term transient thermal effects than are the AFT data.

# Paleotemperature Indicators

## Fission Track Thermochronology (FFT)

- Annealing results in shorter spontaneous tracks, reduced spontaneous track density and a reduction in the apparent fission track age ( $t_a$ ).
- Above a critical  $T$ , fission tracks anneal instantaneously and  $t_a$  is reduced to zero, resetting the fission track clock.

$$t_a = \frac{1}{\psi_D} \ln \left[ 1 + \frac{\psi_D \phi \sigma I \rho_s}{\psi_f \rho_i} \right]$$

$\psi_D$  = total decay constant of  $^{238}\text{U} = 1.55125 \times 10^{-4} \text{ (Ma}^{-1}\text{)}$

$\psi_f$  = spontaneous fission decay constant of  $^{238}\text{U} \text{ (Ma}^{-1}\text{)}$

$I$  = isotopic ratio of  $^{235}\text{U}/^{238}\text{U} = 7.2527 \times 10^{-3}$

$\sigma$  = thermal neutron fission cross section for  $^{235}\text{U} = 580.2 \times 10^{-24} \text{ (cm}^2\text{)}$

$\phi$  = thermal neutron fluence ( $\text{cm}^{-2}$ )

$\rho_s/\rho_i$  = spontaneous/induced track density ratio

A glass standard of known uranium concentration is included with each batch of apatite grain mounts sent for irradiation and is mounted with a muscovite detector.

$$\Phi = B \times \rho_D$$

$\rho_D$  = density of fission tracks revealed on the muscovite detector

$B$  = calibration factor characteristic of the glass

$$t_a = \frac{1}{\psi_D} \ln \left[ 1 + \frac{\rho_s \rho_D}{\rho_i} \zeta \psi_D \right] \quad \zeta = \text{calibration constant}$$



## Paleotemperature Indicators

### Fission Track Thermochronology (FFT)

**Question:** The fission track laboratory at Latrobe University, Australia, has calibrated its technique with a value  $\zeta = 352.7 \pm 3 \text{ yr cm}^{-2}$  (e.g. Beardsmore and O'Sullivan, 1995). The calibration is only valid when the correct units are used for  $\rho$  ( $\times 10^6 \text{ cm}^{-2}$ ) and  $\Psi_D$  ( $\text{Ma}^{-1}$ ). The laboratory examined a population of apatite grains separated from drill cuttings taken from a well in the Browse Basin, Western Australia. The grains revealed fission track densities  $\rho_D = 1.378 \times 10^6 \text{ cm}^{-2}$ ,  $\rho_s = 6.294 \times 10^5 \text{ cm}^{-2}$  and  $\rho_i = 3.618 \times 10^6 \text{ cm}^{-2}$ . What is the apparent fission track age,  $t_a$ , for the sample?

**Answer:** In terms of the correct units for the calibration:

$$\rho_D = 1.378$$

$$\rho_s/\rho_i = 0.5 \times 0.6294/3.618 = 0.08698 \quad (\text{see Note above for explanation of the 0.5 factor})$$

$$\Psi_D = 1.55125 \times 10^{-4}$$

From Equation (5.32):

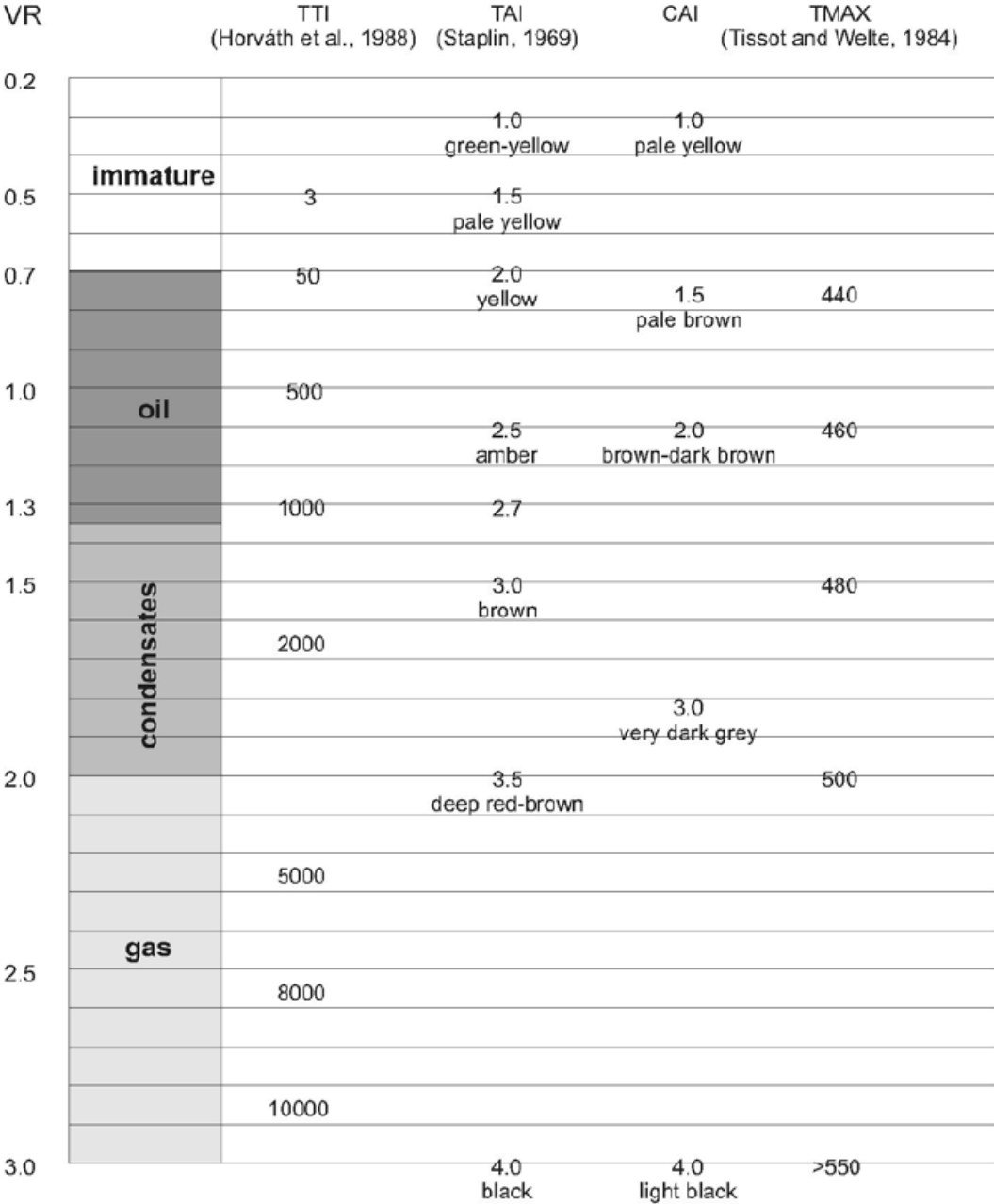
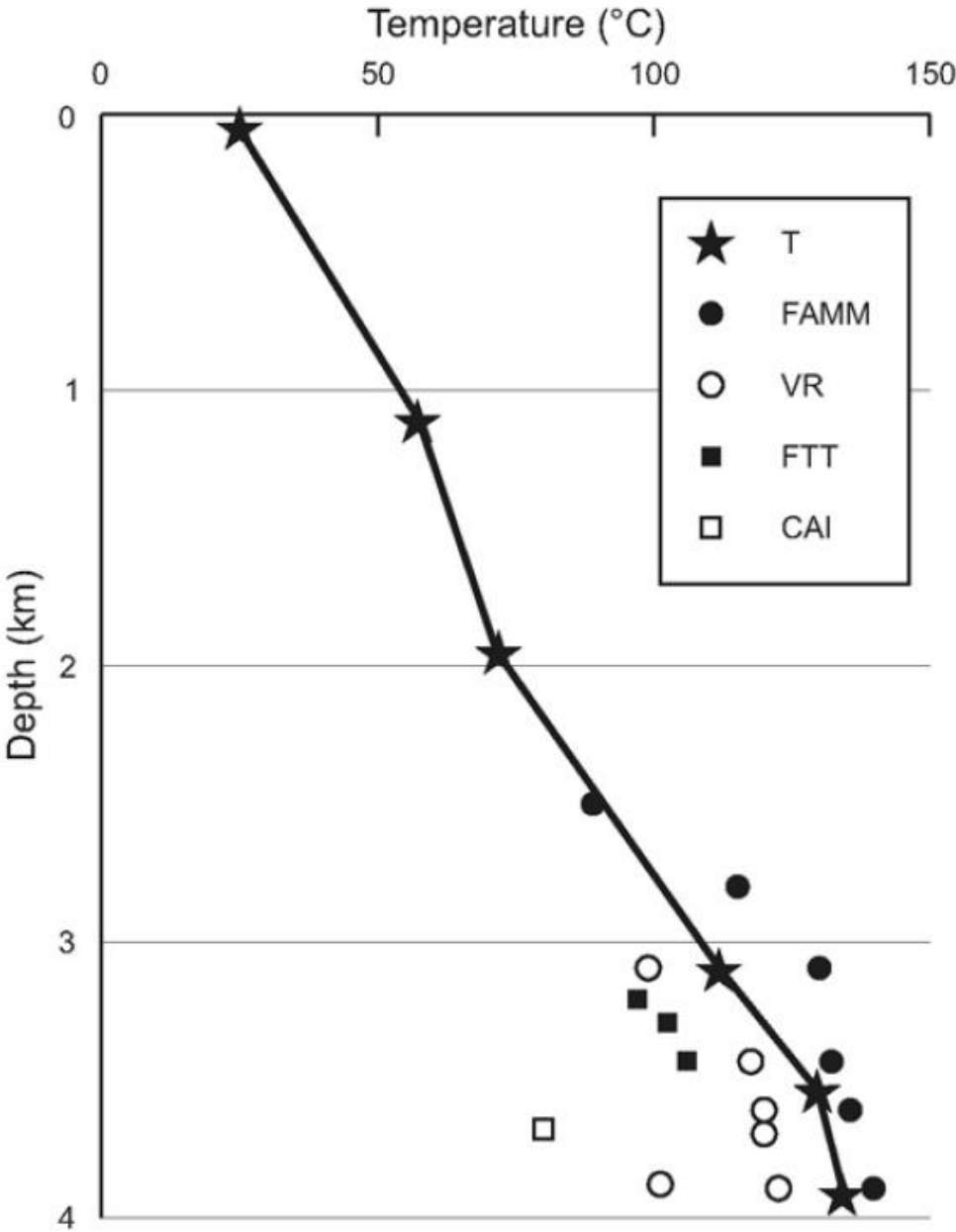
$$t_a = 1/(1.55125 \times 10^{-4}) \times \ln[1 + 0.08698 \times 1.378 \times 352.7 \times 1.55125 \times 10^{-4}] = 6446.4 \ln[1 + 0.00656] = 42.1 \text{ Ma}$$

# Thermal maturity Indicators

	Sediment component analysed	Main rock type	Equipment required	Approximate maturity range (VR <sub>max</sub> )	Palaeotemperature precision	Maximum measurable palaeotemperature (from Equation 5.9)	Time information	Limitations
VR	vitritinite	siltstone, shale	reflecting light microscope, photometer	0.3–5.0	±5°C	>350°C	none	reflectance suppression in marine deposits
FAMM®	various macerals		laser fluorescence microprobe	0.4–1.2		175°C		small number of laboratories set up for technique
TAI	palymorphs		transmitting light microscope, non-colourblind operator	0.3–2.4	±20°C or higher	260°C		subjective, poor temperature resolution
CAI	conodonts	carbonates		0.3–5.0+	±20–50°C	>600°C		subjective, Cambrian-Triassic only, poor resolution
Illite crystallinity	illite	shale		0.4–5.5	±10°C	300°C		clay composition changes due to fluids
Pyrolysis	organic matter	all sediments	GC-MS <sup>a</sup> with pyrolysis inlet, or RockEval tool	0.6–1.4	±5°C	200°C		must use standard heating rate, results vary for sediment type
Fluid inclusions	fluid inclusions in calcite, f'spar, quartz grains	sandstone, limestone	microscope, heating stage	n/a <sup>b</sup>	±2°C <sup>b</sup>	>1000°C	relative timing of different inclusions	only gives temperature at times of fluid migration
Biomarkers	extracts and pyrolysates	all sediment	GC-MS <sup>a</sup>	0.3–2.0	±15°C	240°C	none	most indices are poorly calibrated to VR
FTT	apatite	sandstone	microscope, thermo-nuclear reactor	0.3–1.0	±5°C	130°C	absolute age of last cooling event	composition of apatite must be determined



# Paleo-temperature Indicators



# References

## Main Readings:

### Books:

- Beardsmore and Cull, 2001, Crustal Heat Flow A Guide to Measurement and Modelling, Chapter 5: Thermal Maturity, 146-200.
- Allen and Allen, Basin Analysis, Chapter 10: Thermal history, 343-367.
- Eppelbaum Interpretation of Thermal Measurements, 393-475.

### Further readings:

Armstrong, 2005, Thermochronometers in Sedimentary Basins, Reviews in Mineralogy & Geochemistry, 58, 1-29.