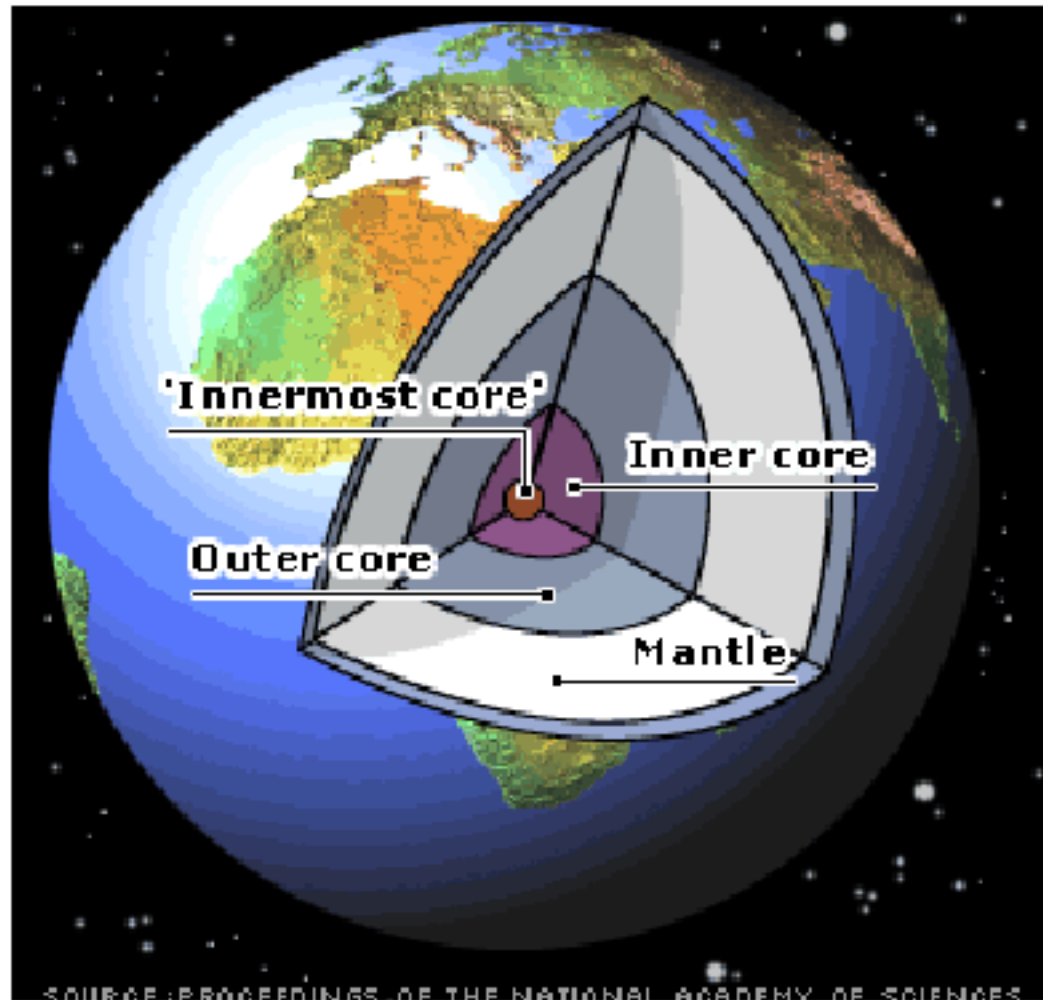


Element to Rock

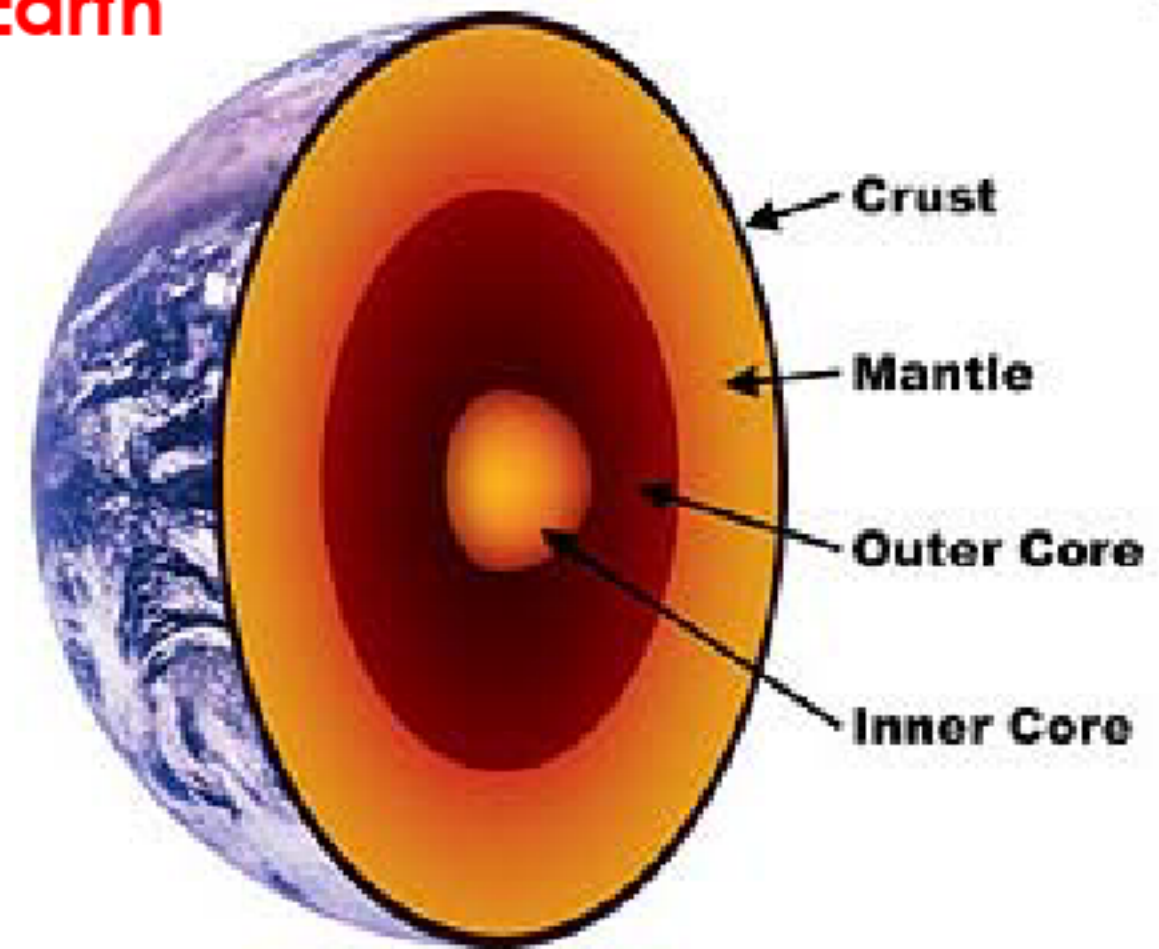
Geology 1

G. Bertotti





The Earth

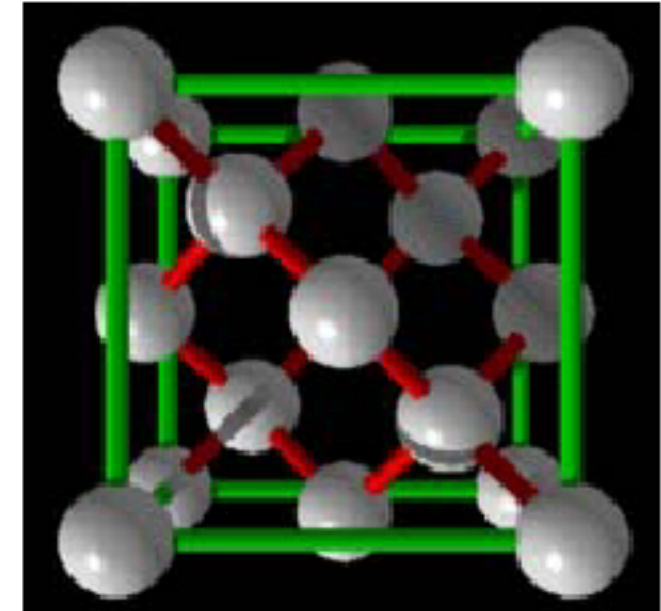


- very little information on elements and minerals (will come in 3rd part, M. van Tooren)
- minerals and rocks forming the Earth (particularly its outer part)

The basic components: **elements**, **minerals** and **rocks**

Elements

the most fundamental substances into which matter can be separated by chemical means



Minerals

Naturally formed, inorganic, solid material with a specific chemical composition and a characteristic crystalline structure.

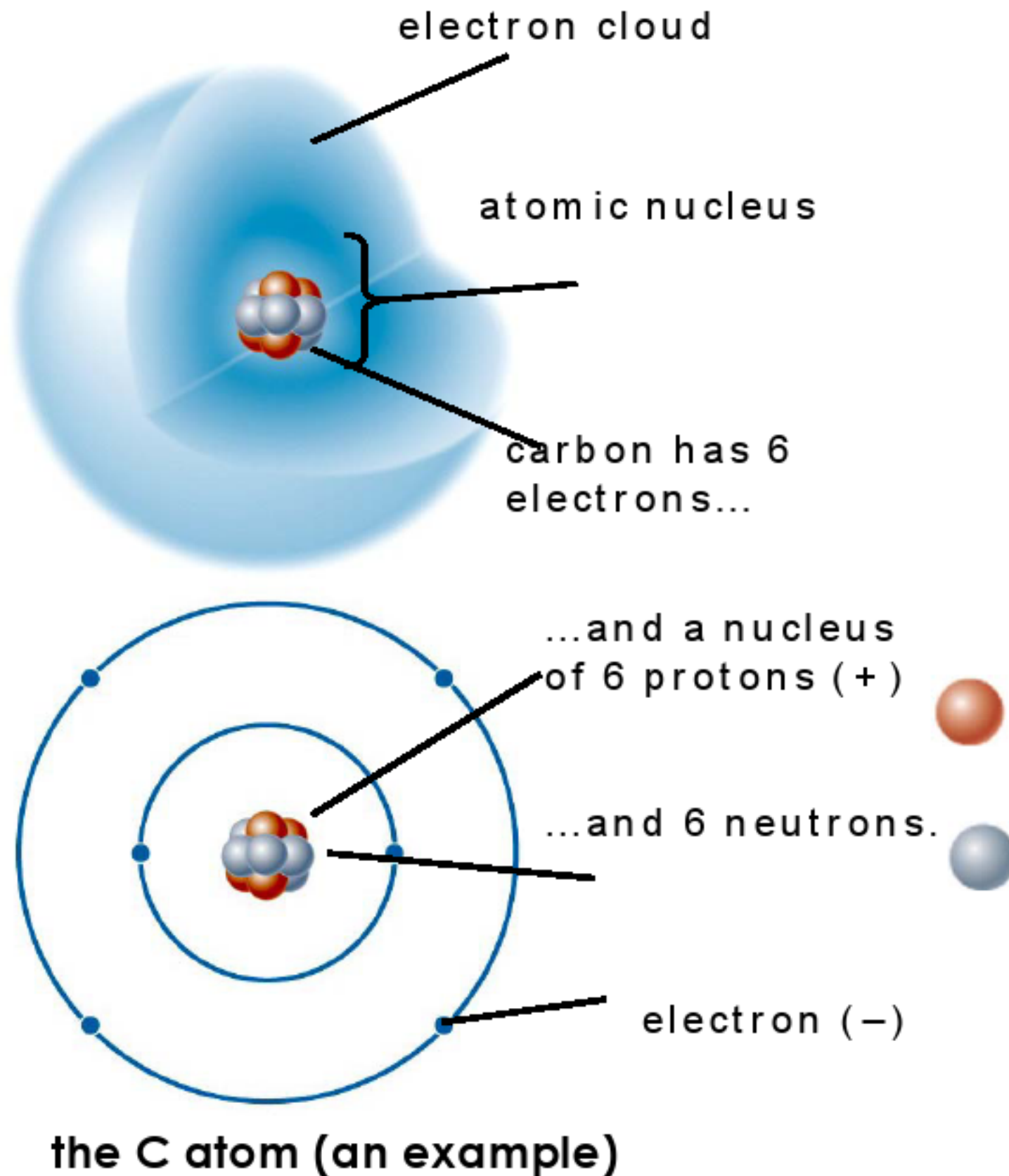


Rocks

Naturally formed, coherent mass of one or more minerals, sometimes including organic debris.



The atom, where everything starts



Atomic number: the number of protons in the nucleus

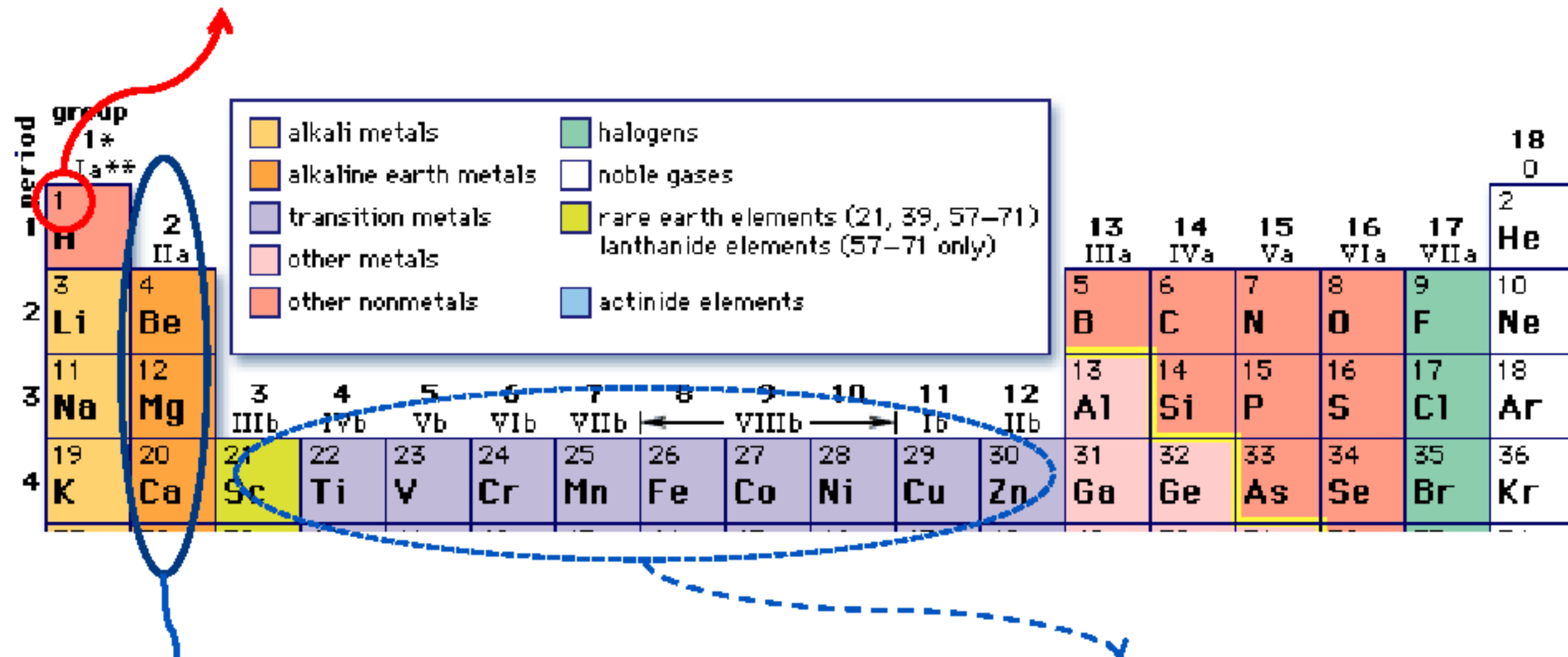
Atomic mass: mass of the neutrons + mass of the protons.

Electrons organized in **shells**.
The electrons in the outer shell give the **valence** which controls interactions

Atoms with the same number of protons but different number of neutrons are called **isotopes**.

The table of elements

Atomic number
(number of protons)



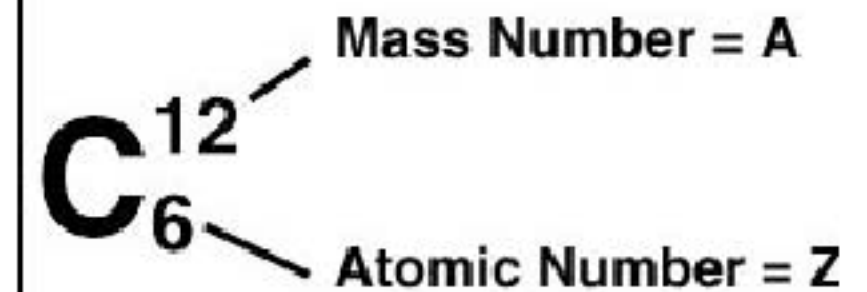
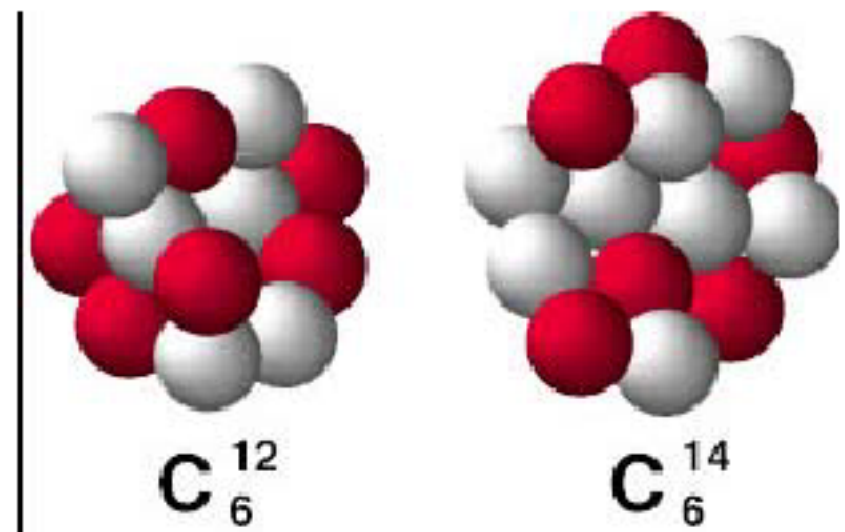
have the same number of electrons in the outside shell

These are exceptions as they have 1-2 electrons in the outer shell (they fill internal shells before the outer one is full)

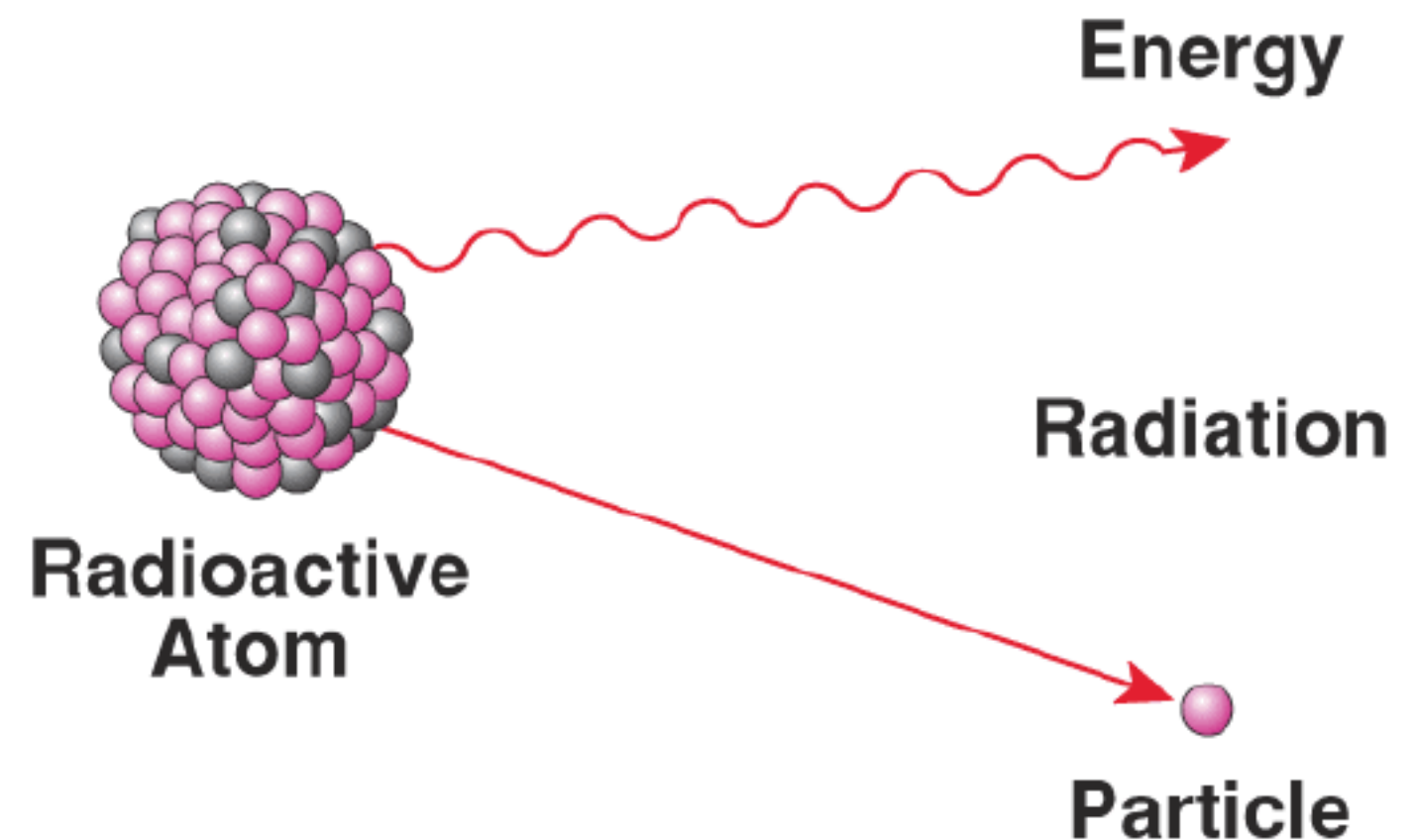
Isotopes

Atoms of the same elements always have the same number of protons but can have different number of **neutrons** leading to isotopes which are characterized by their different **mass number**

Carbon has three **stable** isotopes, C^{12} , C^{13} and C^{14}

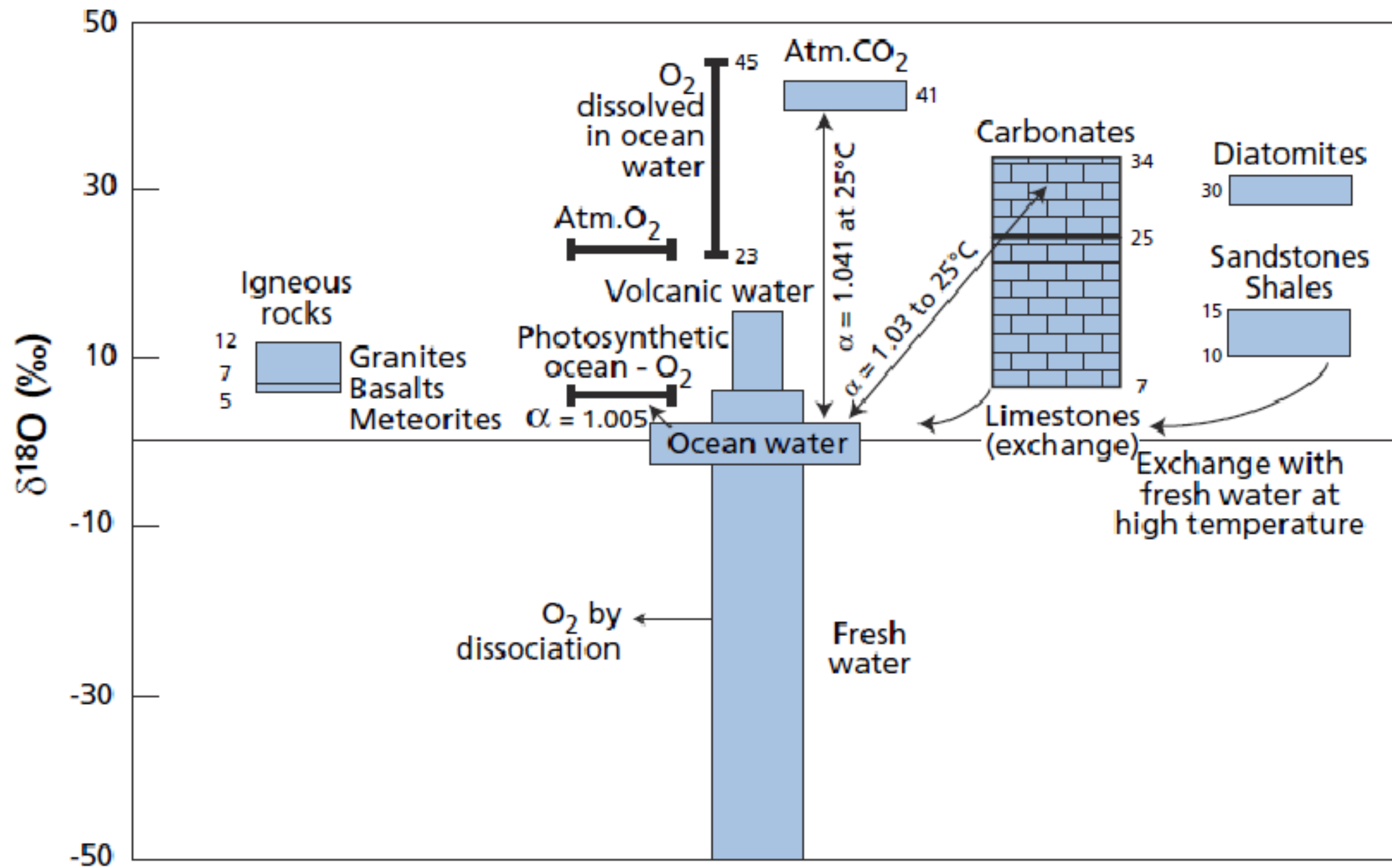


Some isotopes are **unstable** and decay at variable rates radiating energy and particles

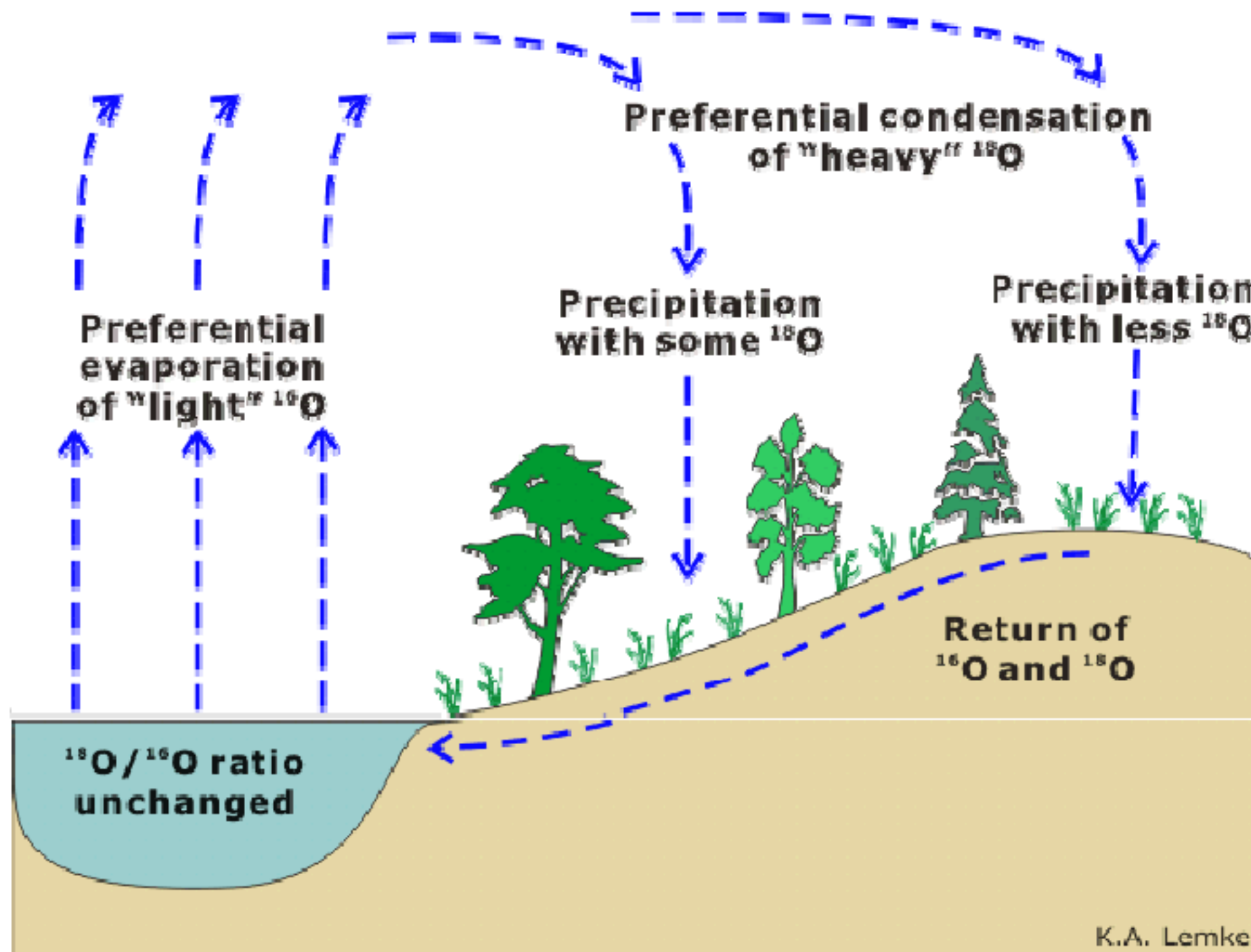


Because of their slightly different mass, isotopes have slightly different behaviour and can **fractionate**. Isotopes are fundamental tracers of numerous natural processes

The example of oxygen isotopes: large variations!



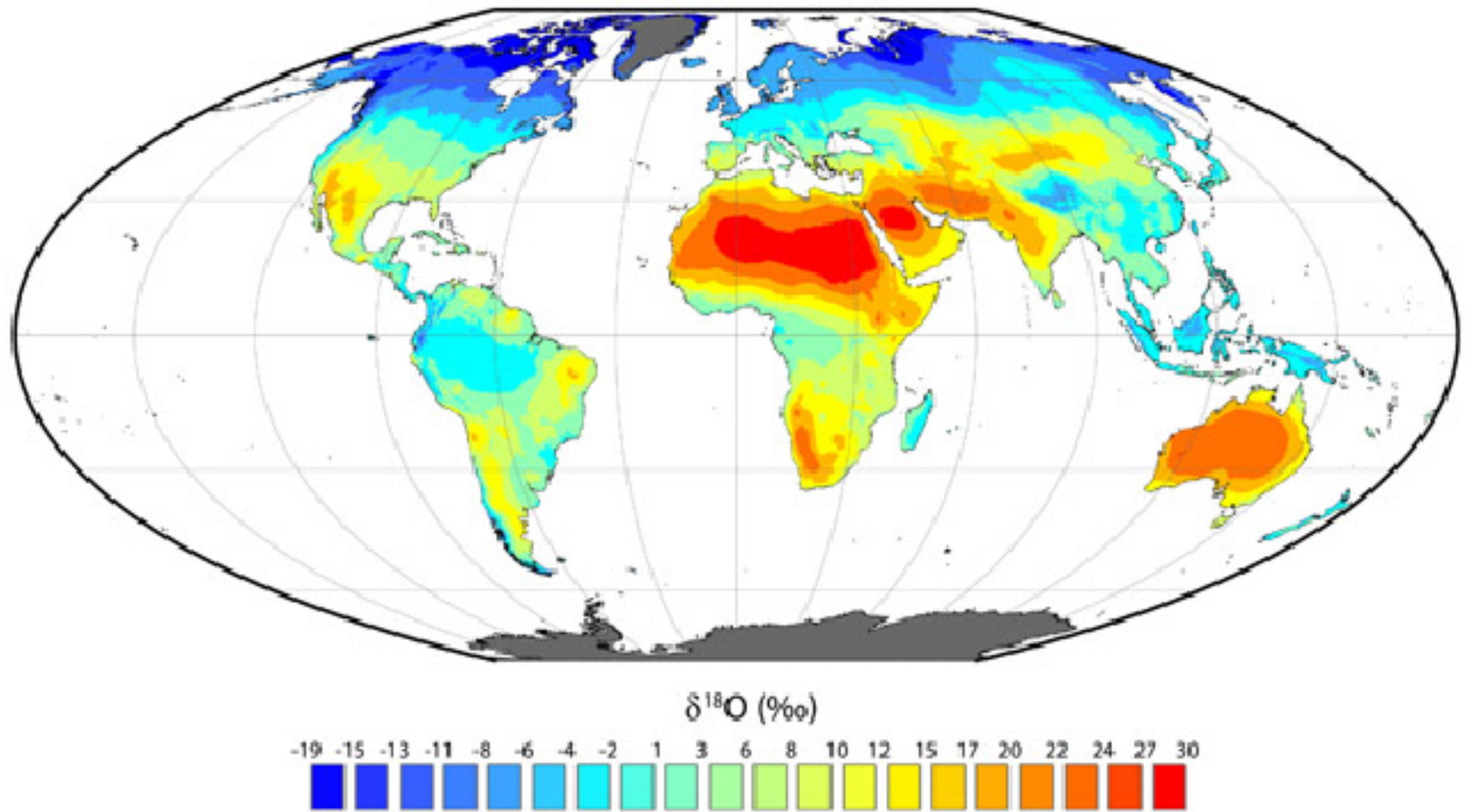
A typical agent of fractionation: **evaporation** and **precipitation**



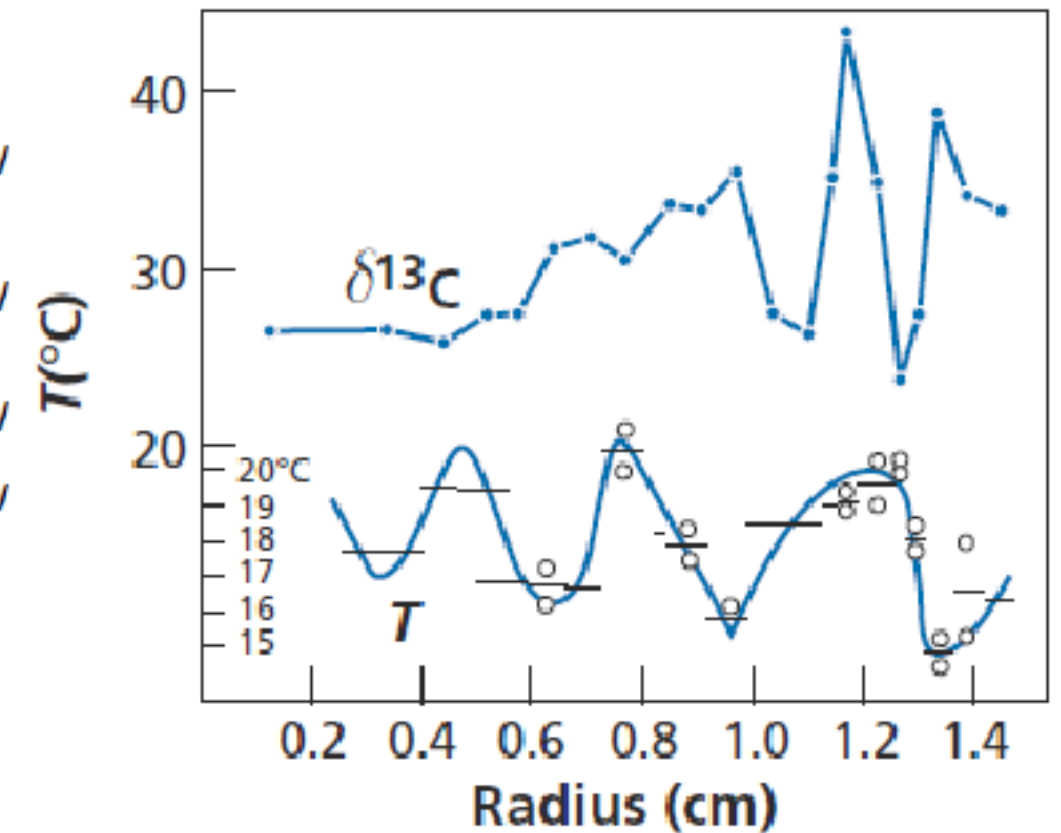
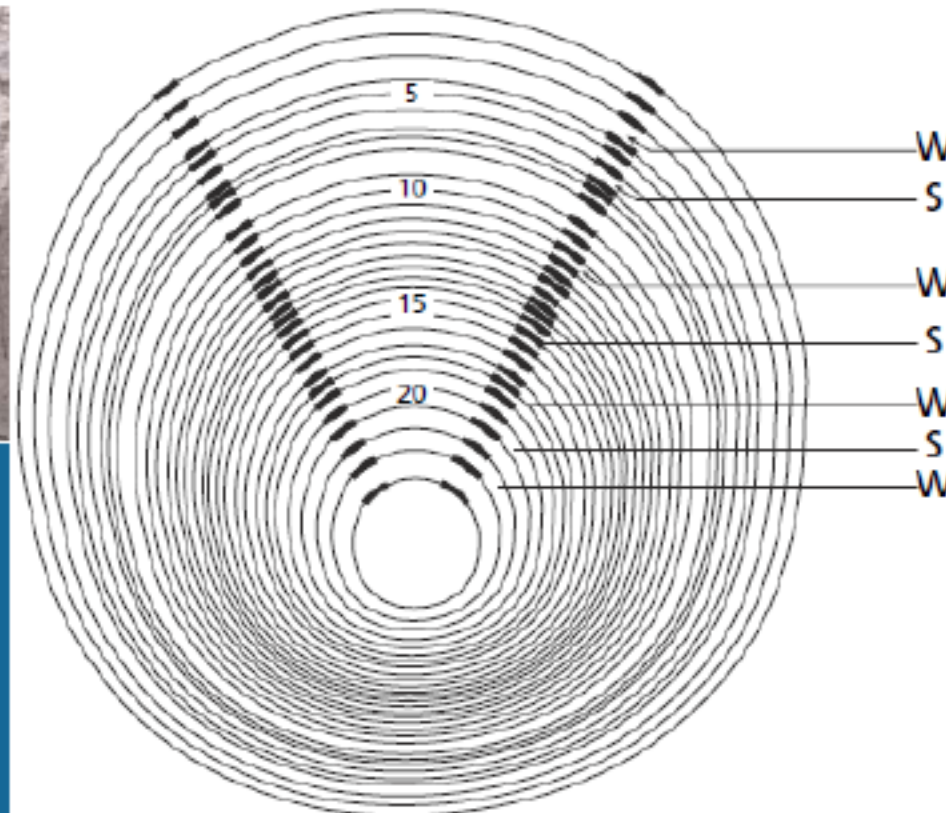
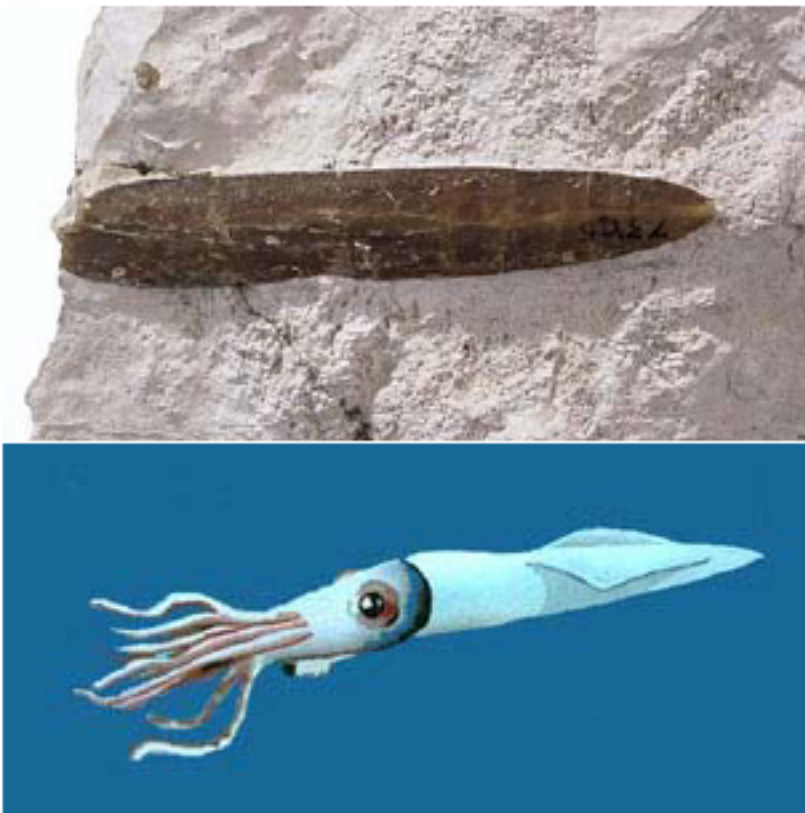
The isotope ratios depend on **temperature!**

Isotopes become tracers and geo-thermometer

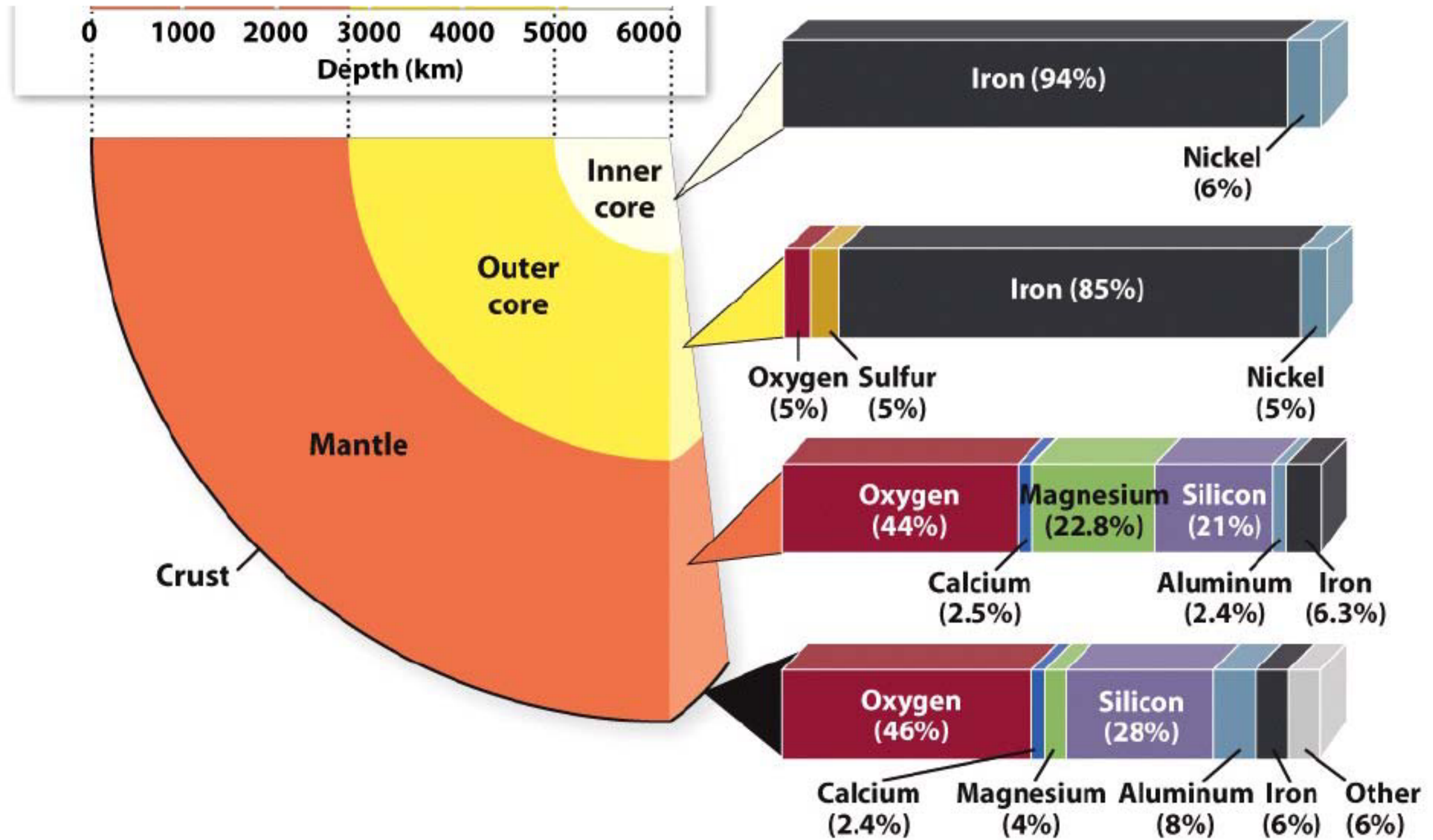
Rain in different parts of the world have different isotopic ratios



Temperatures 180 Million years ago



Distribution of elements in the Earth



Apparently large changes (we will come back to this)

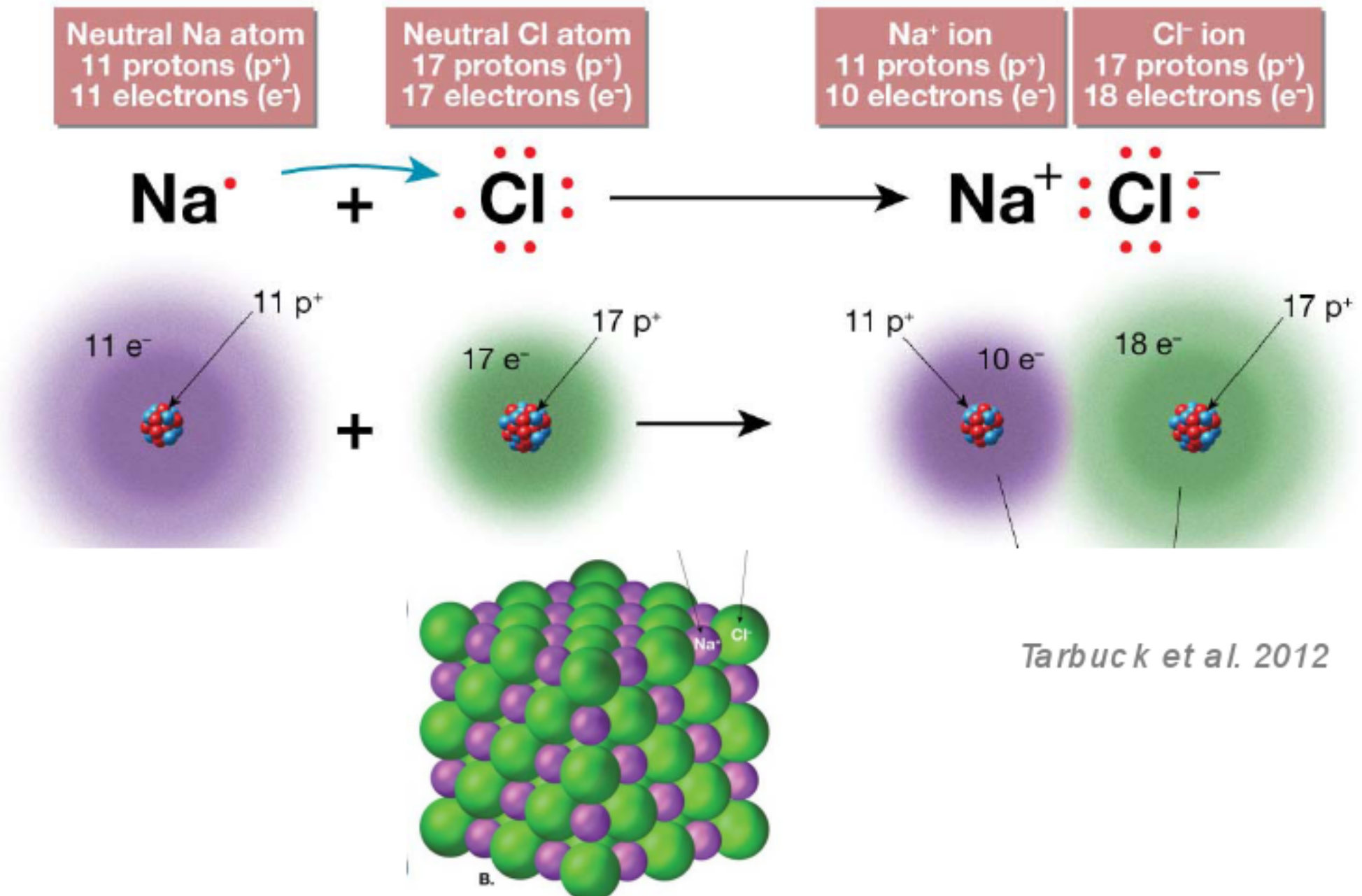
Chemical reactions

With the exception of noble gases, elements combine generally following the **octet rule**:

Atoms tend to gain loose or share electrons until they are surrounded by eight valence electrons

Ionic bonds

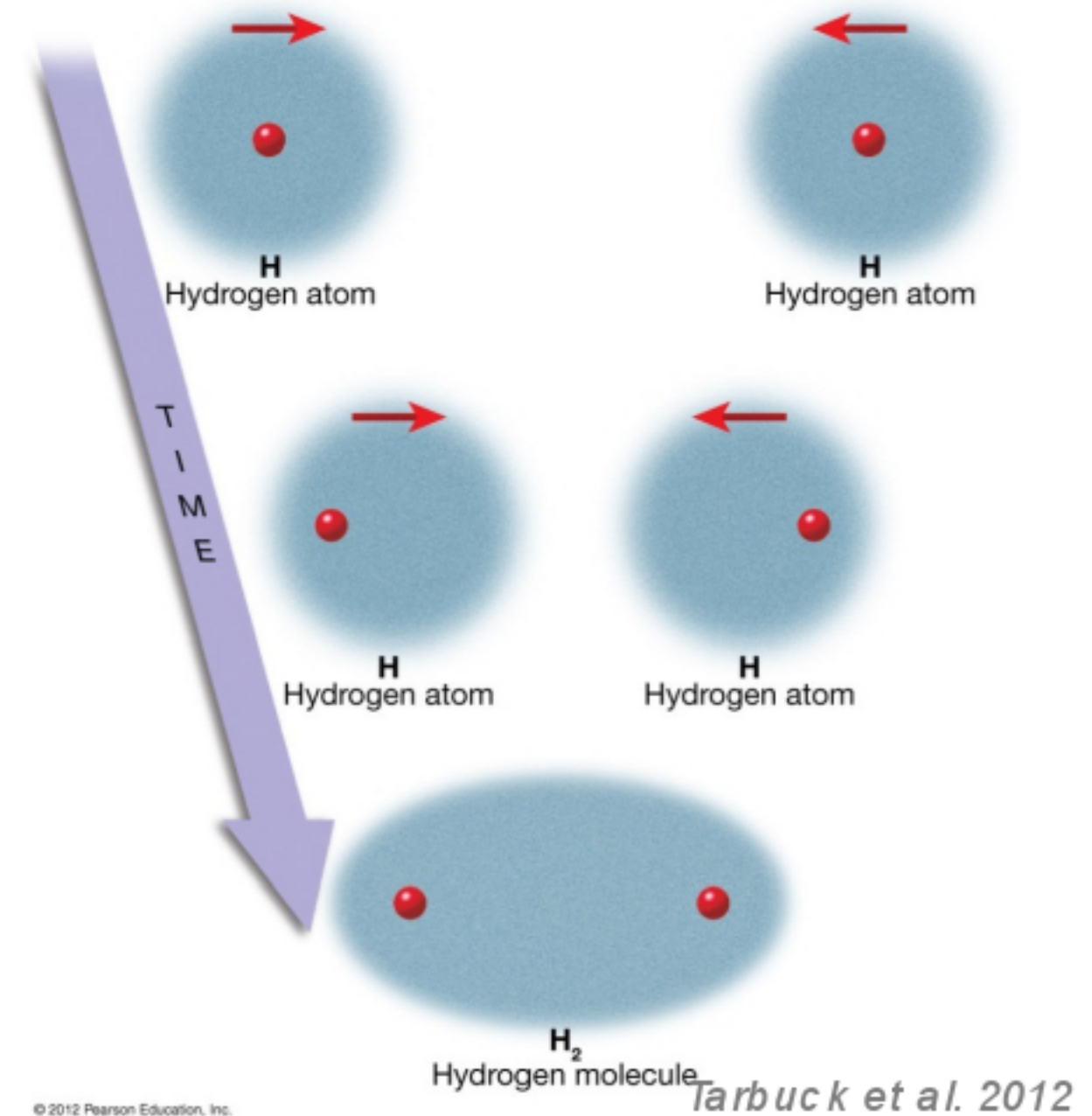
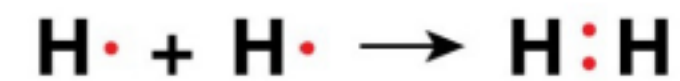
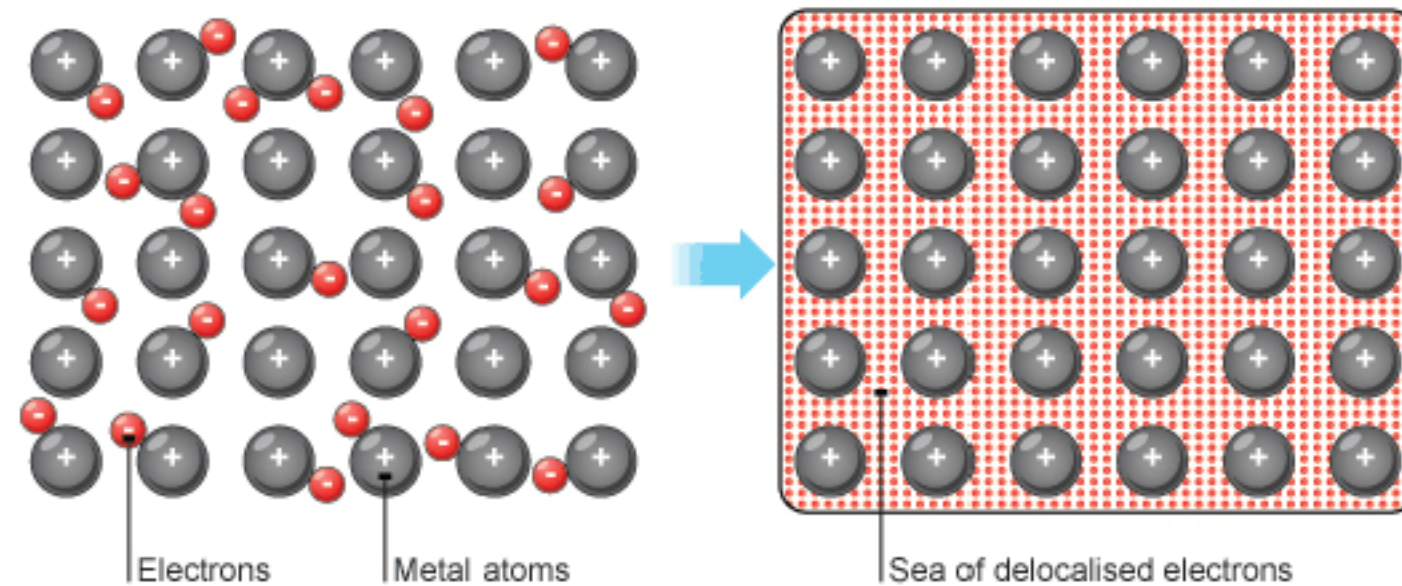
One atom gives up 1 or more of the valence electrons to another atom forming **anions** and **cations** which are then attracted to each other and form a compound



Covalent bonds

When forces holding atoms together are too strong they share a pair of electrons and form a **covalent bond**.

The attraction between protons of one atom and the electrons of the other keeps them together



Metallic bonds

Valence electrons move freely between atoms

Very strong bonds leading to high conductivity, strength etc

Minerals

Minerals

A mineral is a naturally occurring solid chemical substance that :

- is formed through geological processes
- has a **highly ordered** atomic structure
- has a characteristic **chemical** composition
- has specific physical properties.



NB minerals can be formed by 1 element or more elements

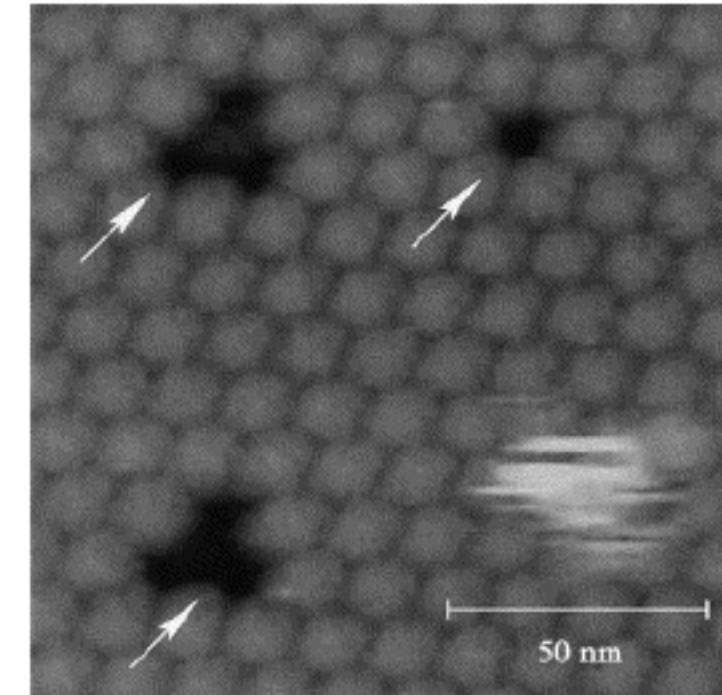
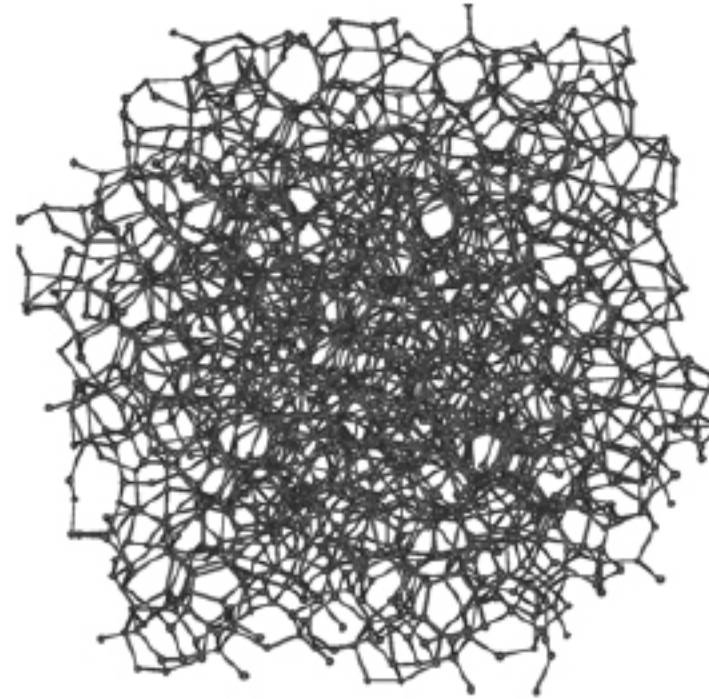


Birth and growth of a crystal

Crystals are mostly born in a liquid solution. Slowly, small nuclei form around which the crystal grows.

If sufficient time is available the crystal will grow developing a perfect structure.

If not atom organization will be **poor** to **non-existent**



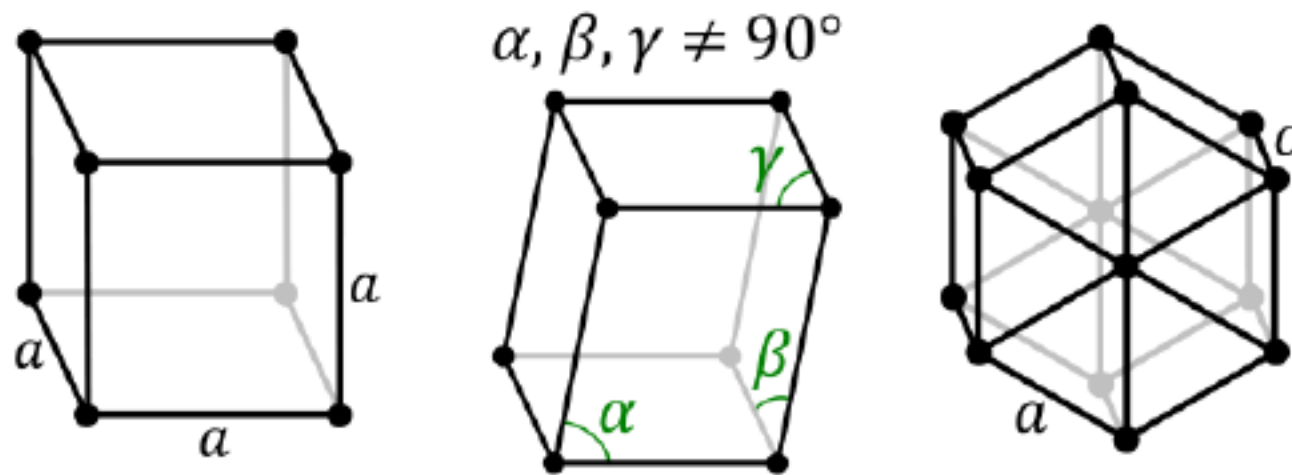
If the crystal can grow freely its crystal habit will be nicely displayed

If not, the crystallographic structure is visible only inside the crystal (x-rays)



Crystal structure and composition

Crystallography. A crystal is composed of **unit cells** stacked in 3D in an **orderly** way. The **unit cell** is a tiny box containing one or more atoms, a spatial arrangement of atoms. Different classes are defined on the basis of **lattice parameters**, (length and angle of cell edges, distances etc).



The way elements are arranged bears consequences on the physical properties of the **mineral**

Mineral classification on the base of composition

Silicates

Carbonates

Sulfates

Halides

Oxides

Sulfides

Phosphates

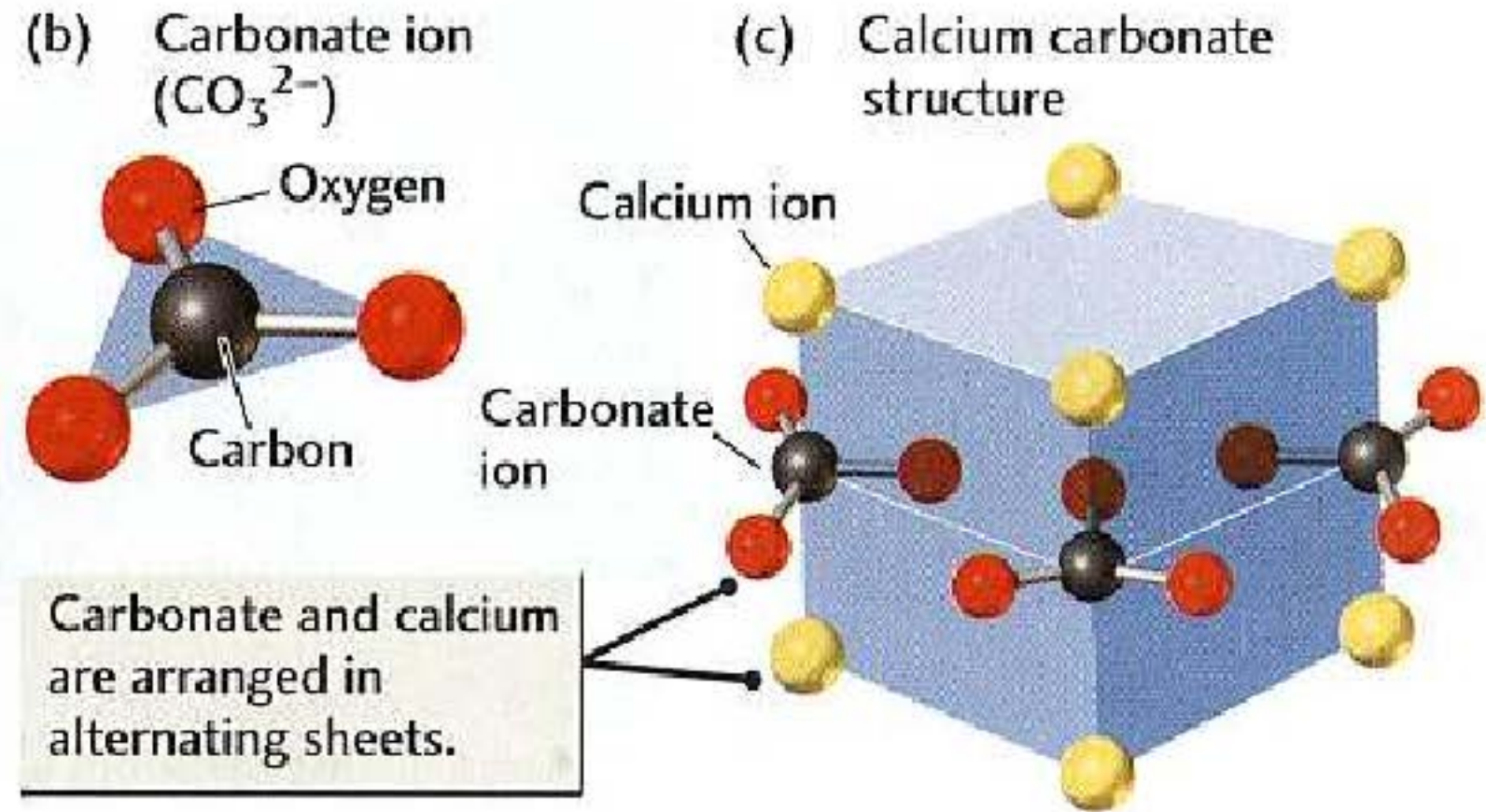
Elemental minerals

Organic minerals

Carbonates (CO_3^{2-})

Carbonate minerals constructed around the carbonate anion, $(\text{CO}_3)^{2-}$ with Ca between the sheet-like layers

- Calcite-Aragonite, CaCO_3
- Dolomite, $\text{CaMg}(\text{CO}_3)_2$

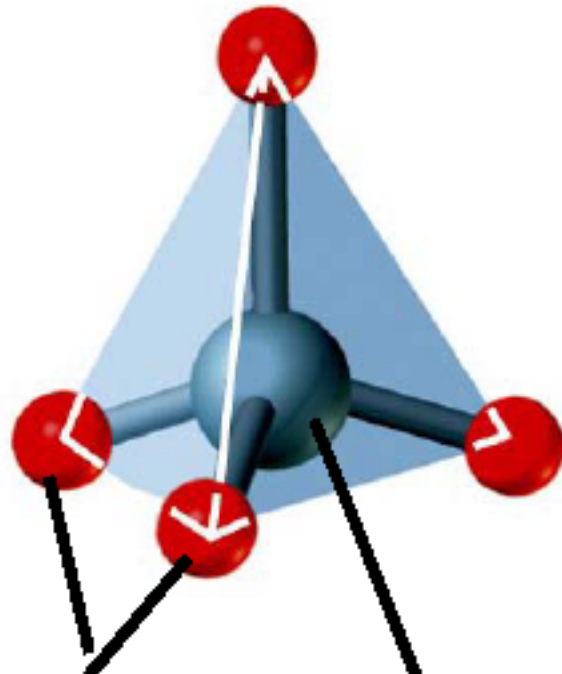
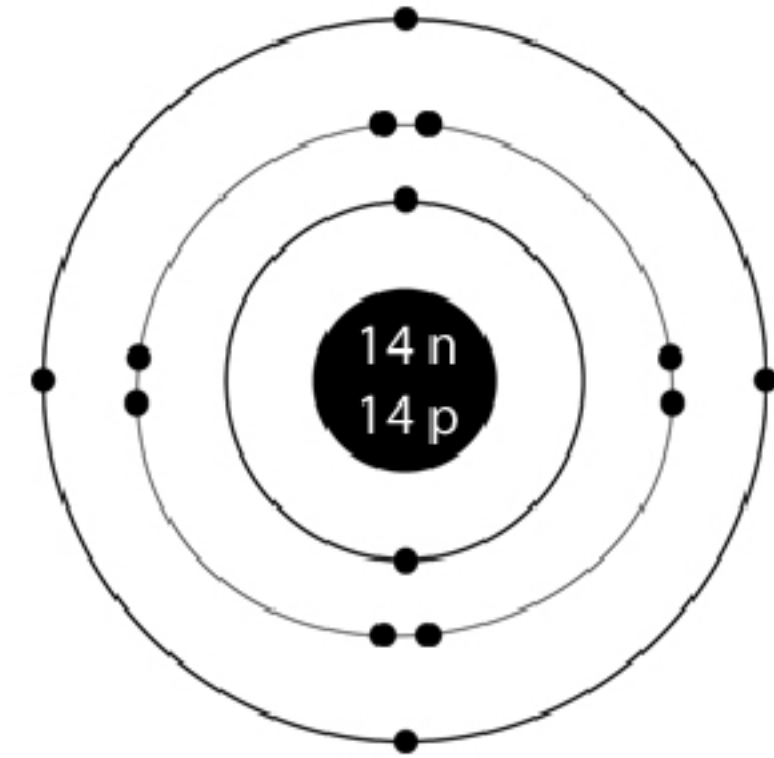


One of the most important minerals created by **living organisms!**



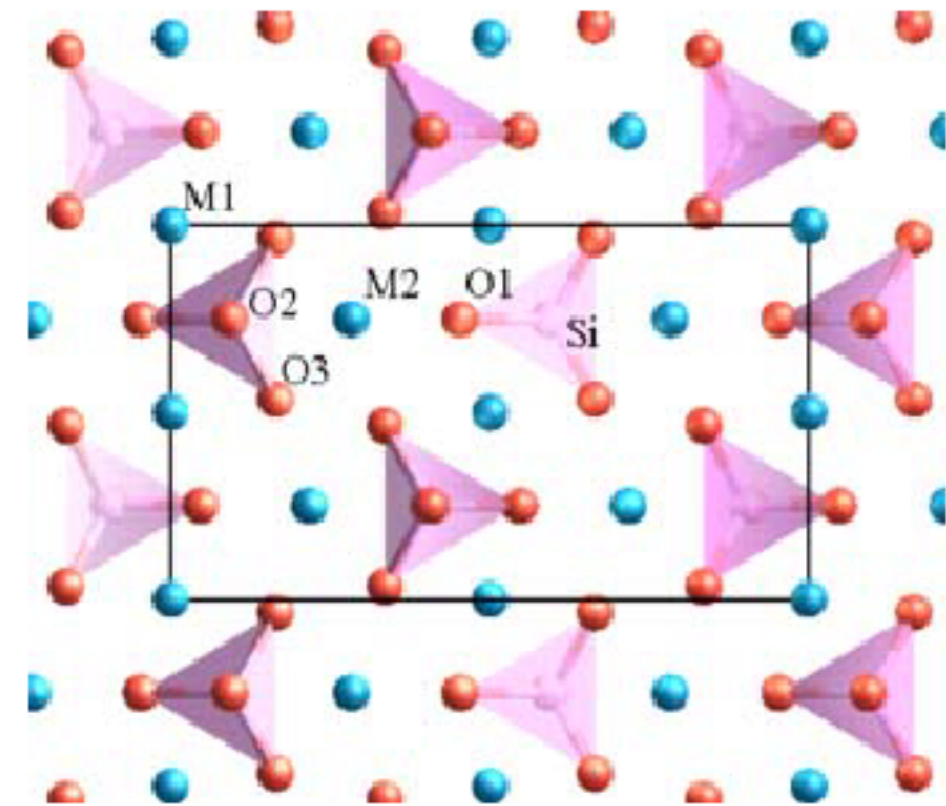
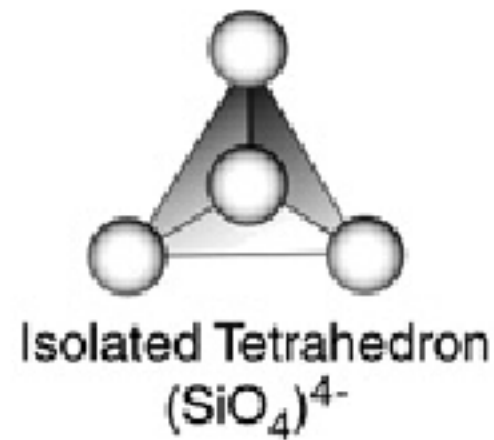
Silicate minerals**The most abundant minerals in Earth's crust**

Si: 4 electrons in the outer shell

oxygen ions (O²⁻) silicon ion (Si⁴⁺)Silicon ties with O to form the very stable (SiO₄)⁴⁻ tetrahedra**The tetrahedra can be linked in many different ways giving rise to very complicated structures****Other elements such as Al, Mg, Fe, Na, K and Ca commonly enter the structure**

Isolated tetrahedra

A Si tetrahedron surrounded by cations (mainly Fe^{2+} and Mg^{2+})



Olivine $(\text{Mg,Fe})_2\text{SiO}_4$ (the most important component of the Earth' mantle)

Elements such as Mg and Fe find their place between tetrahedra (**blue spots**)

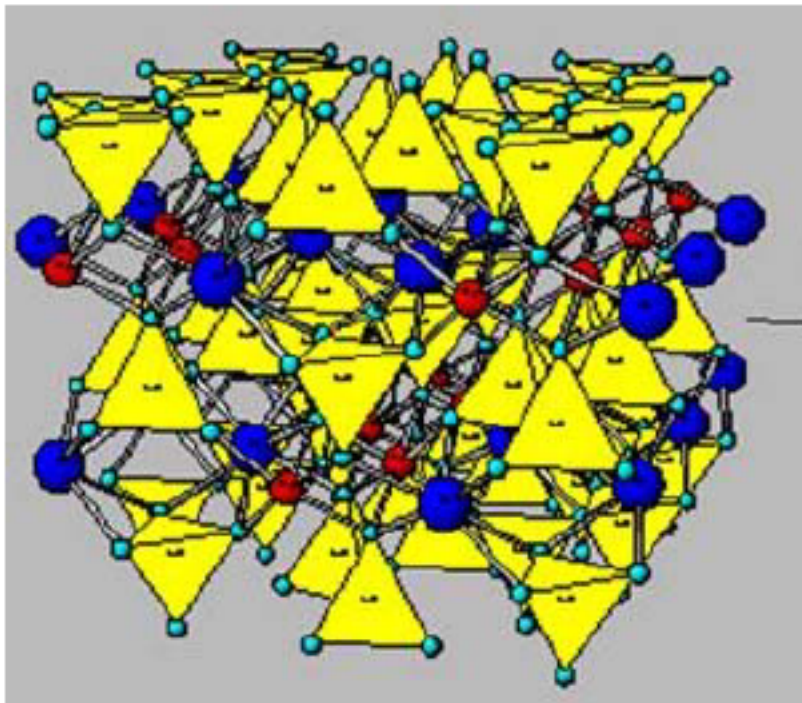
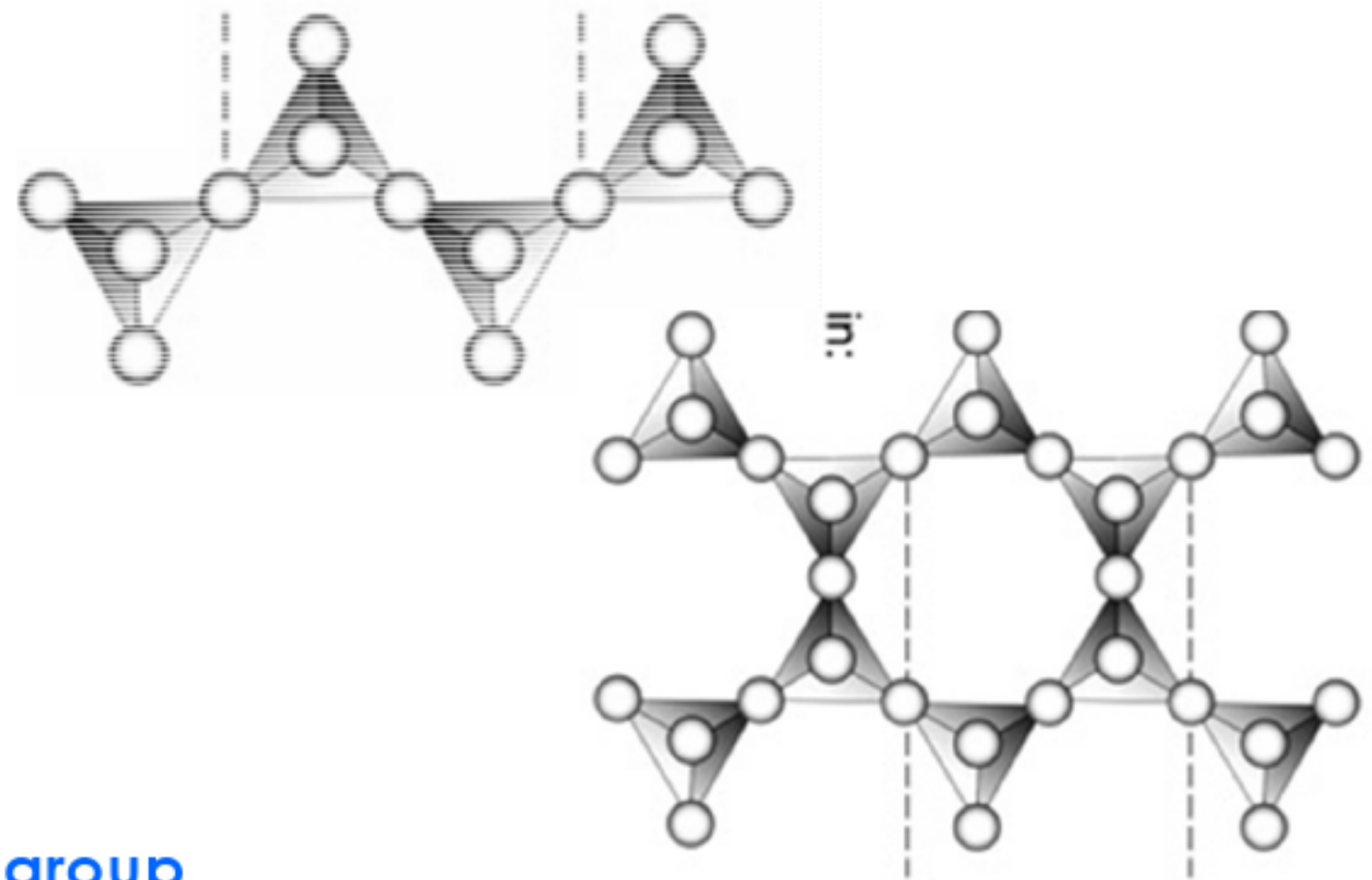


Chain silicates (inosilicates)

Adjoining tetrahedron share 1 oxygen forming chains.

Remaining negative charges are compensated by cations. The loose structure allows many different mineral types

Single or double chains exist



Pyroxene group

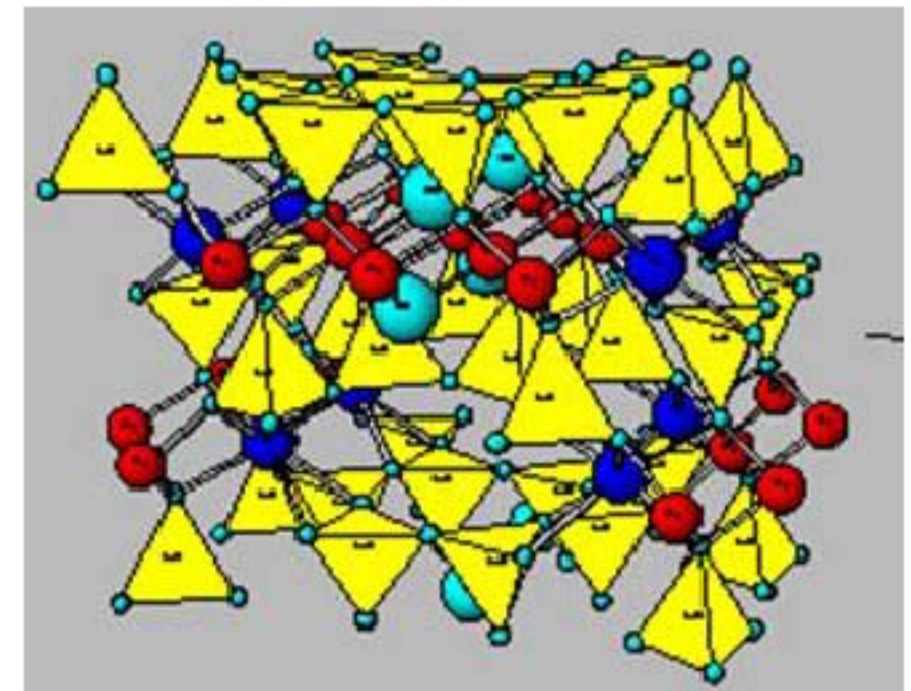


Important in the mantle and elsewhere

Amphiboles

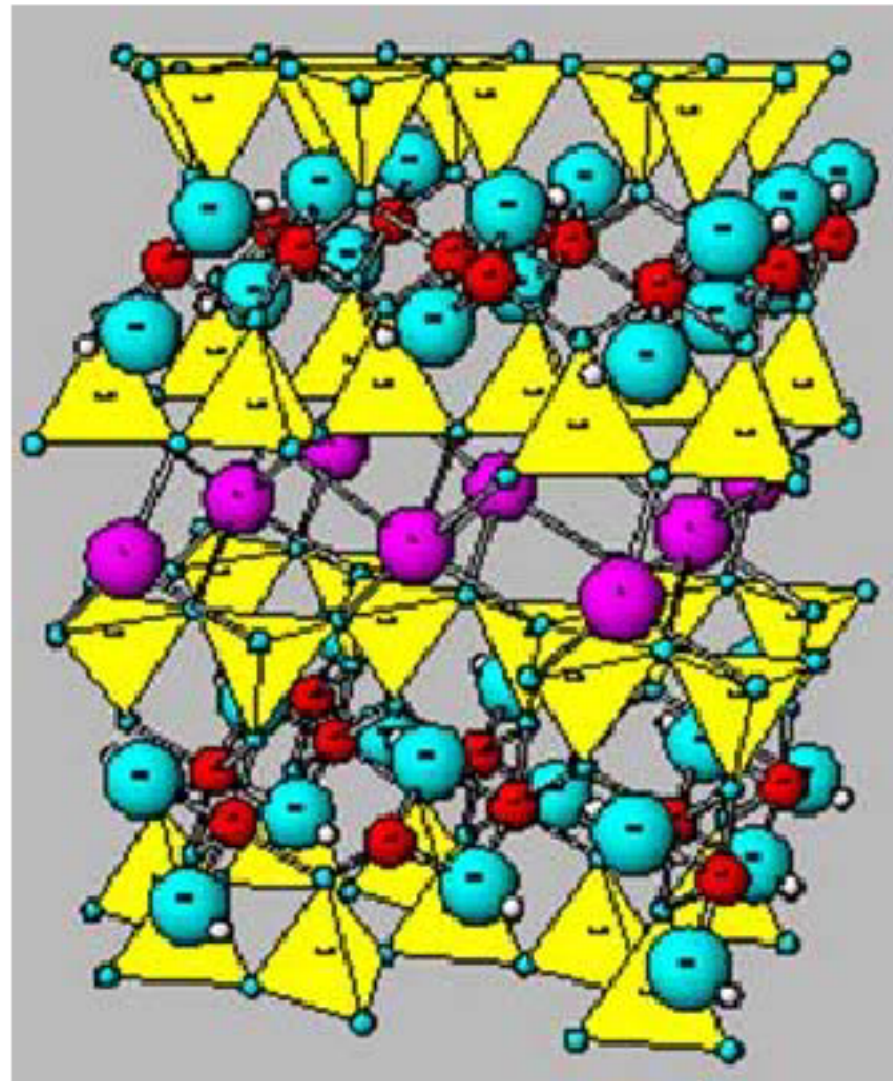
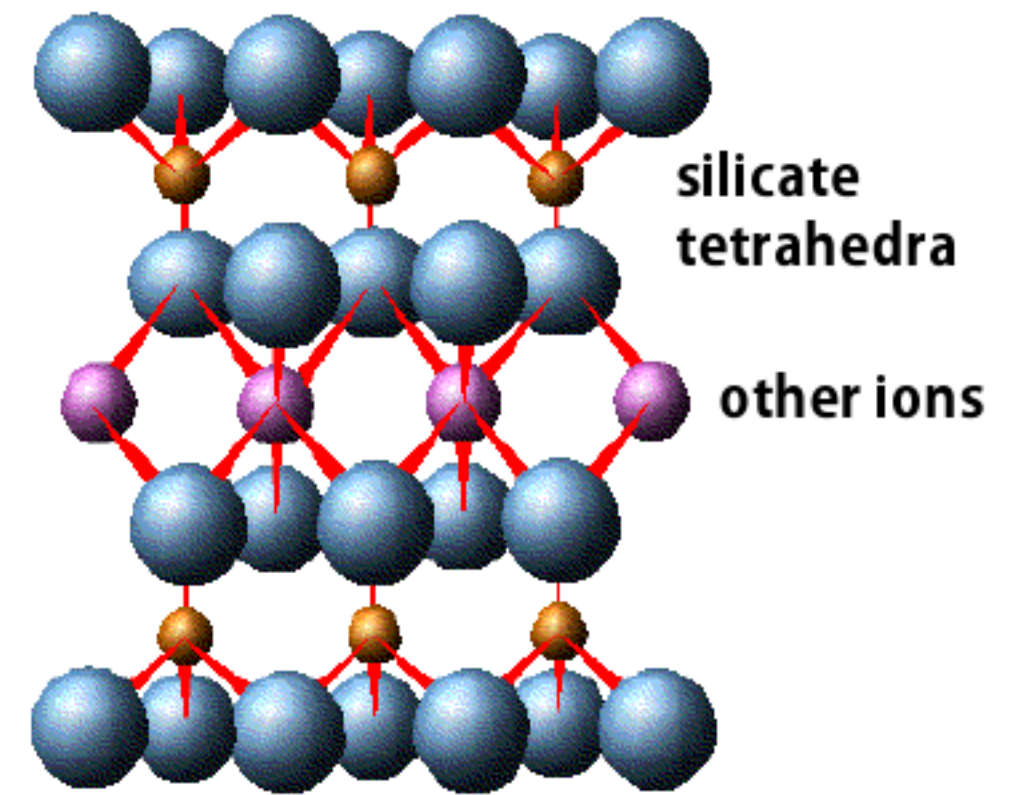


Note the presence of the (OH) oxydrile: it means water!



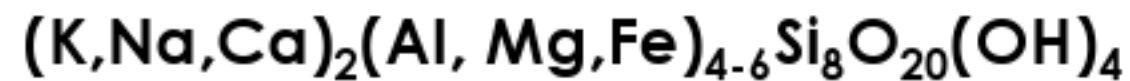
Sheet silicates (phyllosilicates)

Si tetrahedrons are linked 2 dimensionally into sheets. Between two planes of Si tetrahedrons is an octahedral layer (either filled with Al (as one end member) or with some divalent cation (Fe^{+2} , Mg^{+2}) at another extreme).

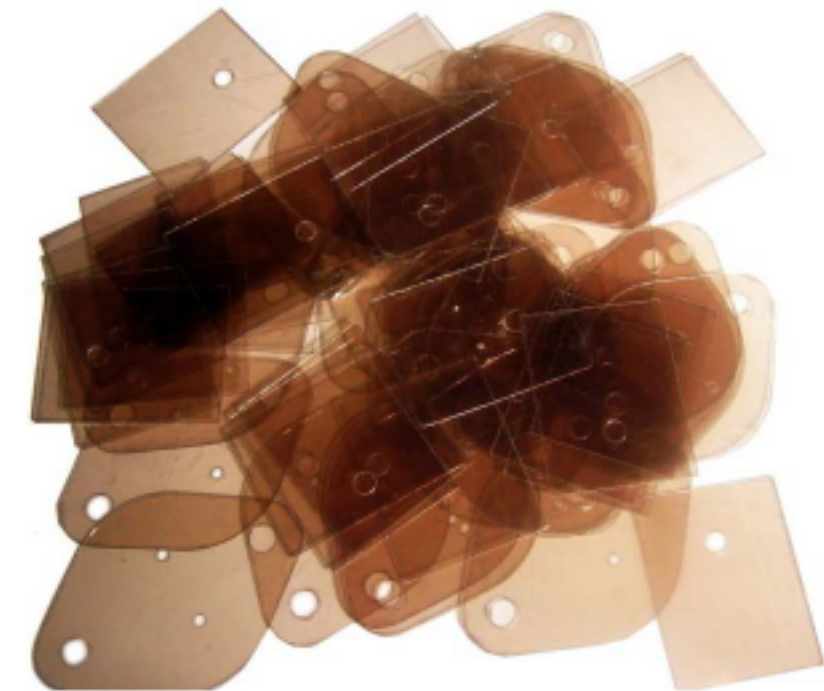


Micas

(e.g. muscovite and biotite)



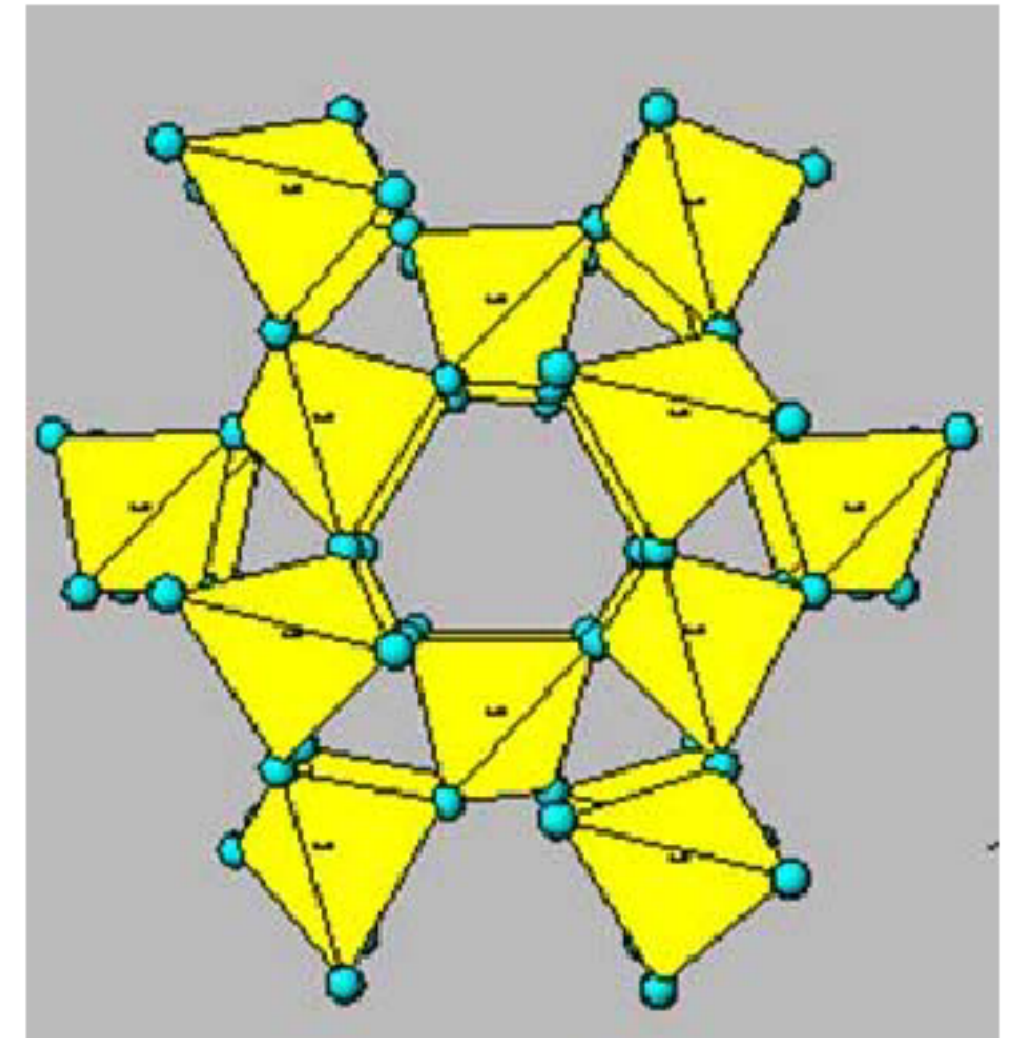
Have a very important basal cleavage



Framework (tecto) silicates

Are the most complex form of Si tetrahedral linkages: a 3D sharing of O.

quartz (SiO_2), a 3D linkage of O which results in a completely neutral charge, no need for counterbalancing cations, and **a very strong resistance to chemical weathering.**



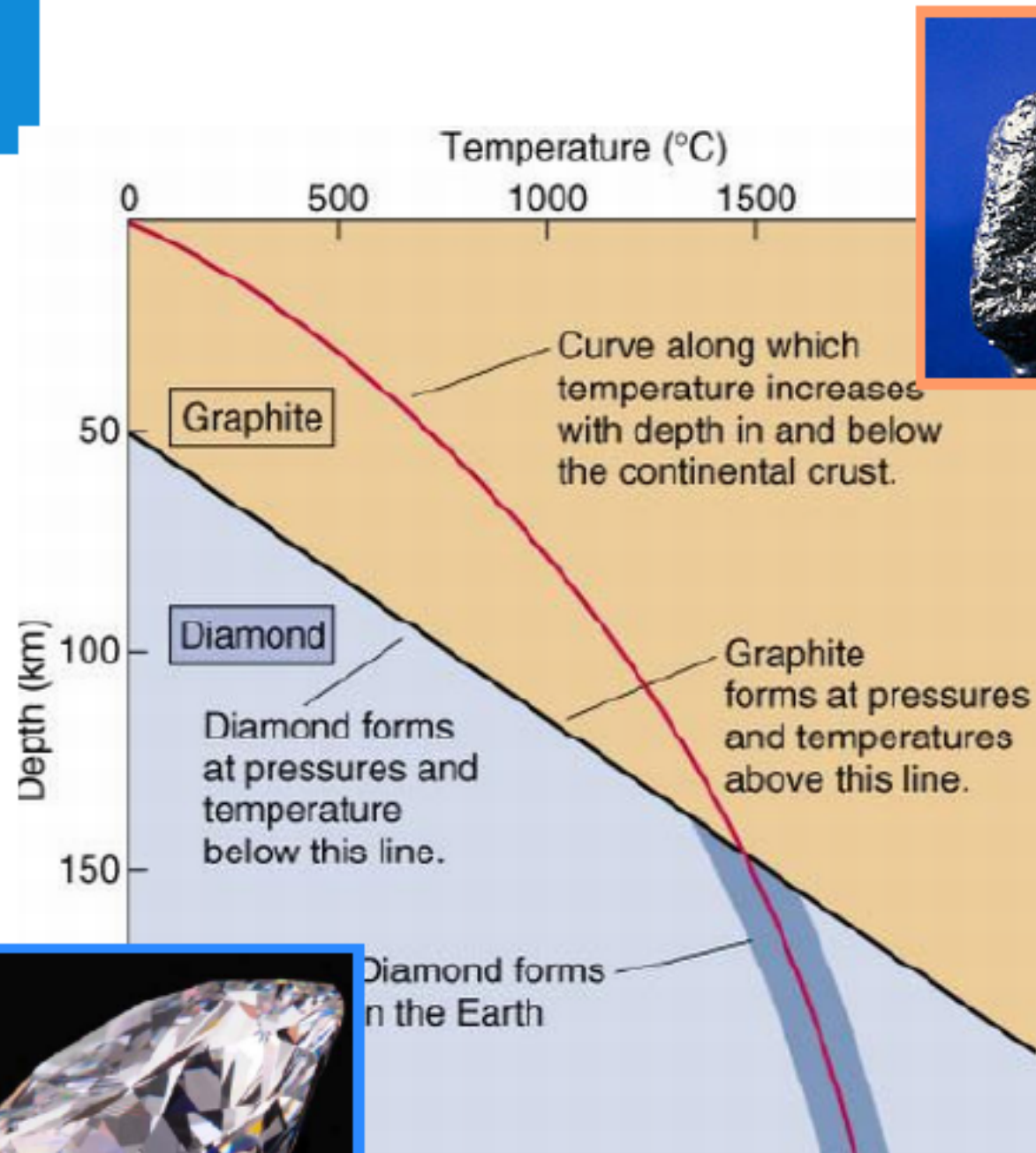
In nature, **Al** can easily substitute Si in the tetrahedral configuration (Al has about the same size and position on the periodic table).

Because Al has one less positive charge than Si, the addition of Al in place of Si creates a net negative charge that must be counterbalanced by a cation.

Tectosilicates with Al are **feldspars (alkali feldspars and plagioclases)**

Stability fields of minerals: single minerals

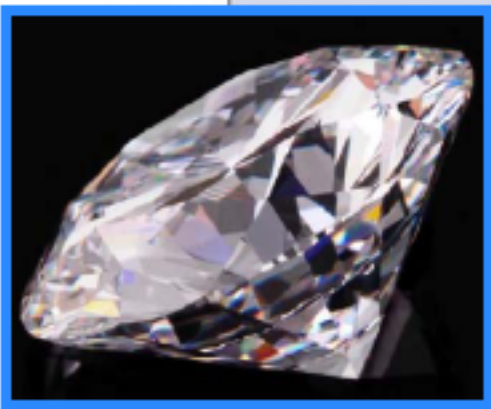
The crystal structure can change with changing pressure and temperature



Polymorphism: Some elements and compounds form two or more different minerals depending on P, T conditions (same composition, different internal structure)

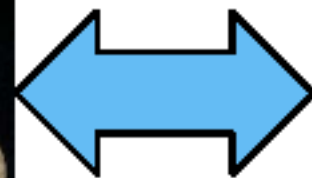
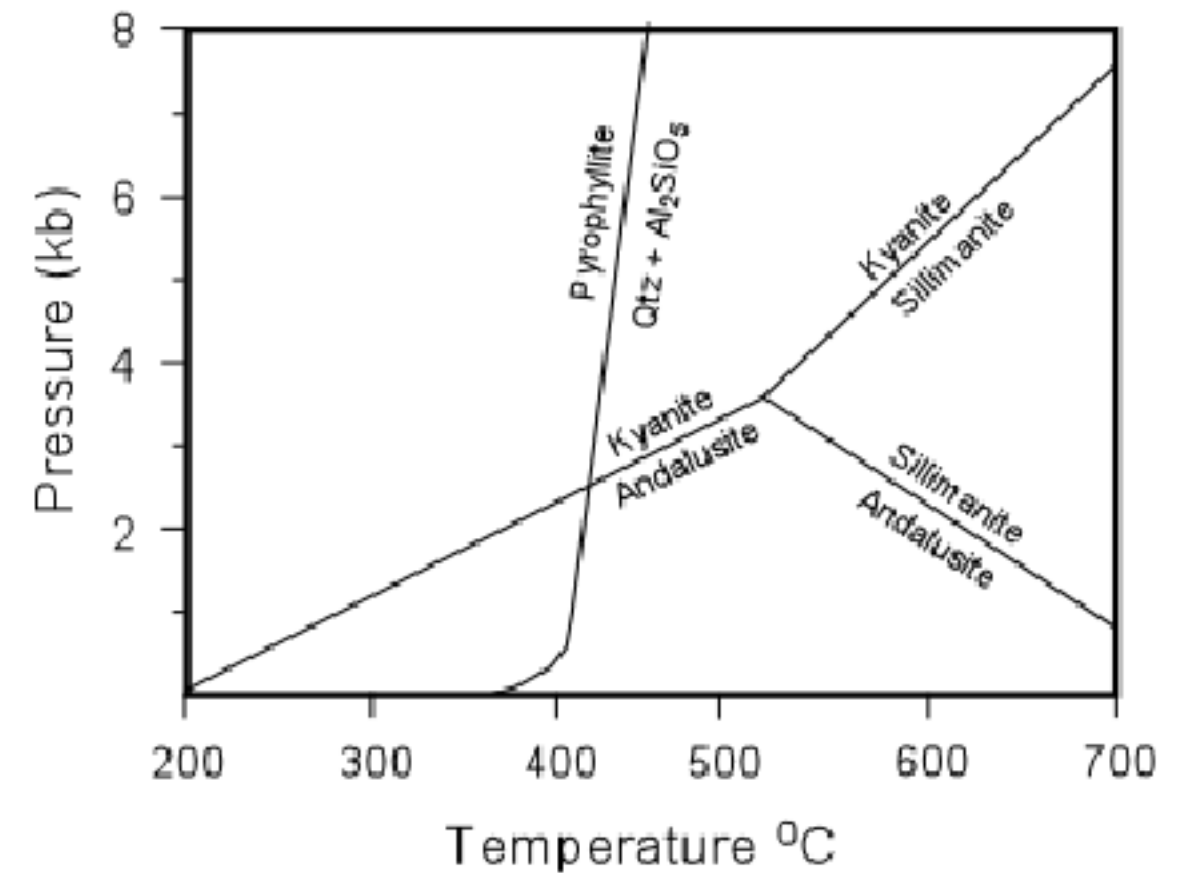
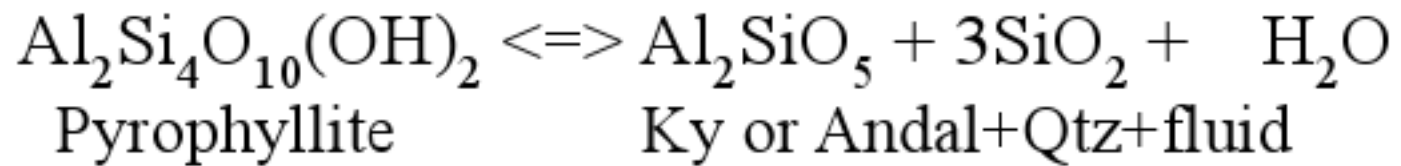
- C graphite, diamond
- CaCO_3 calcite, aragonite
-

very useful as minerals become indicative of specific T, p conditions



Stability fields: reactions

With changing pressure and temperature, minerals can even become unstable and produce other minerals



Pyrophyllite: mineral with many industrial applications (insecticides, thermal insulation ...)

And now..... the rocks

Rocks: naturally occurring aggregates of (one or more) minerals

Igneous rocks

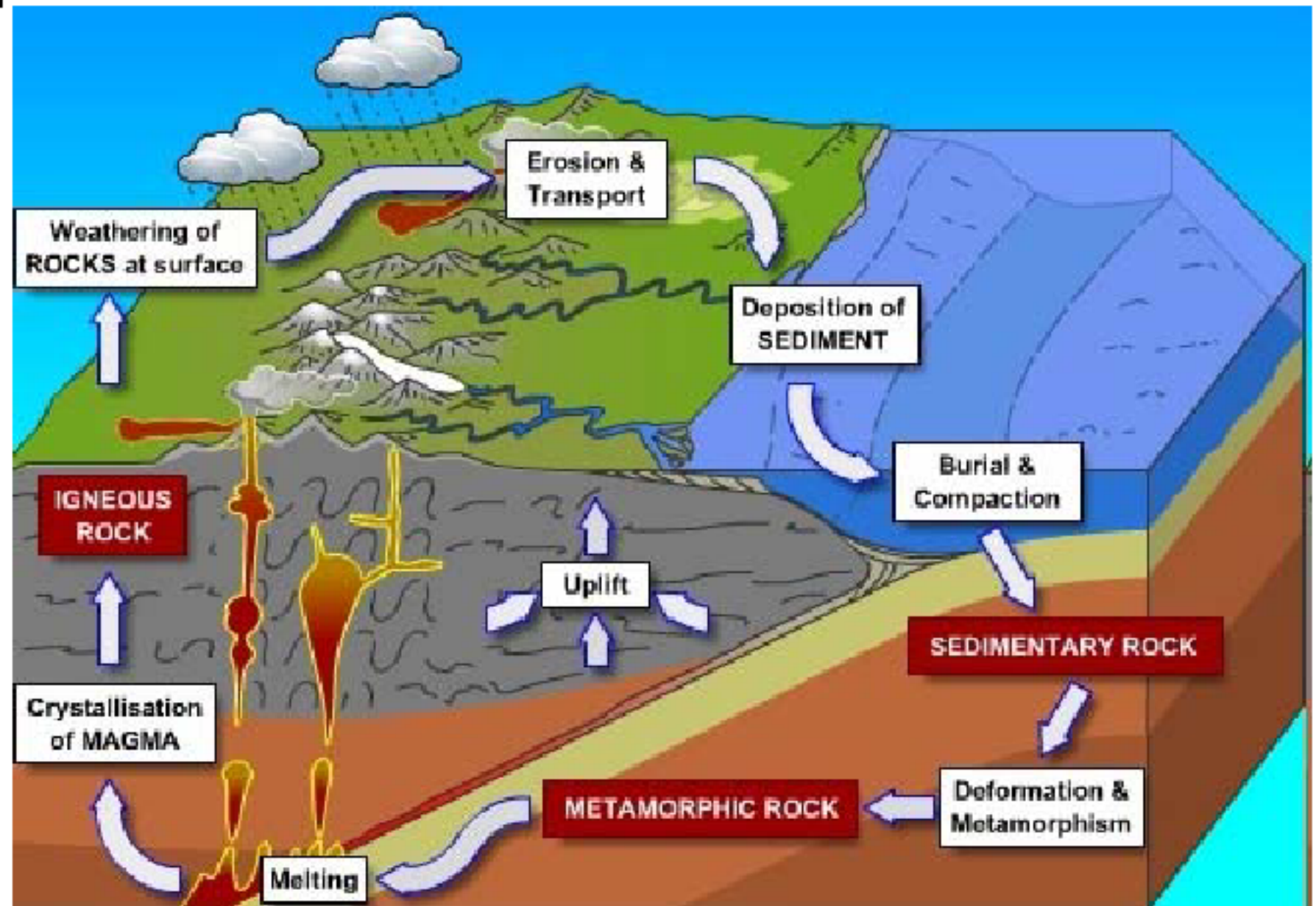
Formed by solidification of magma

Sedimentary rocks

Formed and deposited at the Earth's surface, by erosion of older rocks, biologic or chemical processes

Metamorphic rocks

Formed by the alteration of pre-existing rocks in response to changing pressure and temperature



Magmatic (igneous) rocks

Igneous rocks result from the consolidation of magma at different conditions

Three main stages

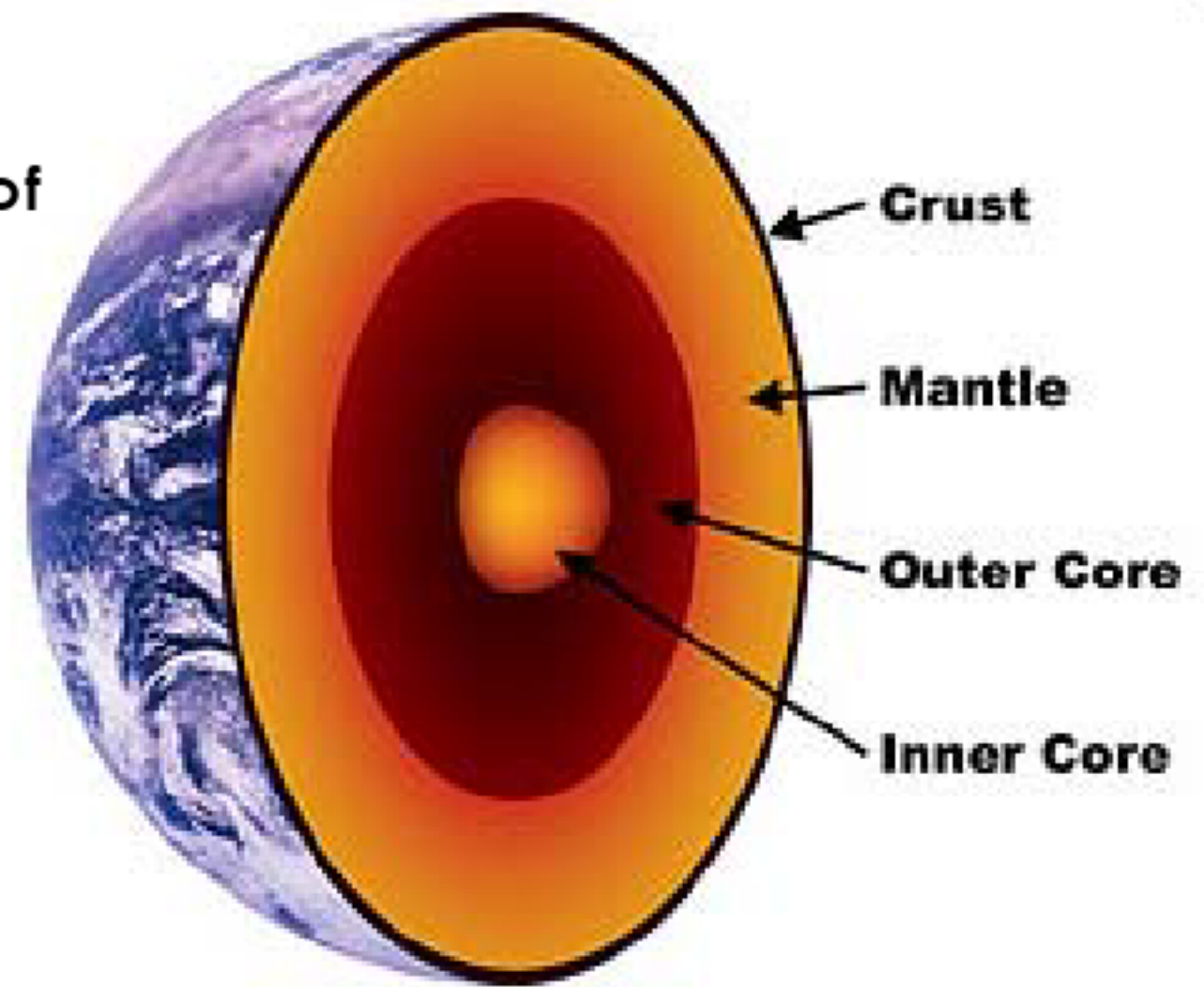
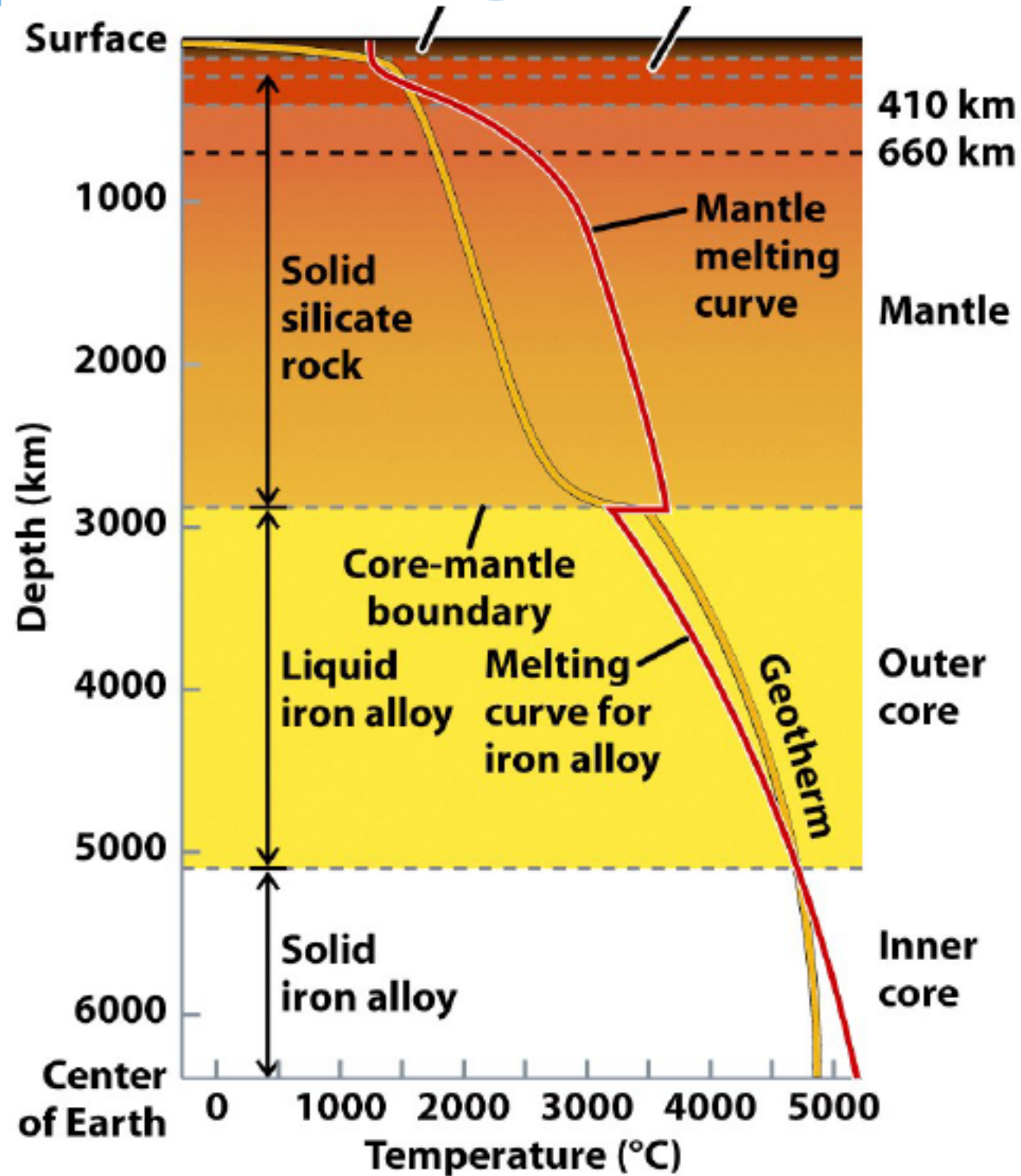
- 1) Magma generation
- 2) Magma transport (ascent)
- 3) Consolidation



Melt generation

The outer 3000km of the Earth are made of **solid rocks**

Where does **magma** come from?



Temperatures do increase with depth
But also the melting point also does



With the partial exception of core,
the Earth is **solid**.



To produce magmatism, one needs
to **melt** rocks.

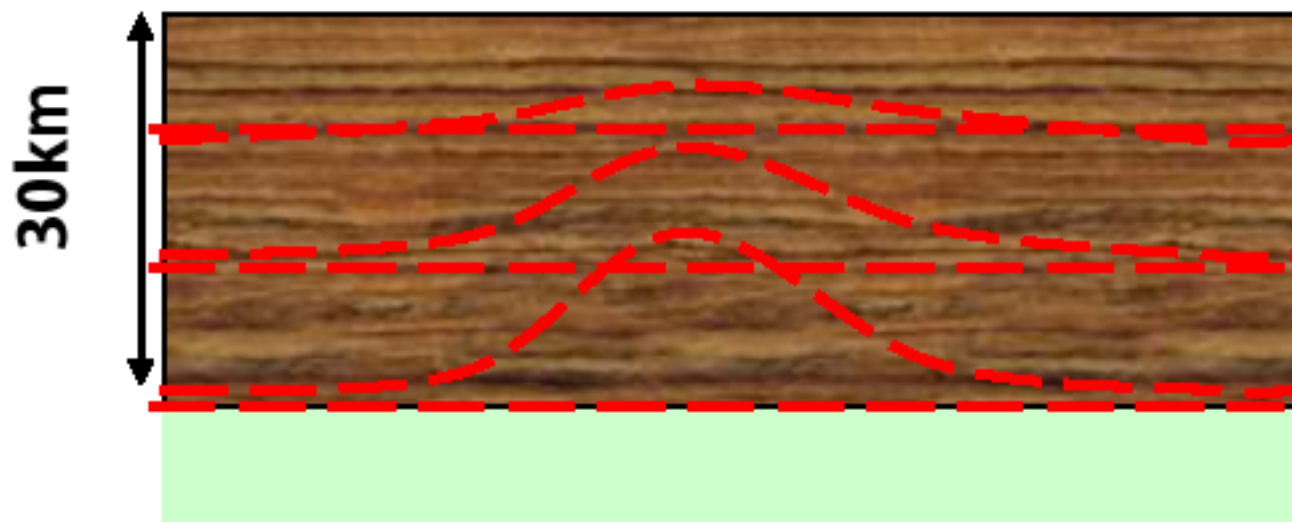
There are 3 different ways to do this

1. Increasing the temperature (=creating a thermal anomaly)
2. Decreasing the pressure
3. Adding fluids (water)

1. Creating a (large enough) thermal anomaly

Implies warming up rocks at a given depth until they start melting

100s of °C are needed!

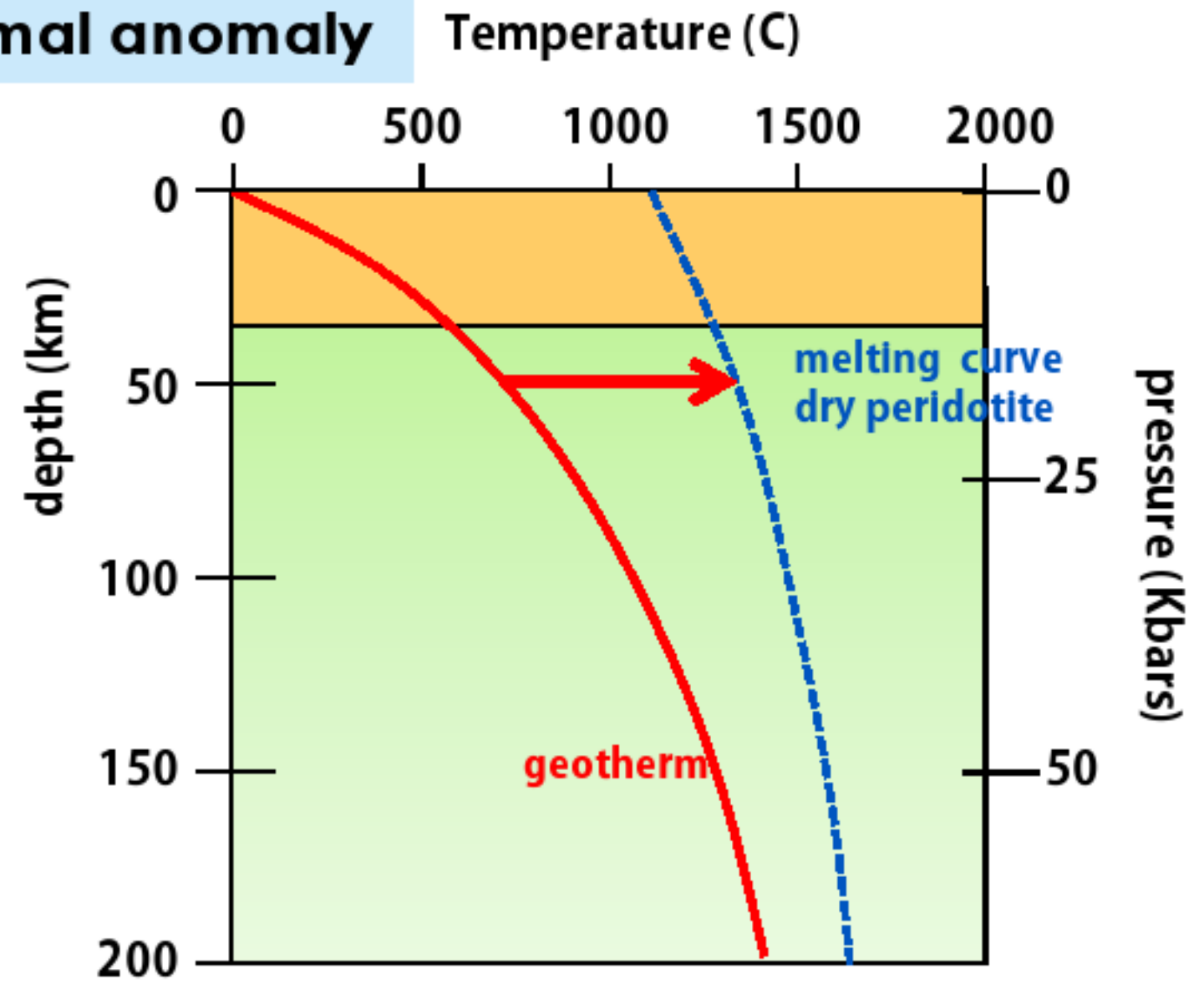


200° C
400° C
600° C

Heating of this magnitude is not trivial

Materials can be heated up by **conduction** or **advection**.

The 1st does not work, the 2nd needs magma



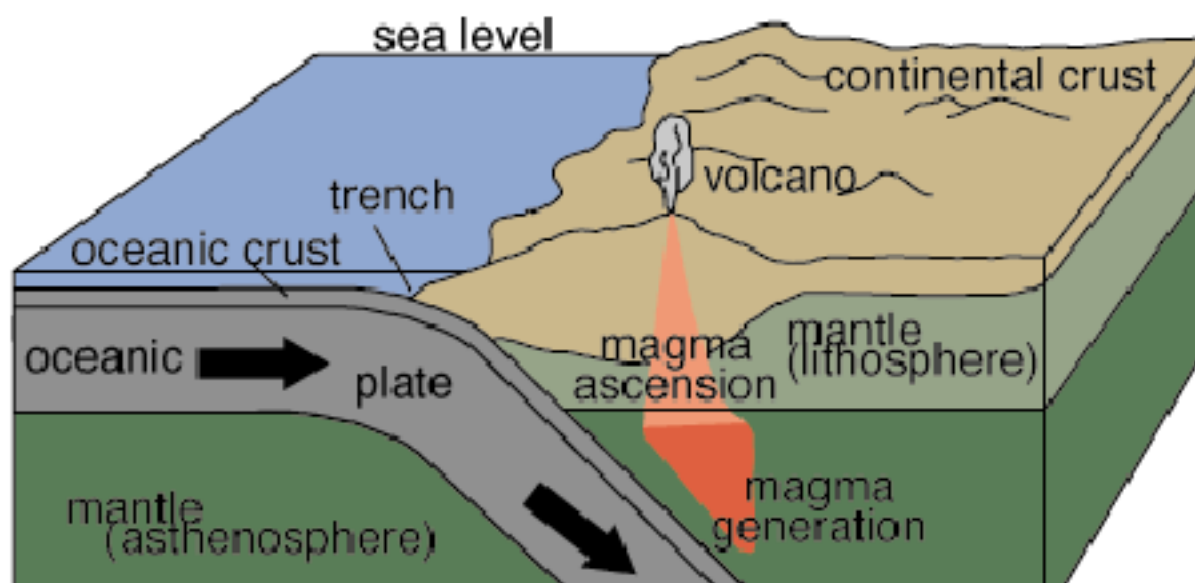
2. Melting by decompression

Melting temperatures are pressure-dependent

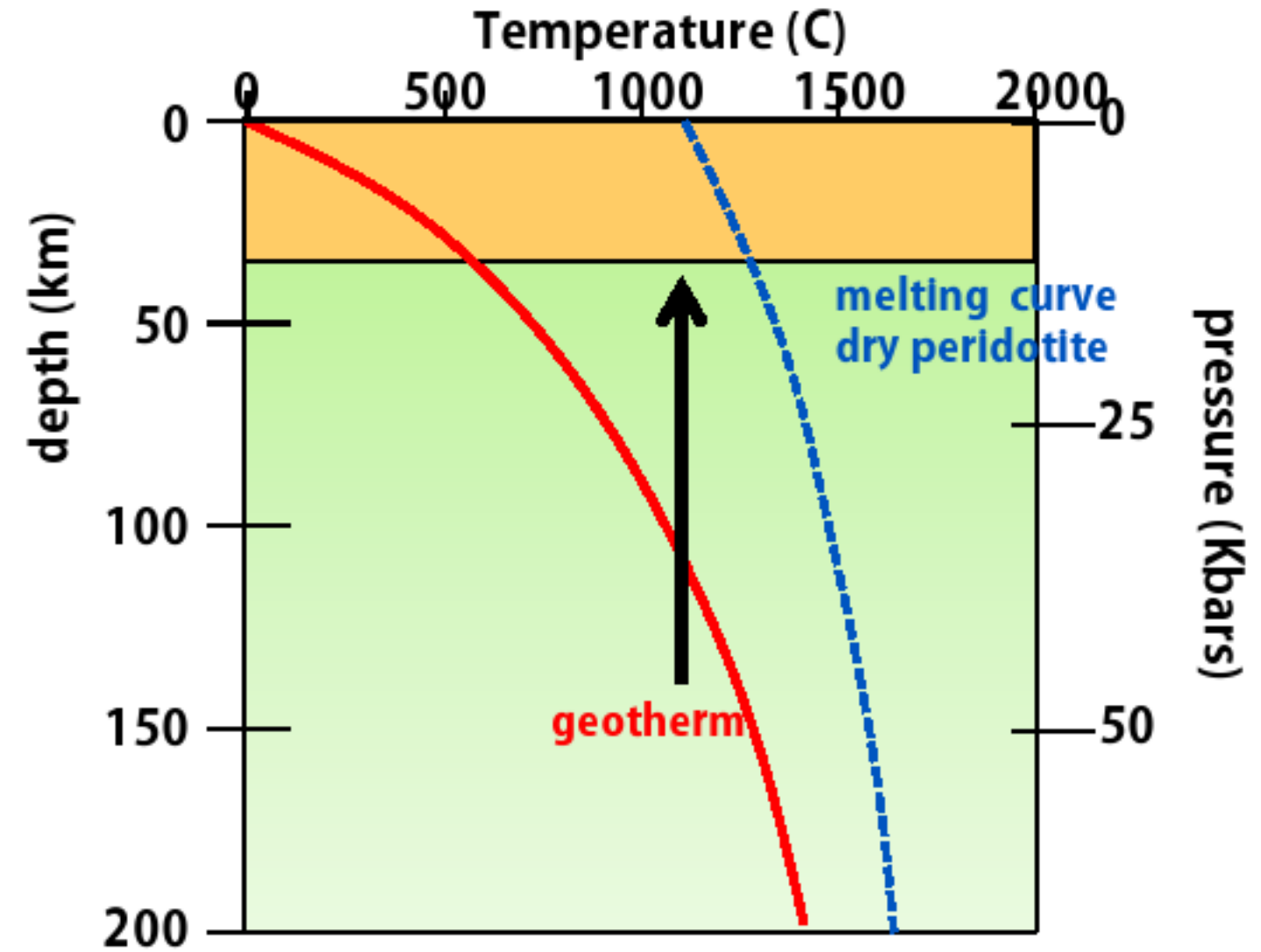
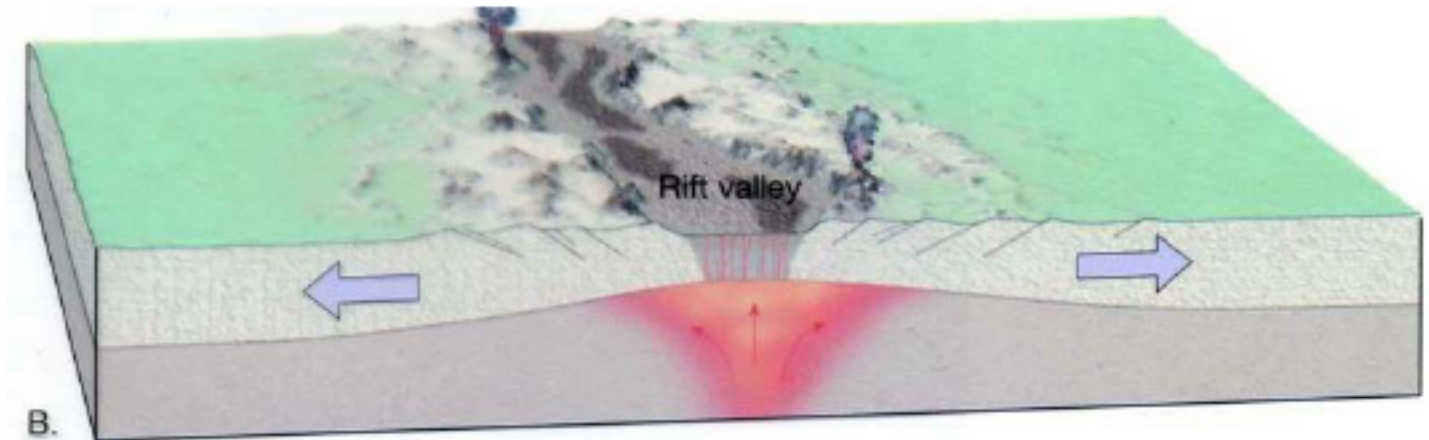
If **pressures** decrease enough, while **temperatures** do not, melting can take place

Two ways to do this:

Move rock upward **very rapidly**



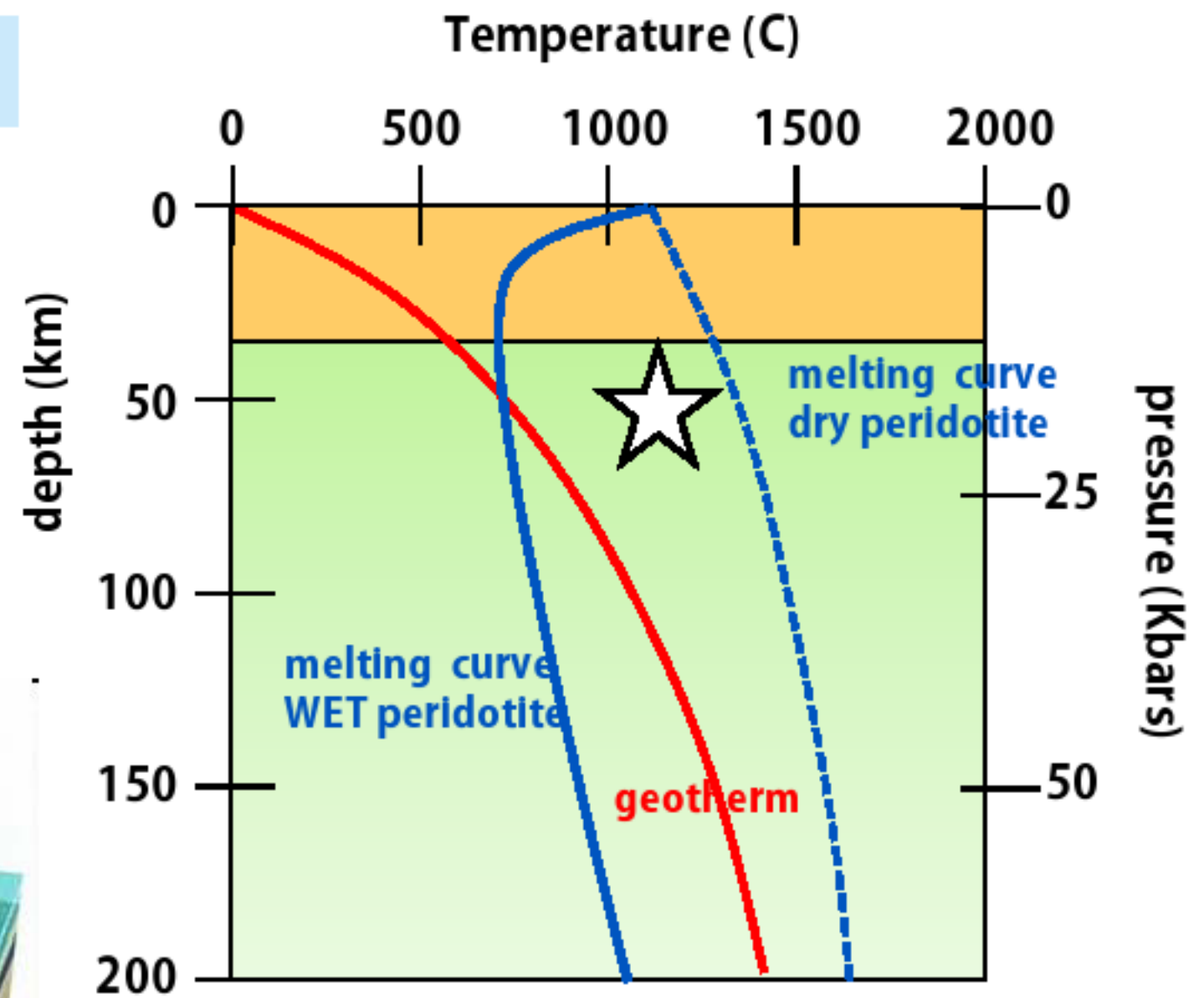
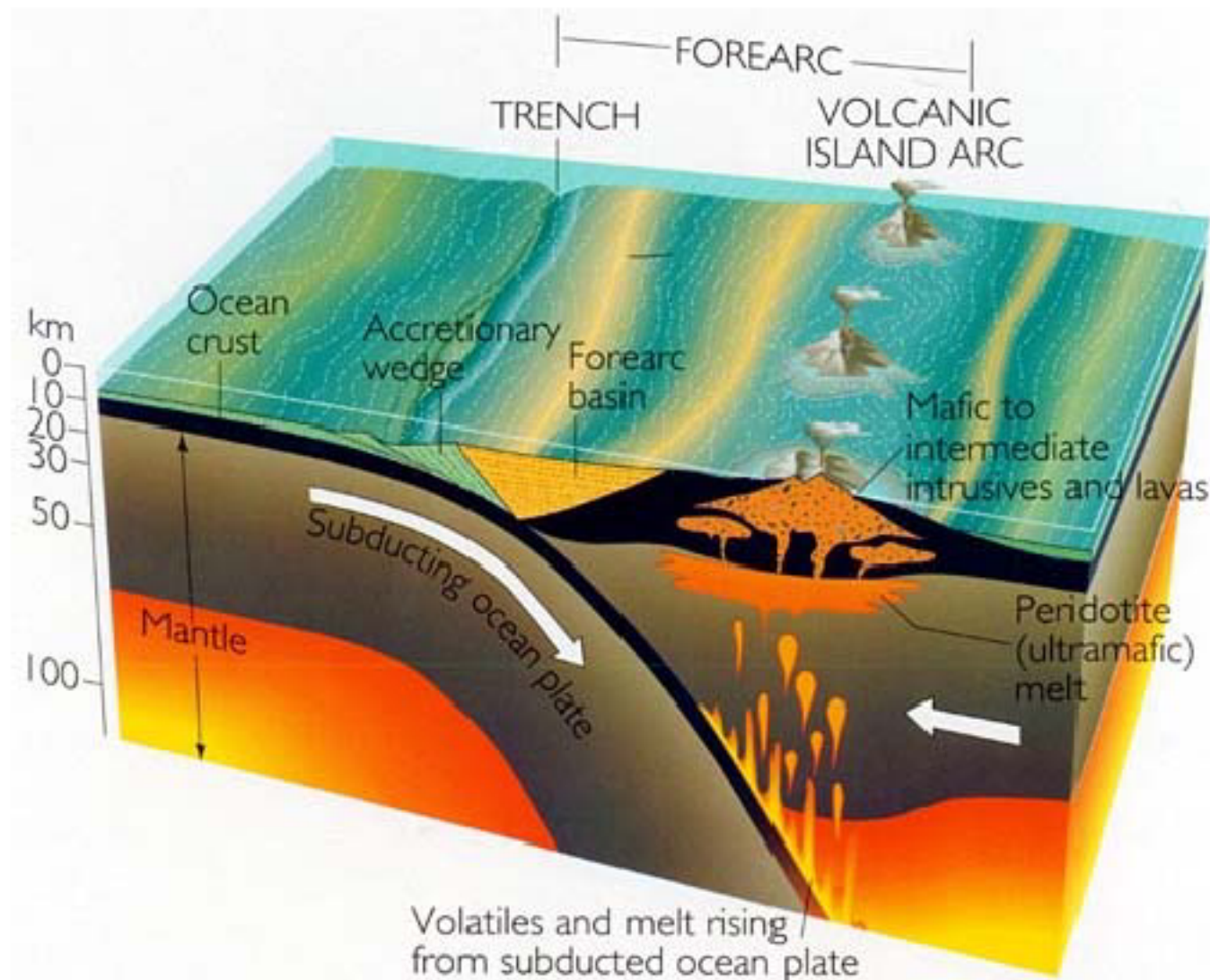
Decrease the overburden



3. Melting by addition of fluids

Fluids provoke a substantial decrease in melting temperatures

Rocks start melting even if they do not move



This is what happens above subduction zones

(any idea where do the fluids come from?)

Melting starts! What kind of magma is produced?

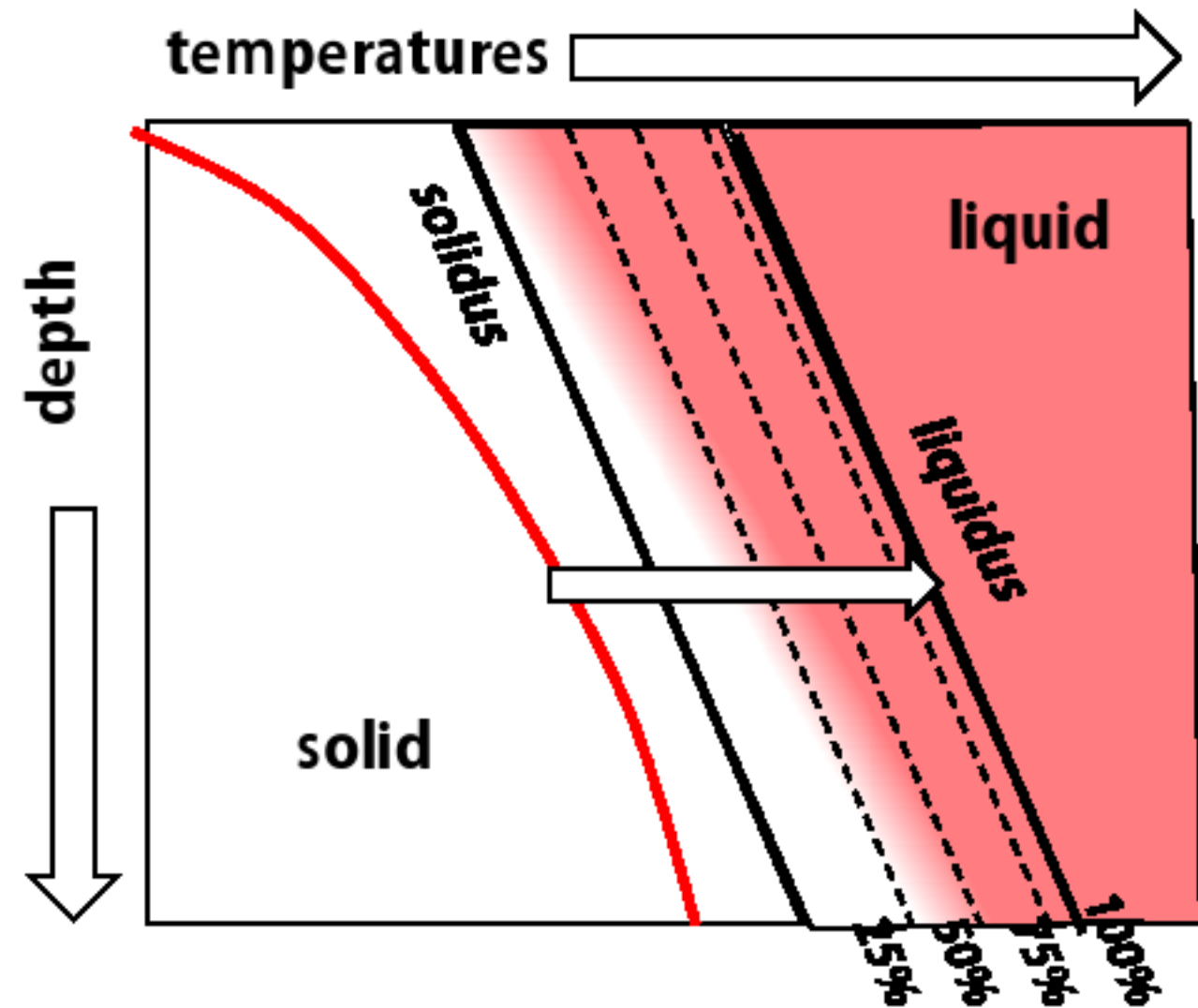
for **1 mineral** rock:

- melting takes place at one T
- the melt will have the **same composition** of the rock

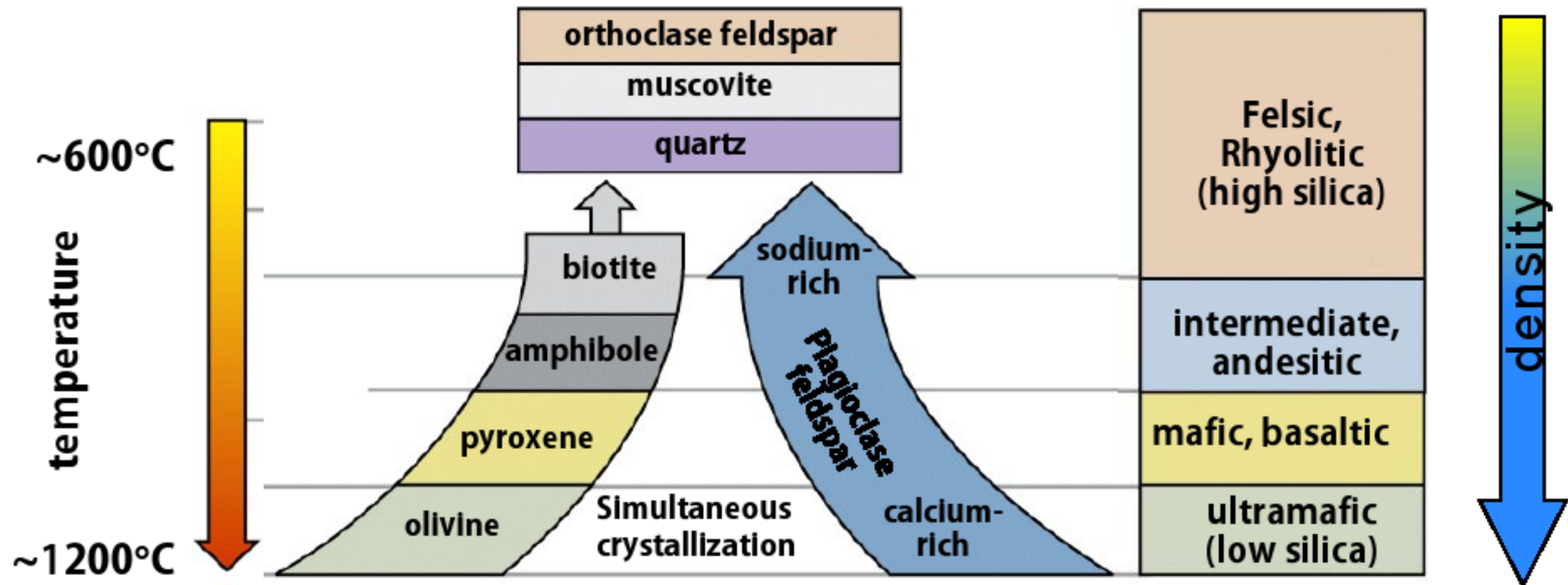
olivine	amphibole plagioclase	quartz
1000 °C	800 °C	600 °C

If rocks are composed of **>1 mineral**

- each mineral will melt at **different temperatures**.
- temperature at the first drop of melt is the *solidus*, that at total melt is *liquidus*
- The composition of the melt will **not coincide** with that of the rock unless melting is complete. This is **differentiation**

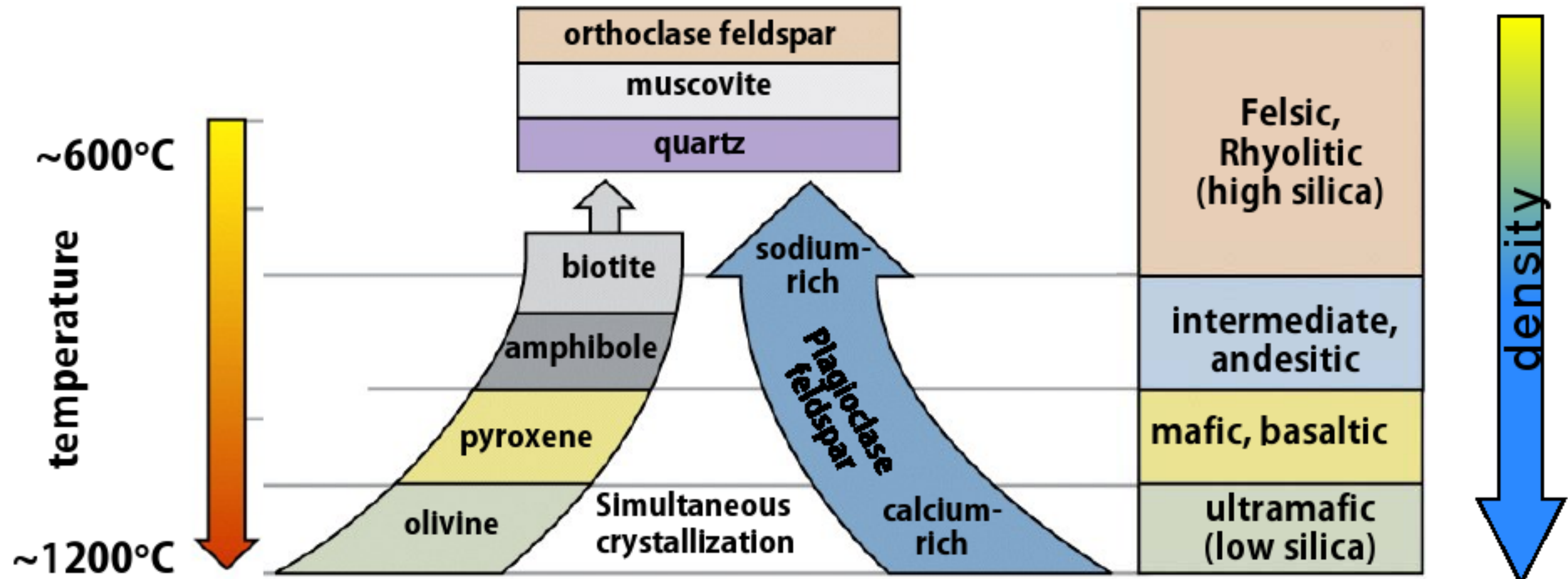


Predicting **differentiation**: Bowen's reaction series



The composition of the magma depends on a) the **primary composition** of the rock and b) the **degree** of melting

Silica rich magmas are **lighter** than silica-poor ones



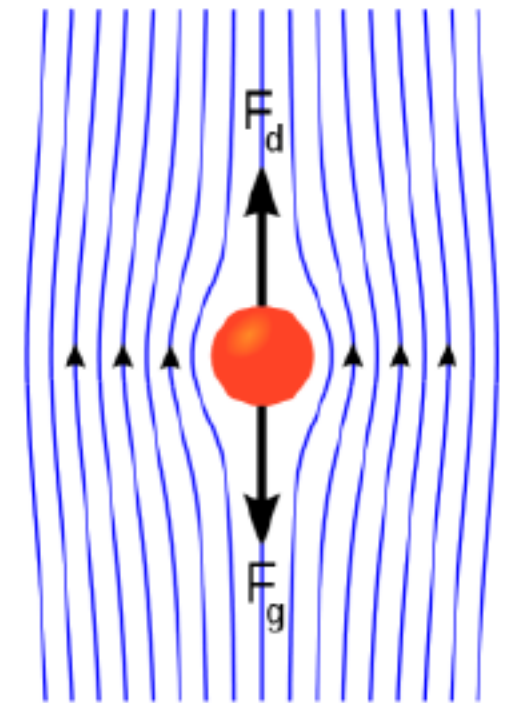
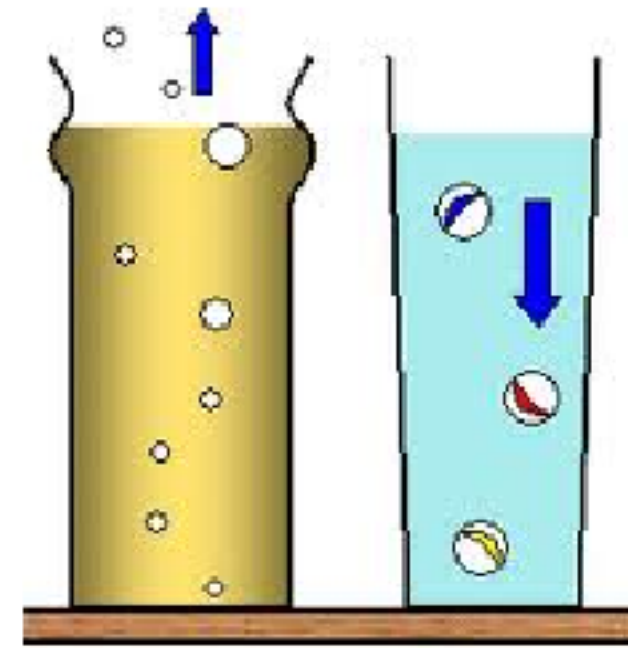
Initial magmas are light magmas and tend to escape easily

feldspar	quartz	biotite	pyroxene	olivine
2.56	2.62	3.09	3.2	3.32
magma	basalt	andesite	rhyolite	
	2.6-2.8	2.45-2.50	2.18-2.25	

The upward movement of the magma

Magma starts moving upward driven by density difference (**buoyancy**).

A simple description is **Stoke's law** which describes density driven movements



$$v_s = \frac{\sigma_p - \sigma_f}{\mu} \frac{2}{9} g R^2$$

v_s = velocity
 σ_p, σ_f = density of particle and fluid
 R = particle diameter
 μ = fluid viscosity

Some values $v_s = ?$

See exercise

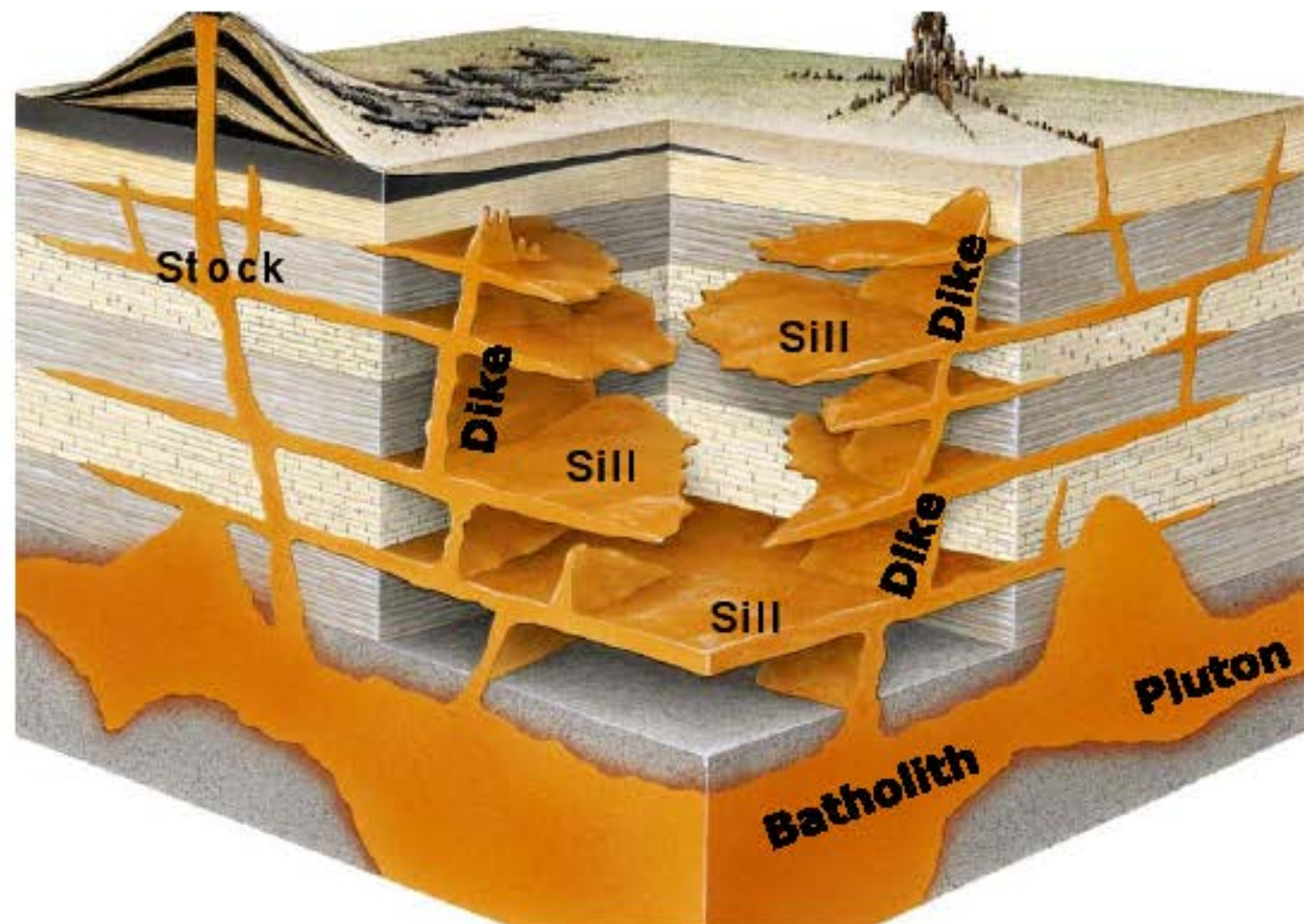
The good news is that rocks, undoubtedly poor fluids, can break creating pathways for the magma to move
 (obviously extra energy is needed)

Where does the magma does stop?

While moving upward, magma will cool and the $\Delta\rho$ will also decrease.

If the $(\delta_p - \delta_f)$ remains large (negative) the magma can reach the surface (volcano, lava).

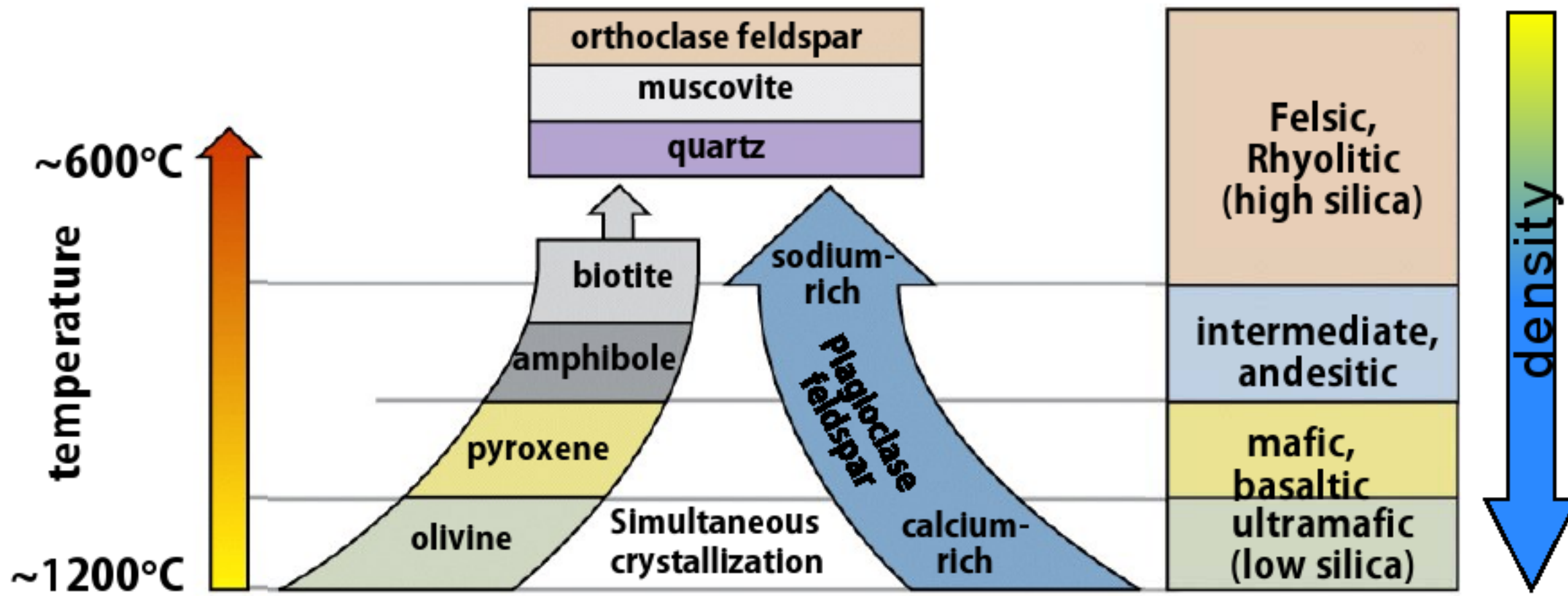
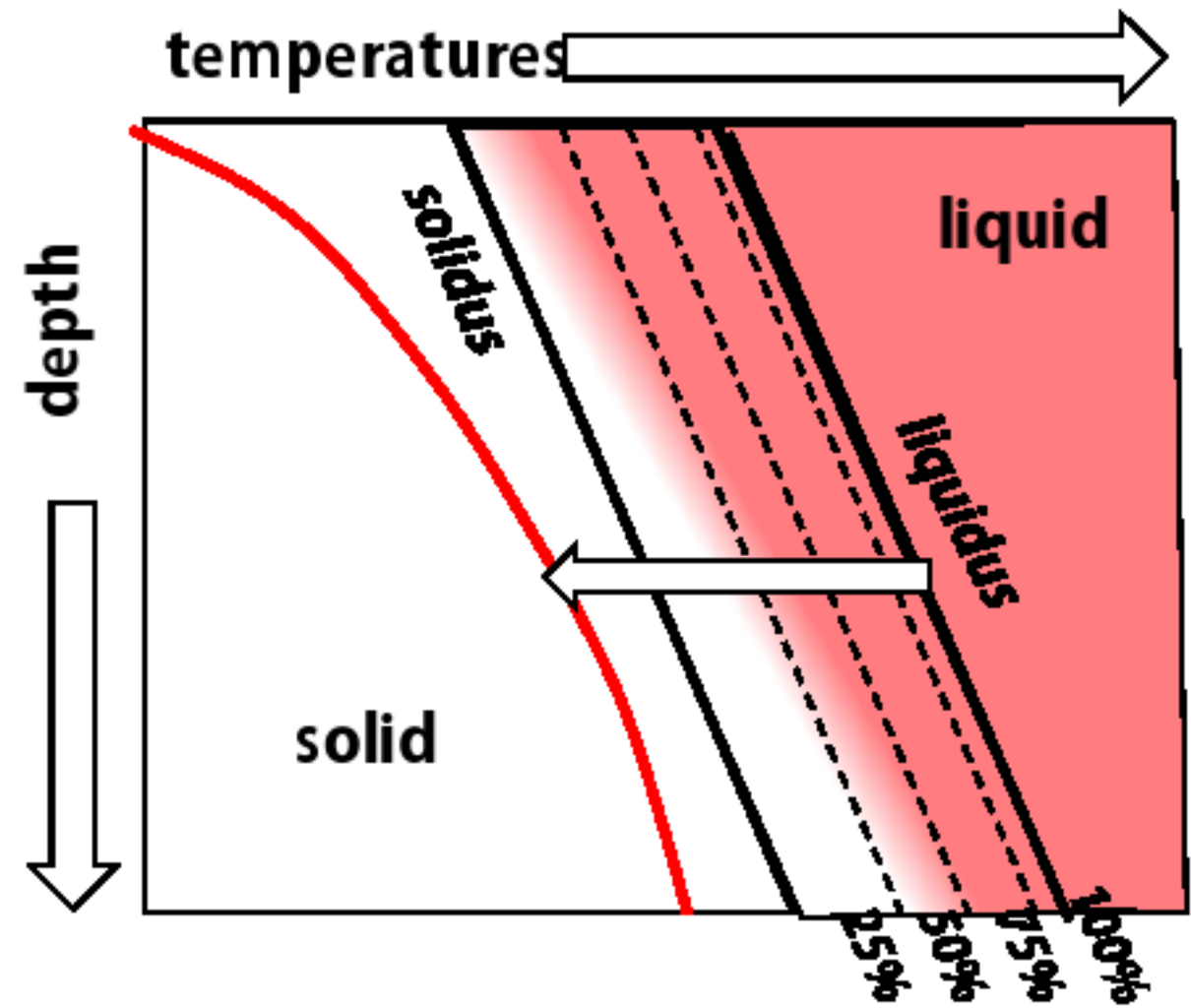
If it becomes too small, magma will stop somewhere on its way up, completely cool and solidify. An **intrusive** rock is born



Consolidation can happen in a variety of settings

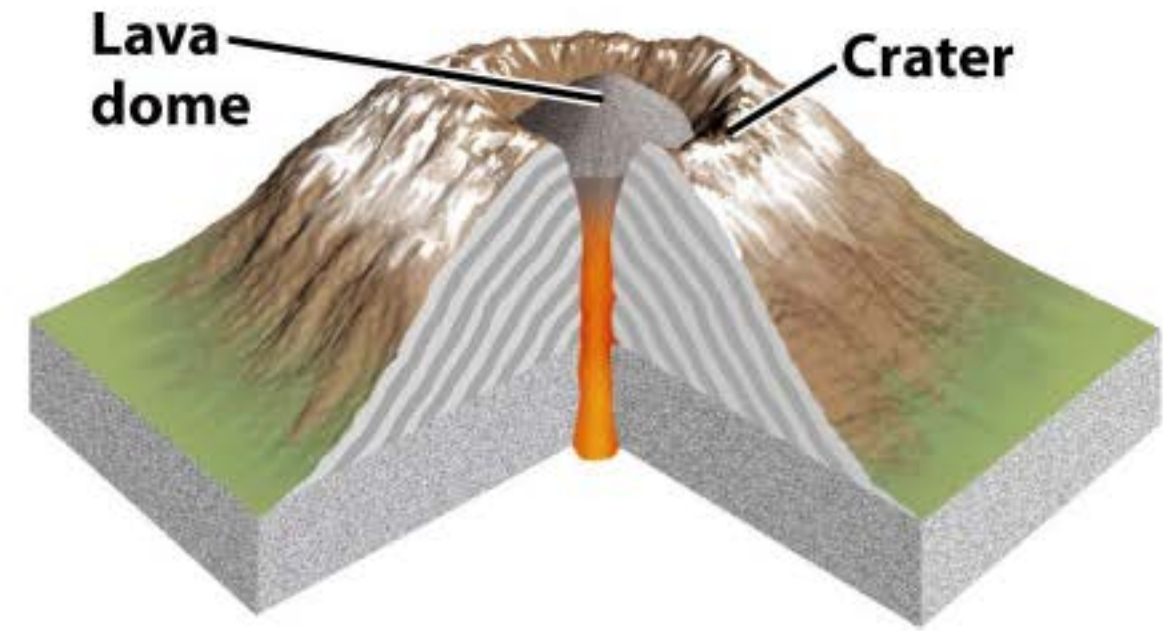
Different names are adopted

Magma starts cooling and solidifies
 Progressive cooling will cause precipitation of high-T (heavy) phases first and lower-T (lighter) later.
 Differentiation (segregation) takes place



Acid magmas

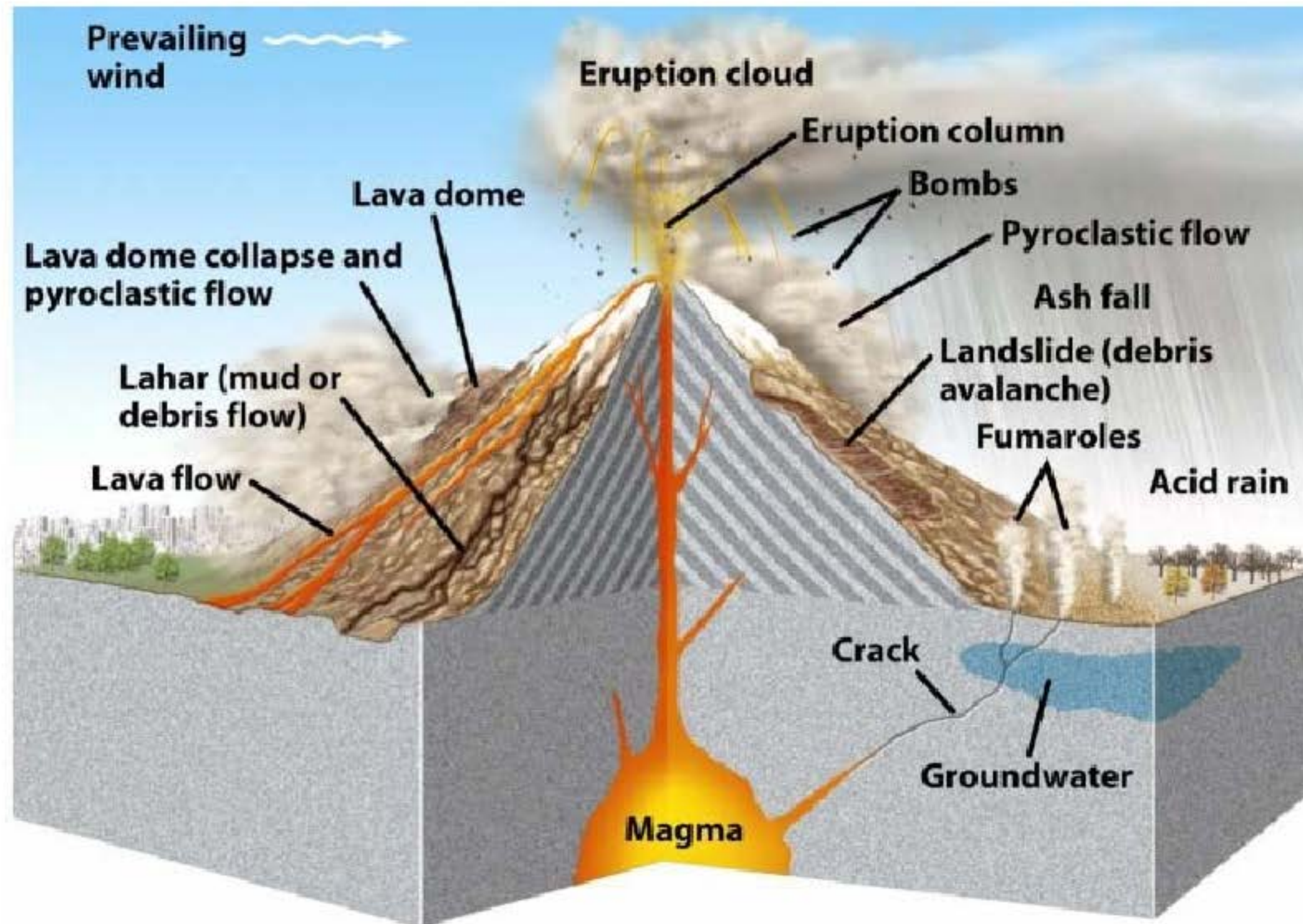
More acid magmas (>70% SiO₂) have lower temperatures (600-800°), are more viscous



Volcanoes are steep and aerially restricted

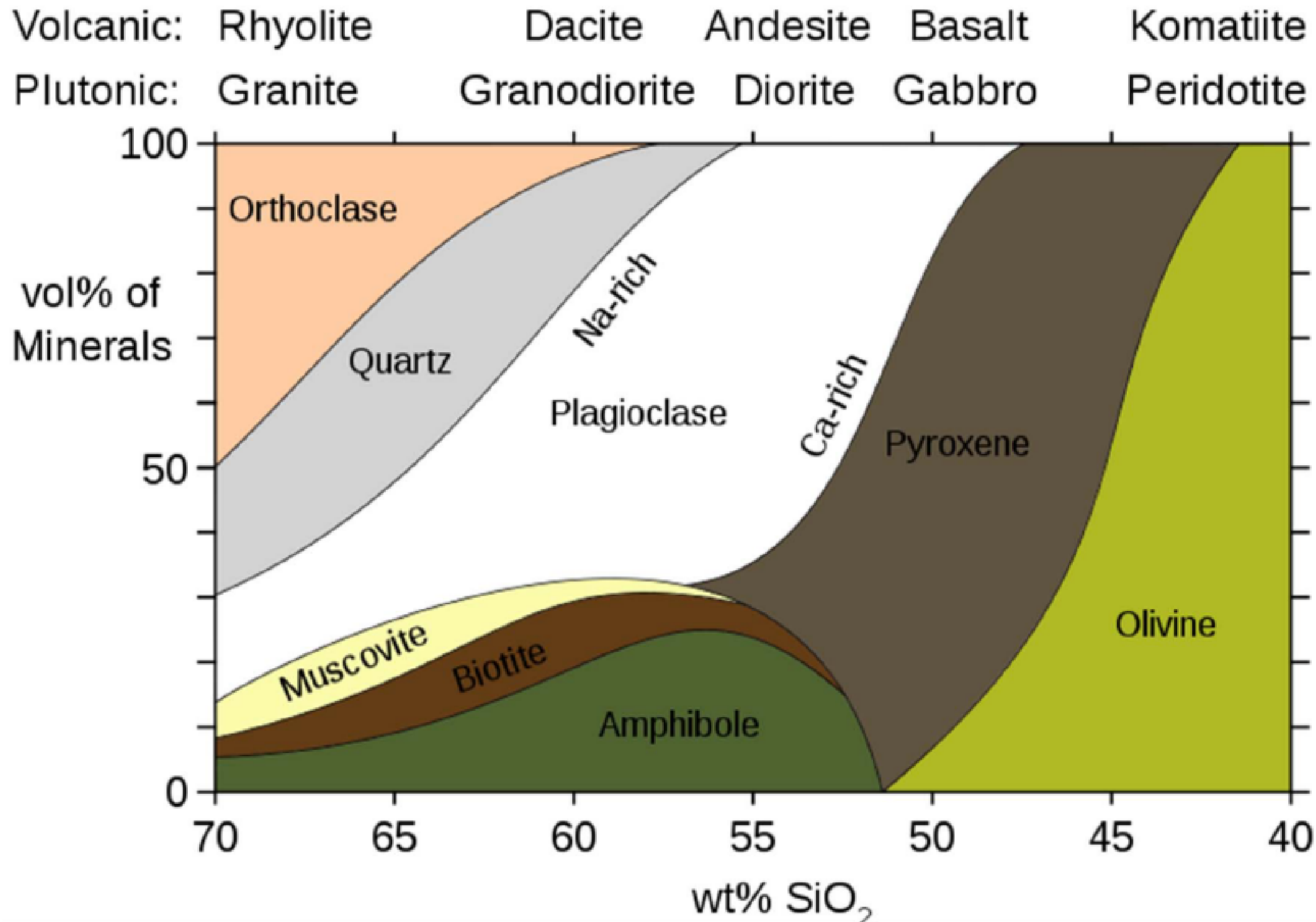
Eruptions are very explosive (thereby much more dangerous than basic volcanoes)

Volcanoes ejects not only lava but a lot of pyroclastic stuff (bombs, ashes...)



Magmatic rocks: classification

- Names of magmatic rocks depend on their mineralogy
- Different names are given to intrusive (plutonic) and extrusive (volcanic rocks)



Two rocks with the same composition but different names

Intrusive rocks

Typically coarse-crystalline (slow-cooling).

Extrusive rocks

Typically microcrystalline. Magma solidifies on the surface cooling rapidly, with insufficient time for crystals to grow



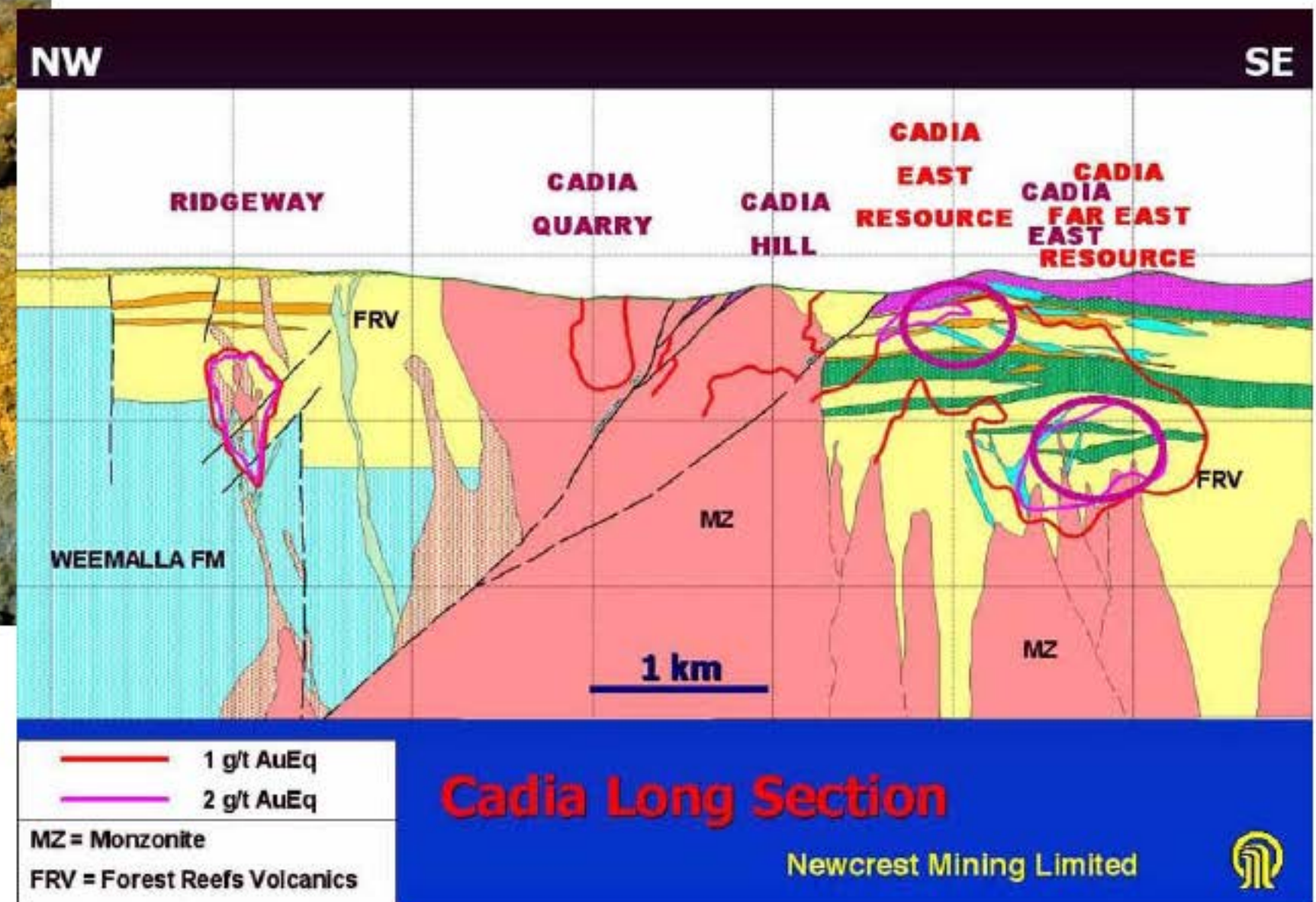
Some magmatism-related issues

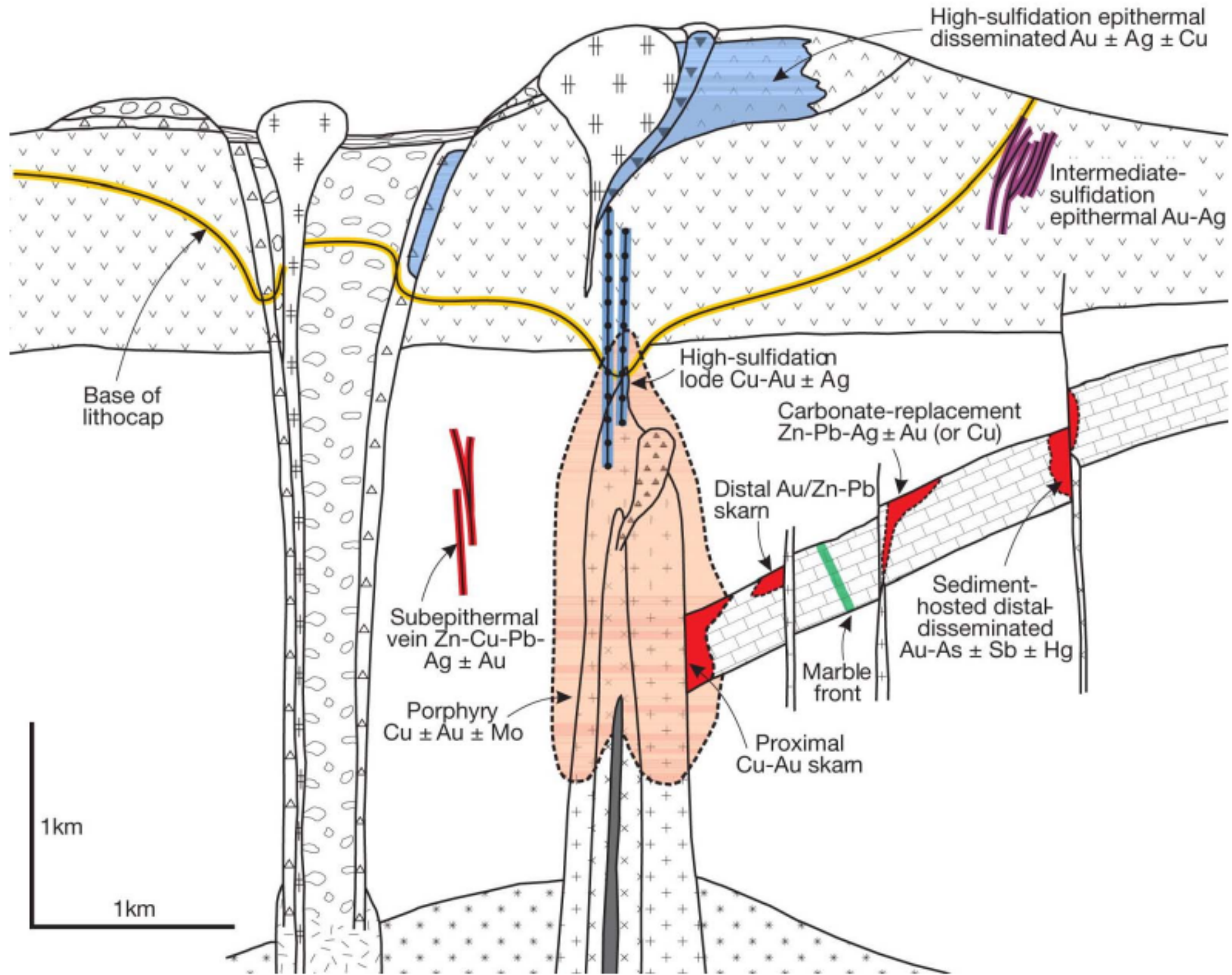
Magmatism is the major source of **heat** at shallow crustal levels and of hot, chemically-loaded **fluids**. Great economic importance

mineralization

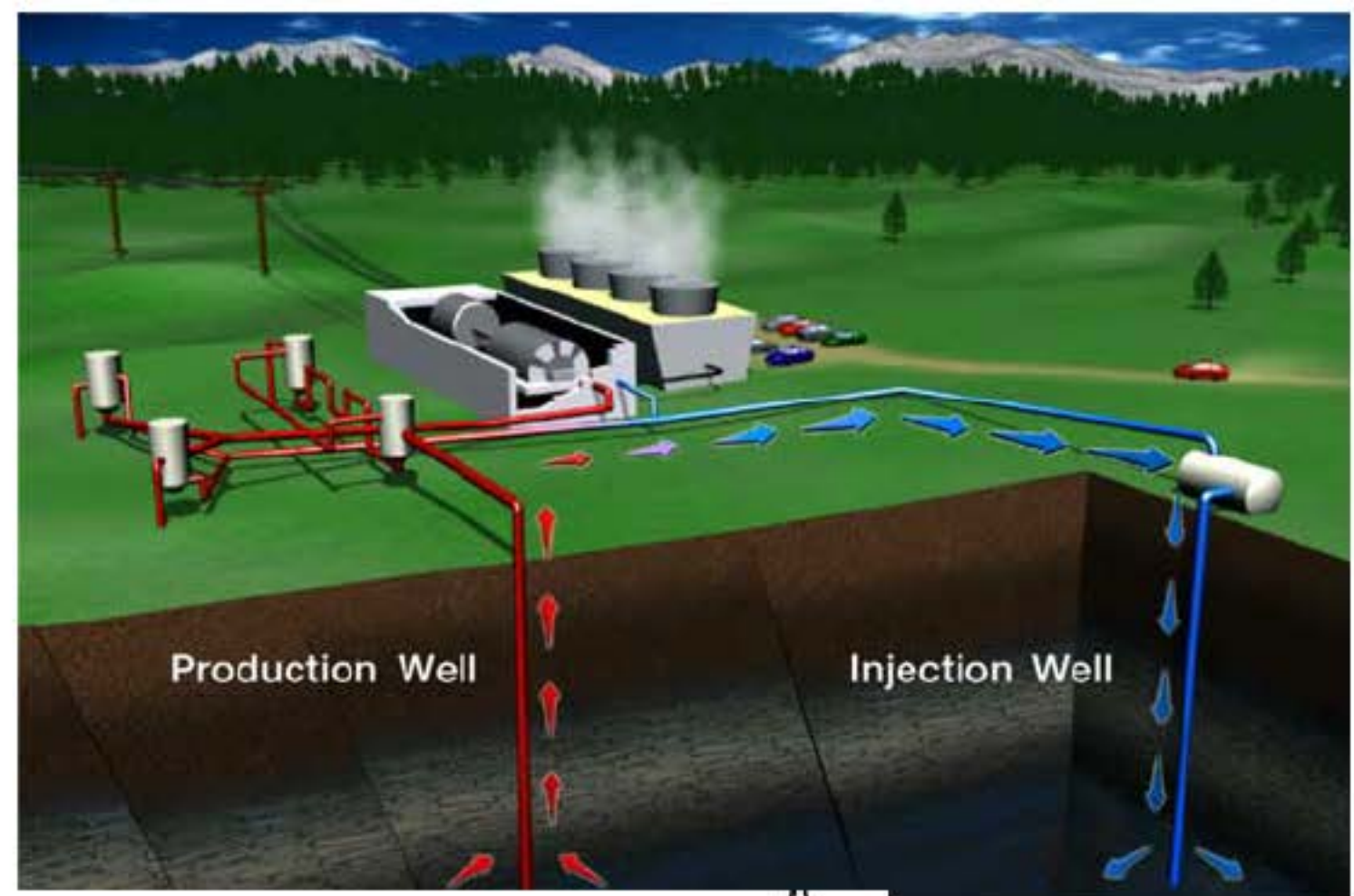


Porphyry coppers

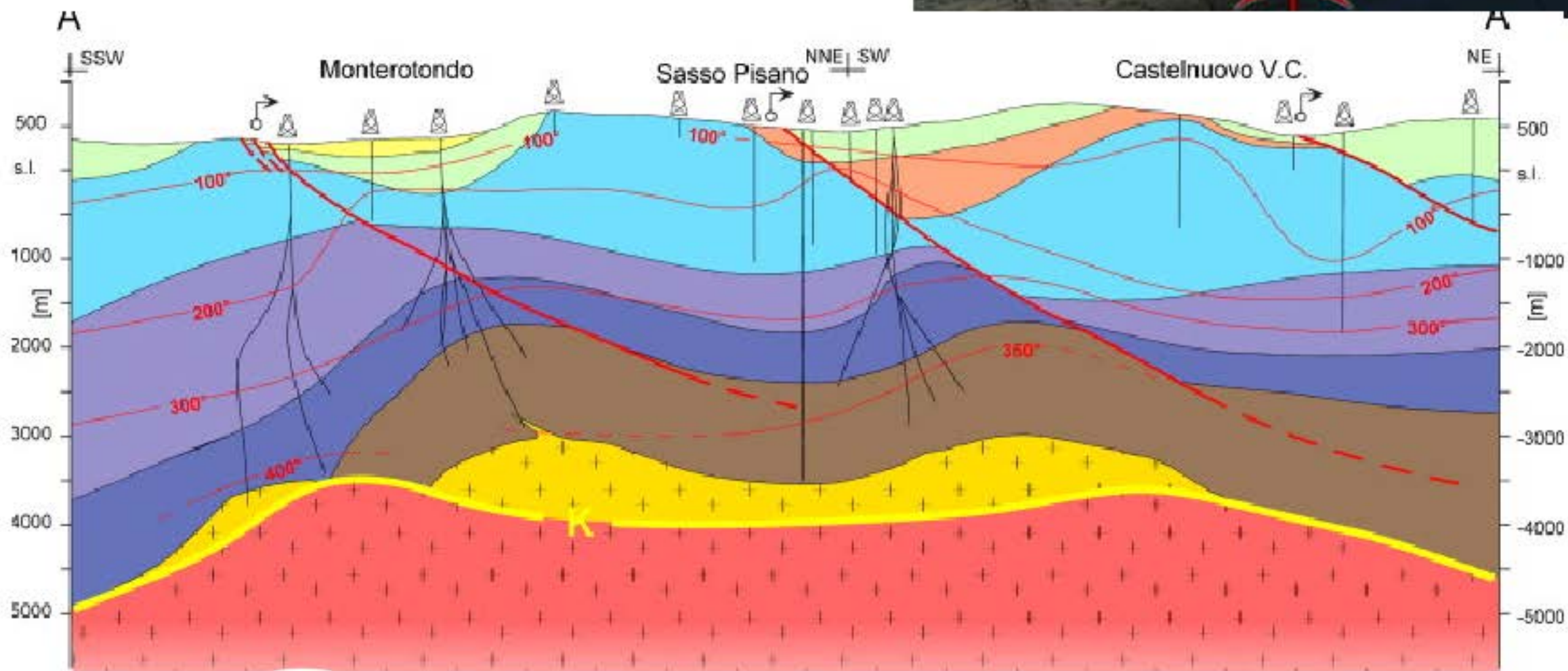




geothermal energy



The Larderello geothermal field



Sedimentary rocks

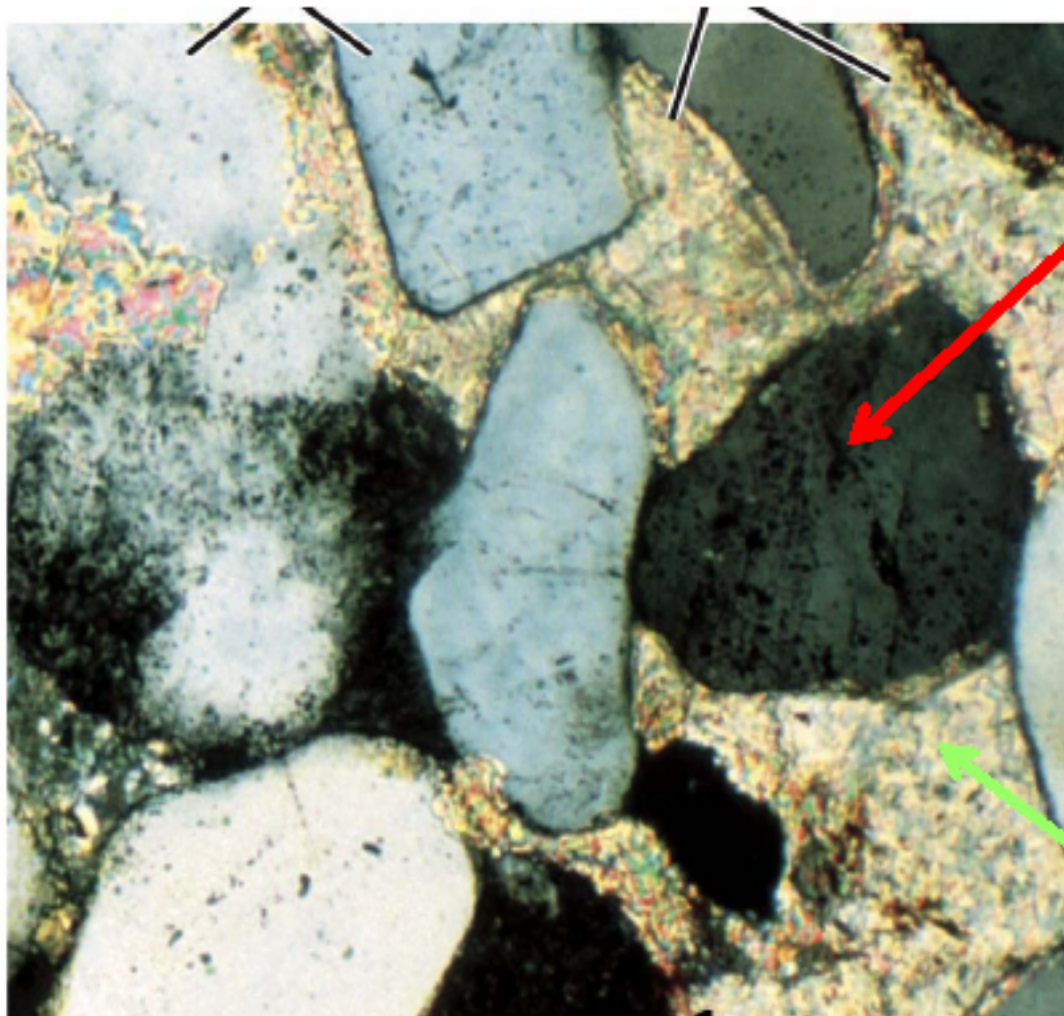


Rocks which form at the **surface** of the Earth (inclusive of **water masses**) through a) erosion of pre-existing rocks, b) life-related processes and c) purely chemical processes

Terrigenous rocks

Formed from the erosion and mobilization of older rocks

Composed of **grains** (=clasts) and (filled) **space** between them

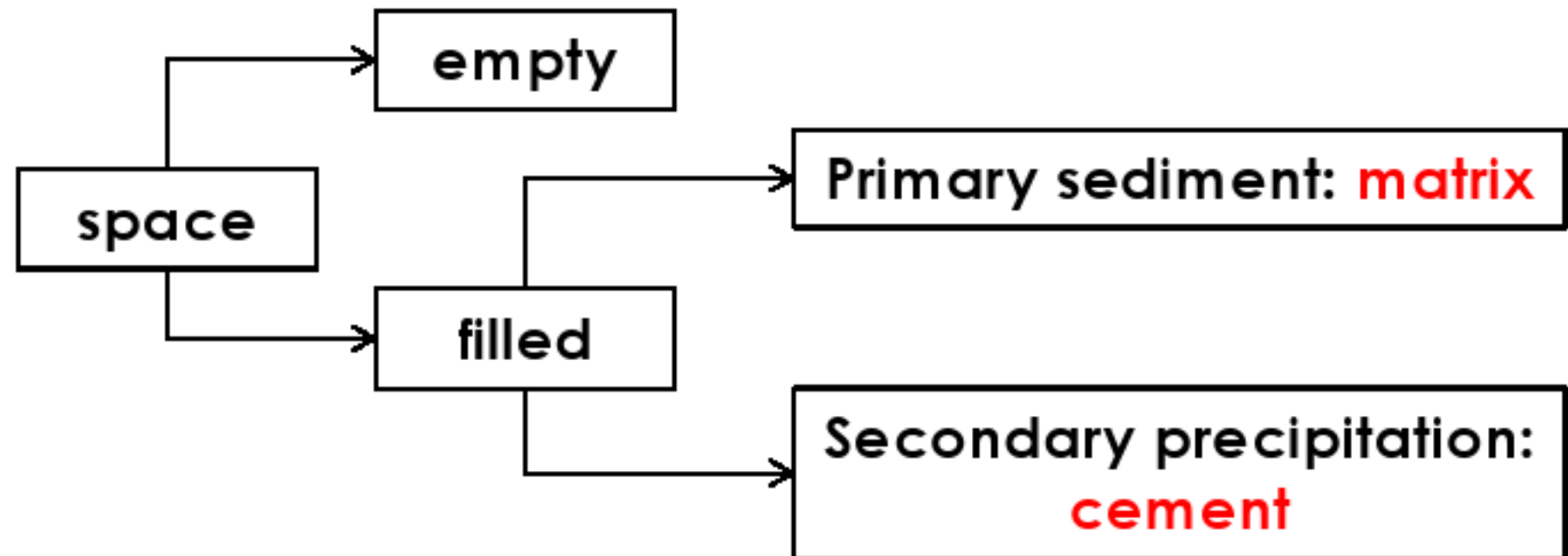


Clasts: typically pieces of rocks but can be also organic material etc

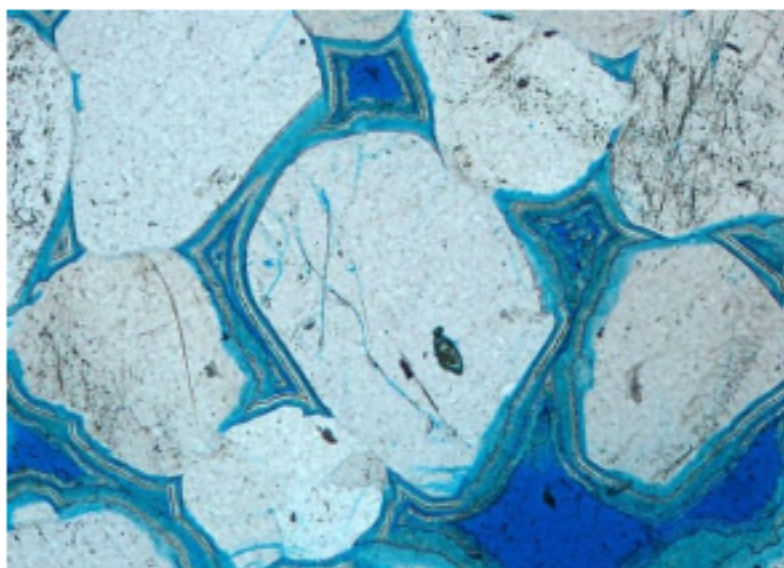
Interclast space

NB: clasts and matrix are **not** related to any absolute dimension!!

Space **between** clasts:



with matrix (= **finer-grained** portion of the primary sediment)



with cement (= newly precipitated minerals)

Defining and understanding matrix, cement etc is crucial for determining porosity and permeability

Classifying sedimentary rocks in terms of **clast** size

Particle Size	Sediment	Rock
COARSE Larger than 256 mm 256–64 mm 64–2 mm	GRAVEL Boulder Cobble Pebble	Conglomerate
MEDIUM 2–0.062 mm	SAND	
FINE 0.062–0.0039 mm	MUD Silt	Siltstone
Finer than 0.0039 mm	Clay	<ul style="list-style-type: none"> Mudstone (blocky fracture) Shale (breaks along bedding) Claystone

- Rock names are on the basis of the size of the clasts not of the matrix, which will be finer-grained (**no absolute size implied**)
- The **composition** of the clasts is **irrelevant**



The **composition of grains** can also be added

quartzitic sand (quartzarenite)

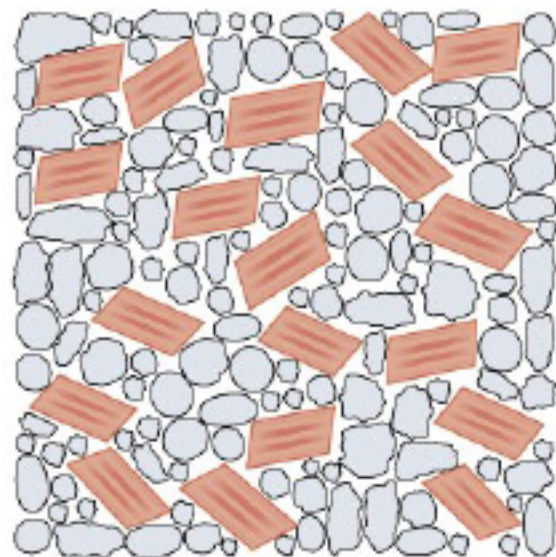
- feldspar rich sand

-

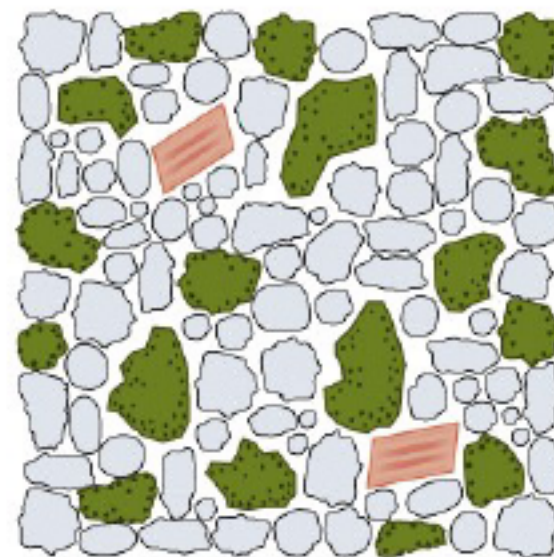
-

and some traditional names

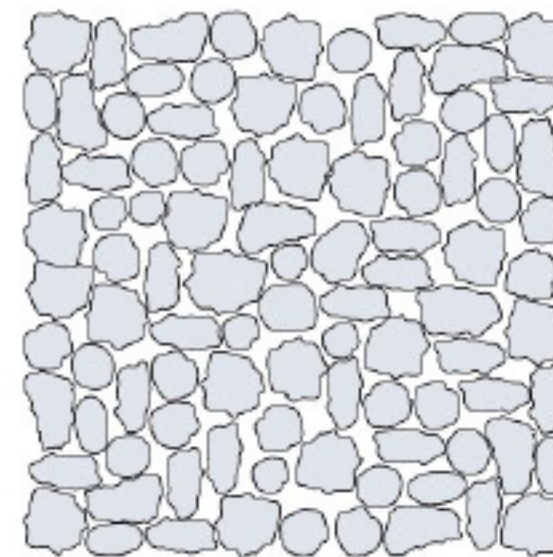
Arkose:
feldspar-rich



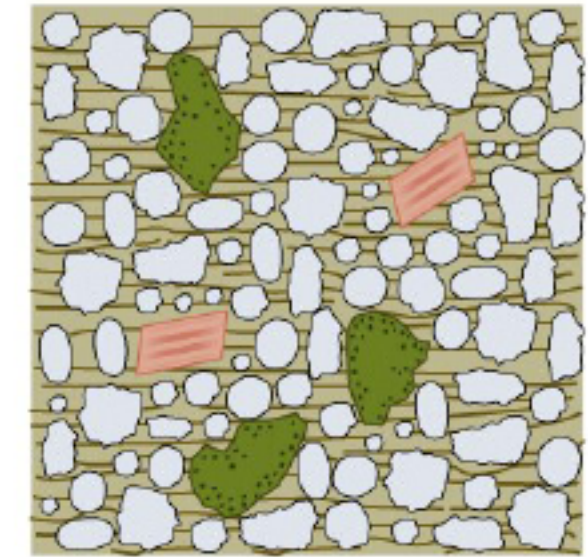
Lithic sandstone:
rock fragment-rich



Quartz arenite:
pure quartz



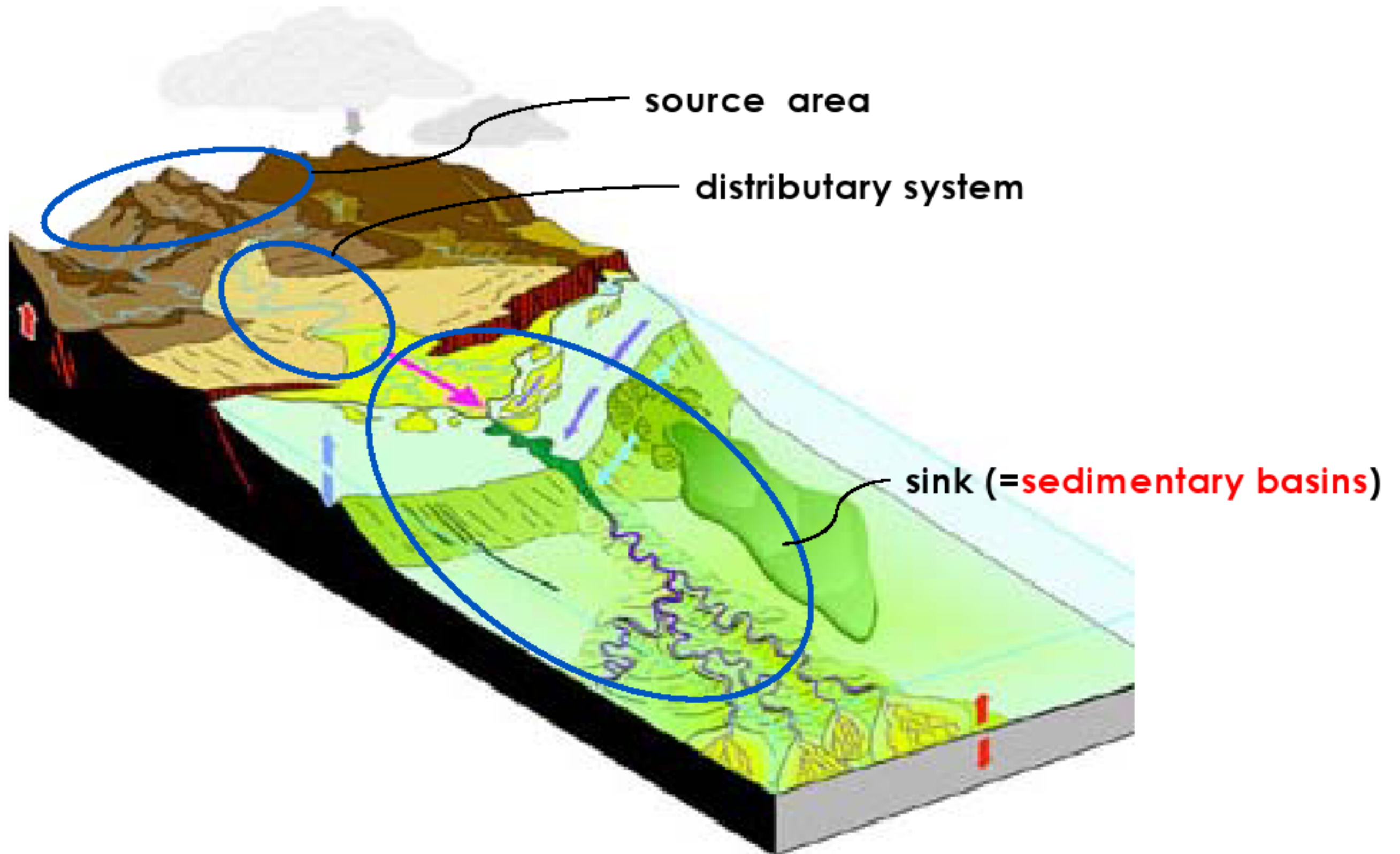
Graywacke:
matrix-rich



1 mm



The **source-to-sink** approach



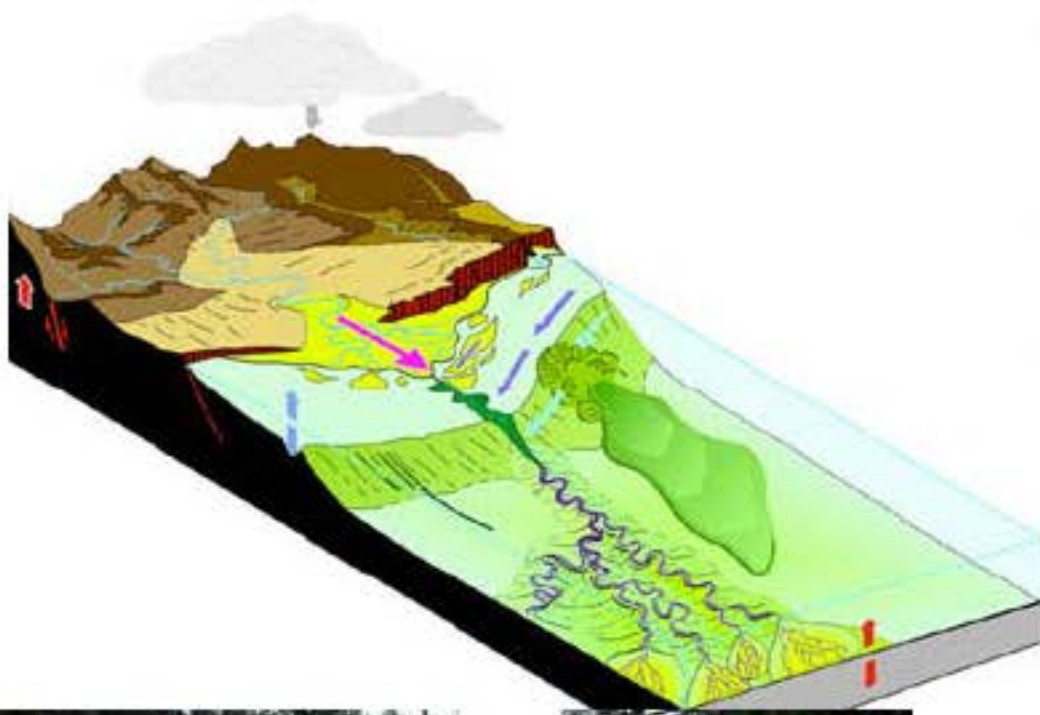


The source area

Here it is decided **how much** and **what type** of sediments is produced, depending on

- kind of rocks exposed
- Amount of uplift/exhumation
- climate

Note: all these **change** through time



The distributary system

Gravity, water, wind etc will remove the sediments from the production area and transport them until they find a place to rest (the **sink**)

Rivers are a major part of the distributary system



The Rhine



The Mississippi

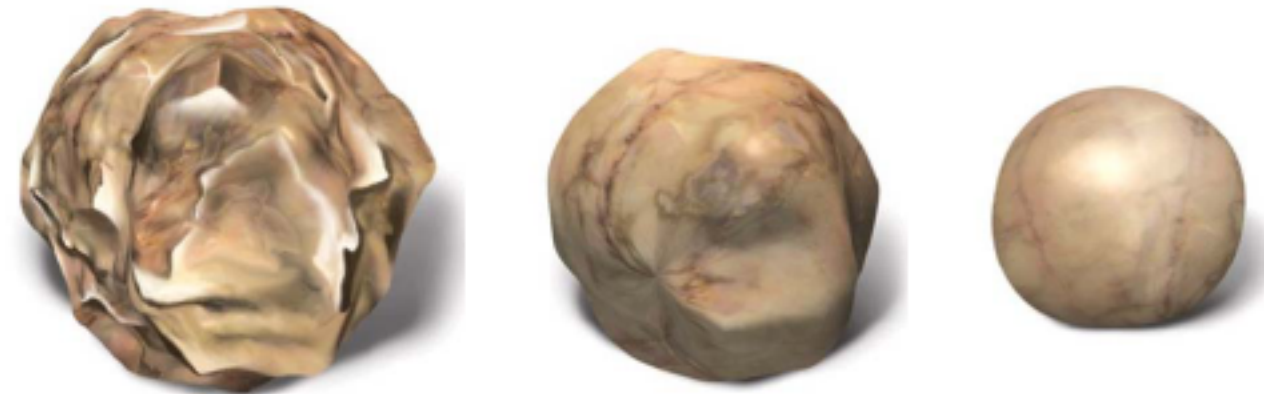
Transport and movement can modify the sediment delivered to the sink
(=sediment characteristics in the sink depend also on transport)

Mechanical modifications

Movement (collisions) **rounds** clasts

Movement (collision) breaks particles making them smaller

Currents select grains on the basis of their weight (volume and density)



Chemical changes

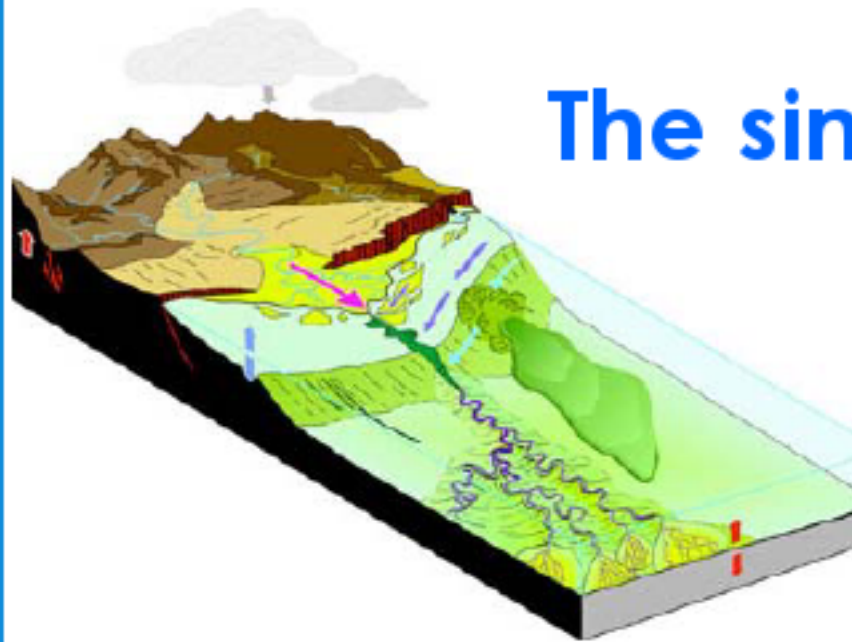
affect clasts during transport (dissolution etc)

In an **immature** rock, clasts are not rounded, poorly sorted and of different types



In a **mature** rock, clasts are rounded, well sorted and of one dominant type





The sink:

A place where the transport agents can dump their sediments

=

a place where **accommodation space** is available

The best accommodation space is between the top and bottom of a water column (sea better than lake)
(sediments dumped above sea level are very unstable)

sea level

sea floor



1

water column

2

sediments

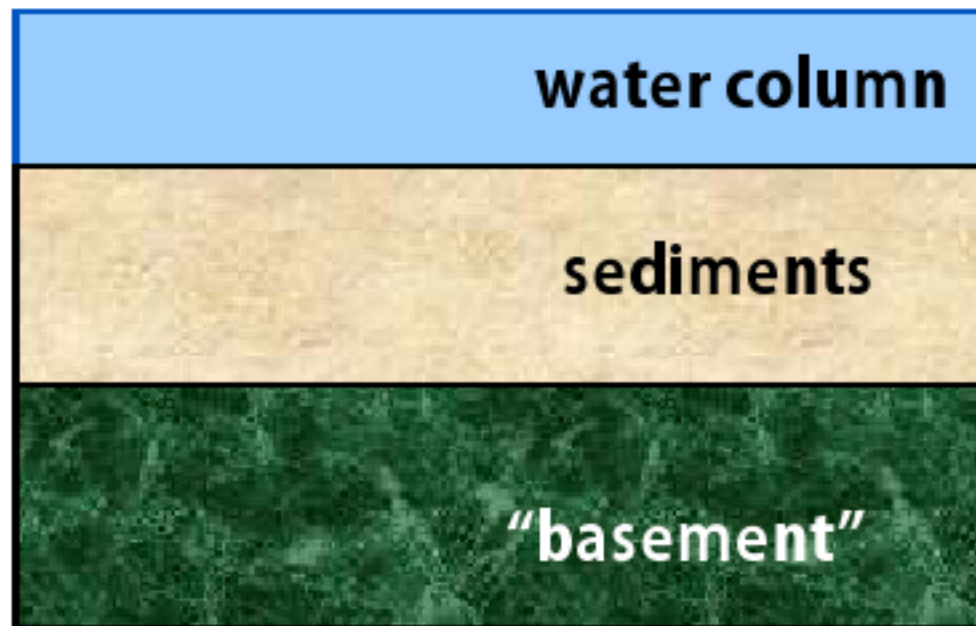
3

"basement"

A first layer of sediment is deposited

Three relevant surfaces

- 1) sea level
- 2) sea floor
- 3) bottom of the sedimentary succession



1

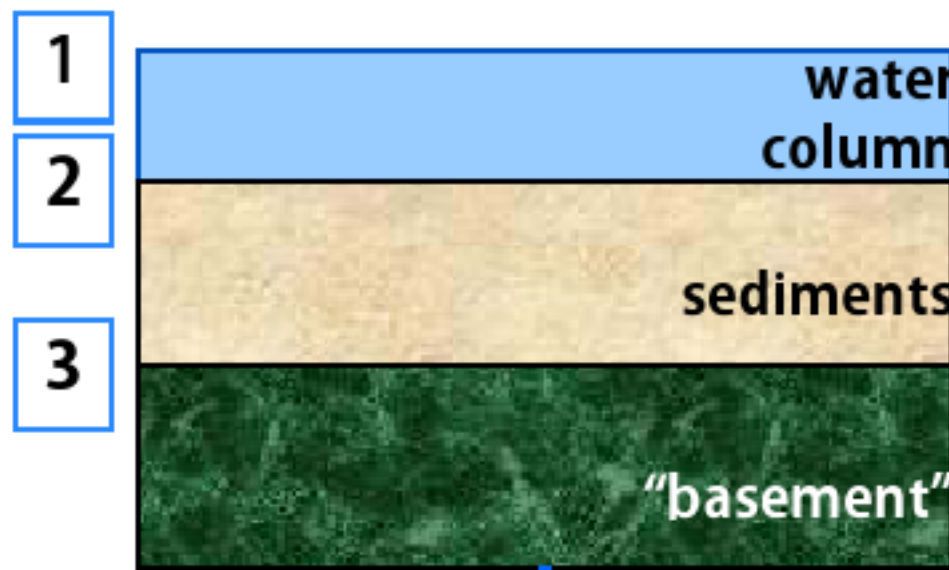
Controlled by global sea level changes

2

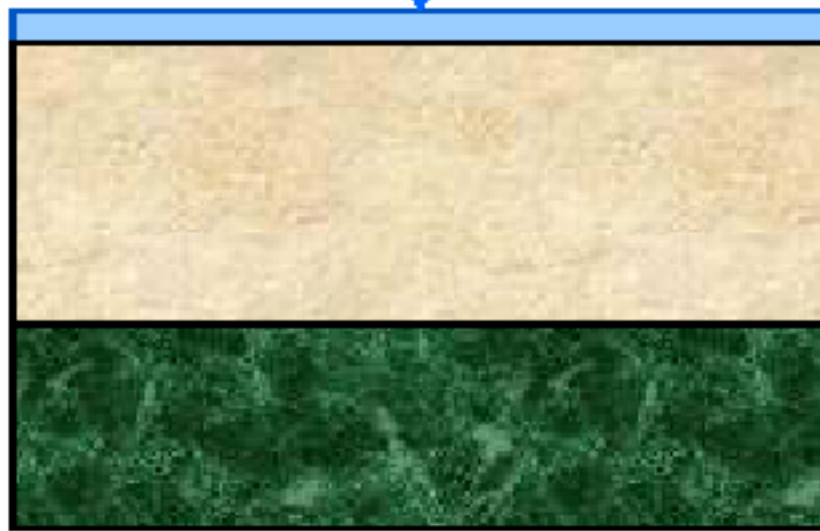
Controlled by the amount of sediments. Controls the **depositional environment**, also called **facies** (shallow vs deep water)

3

Controlled by **tectonics**
Controls the **thickness** of the deposited sediments



If 3 does not move, the water column will be progressively filled



If 3 moves downward (subsidence) a lot of sediments can be accumulated



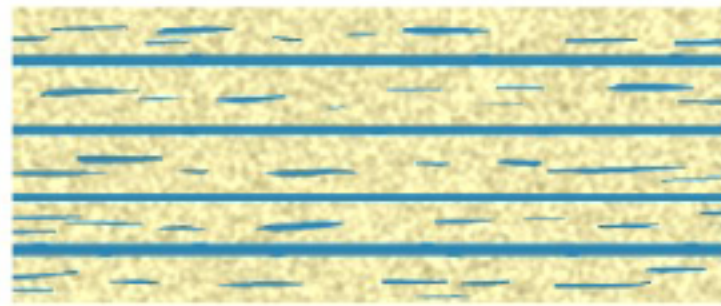
- the water column=maximum thickness of sediments
- the facies of the sediment will change through time

- Thicknesses can be very large (kms)
- Facies do not change necessarily

Transforming loose sediments in **rocks**

During subsidence, sediments will be gradually covered by other, newly arrived sediments and, slowly but surely, will become rocks (lithification).

Compaction



50–60% water

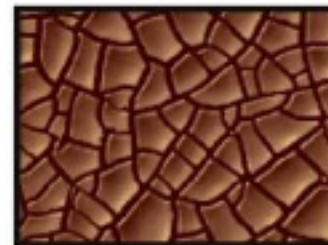


10–20% water



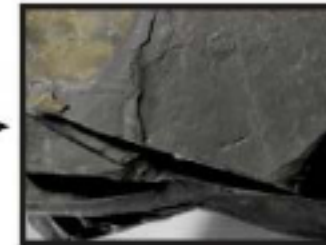
Lithification

Mud



Pressure

Shale



Silt and siltstone, mudstone and shale, clay and claystone

Sand



Pressure

Sandstone



Gravel



Pressure

Conglomerate



cementation can provide an important contribution to lithification



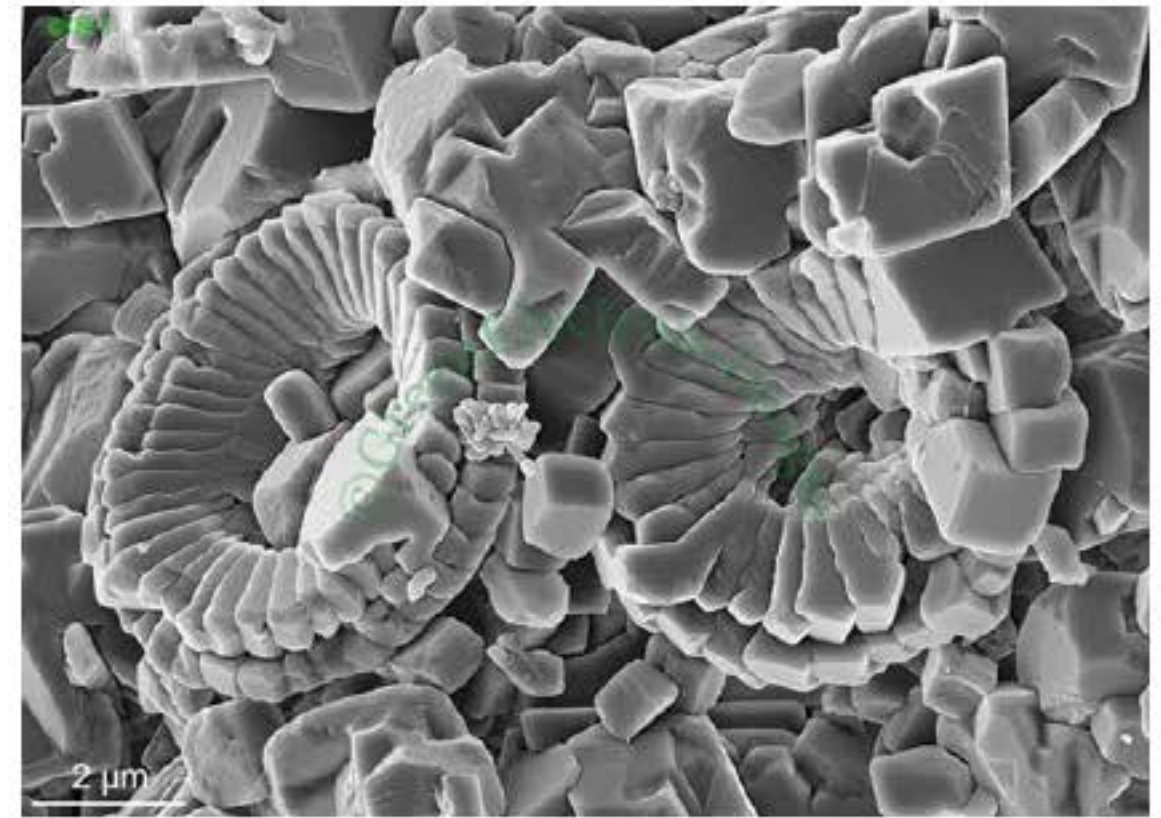
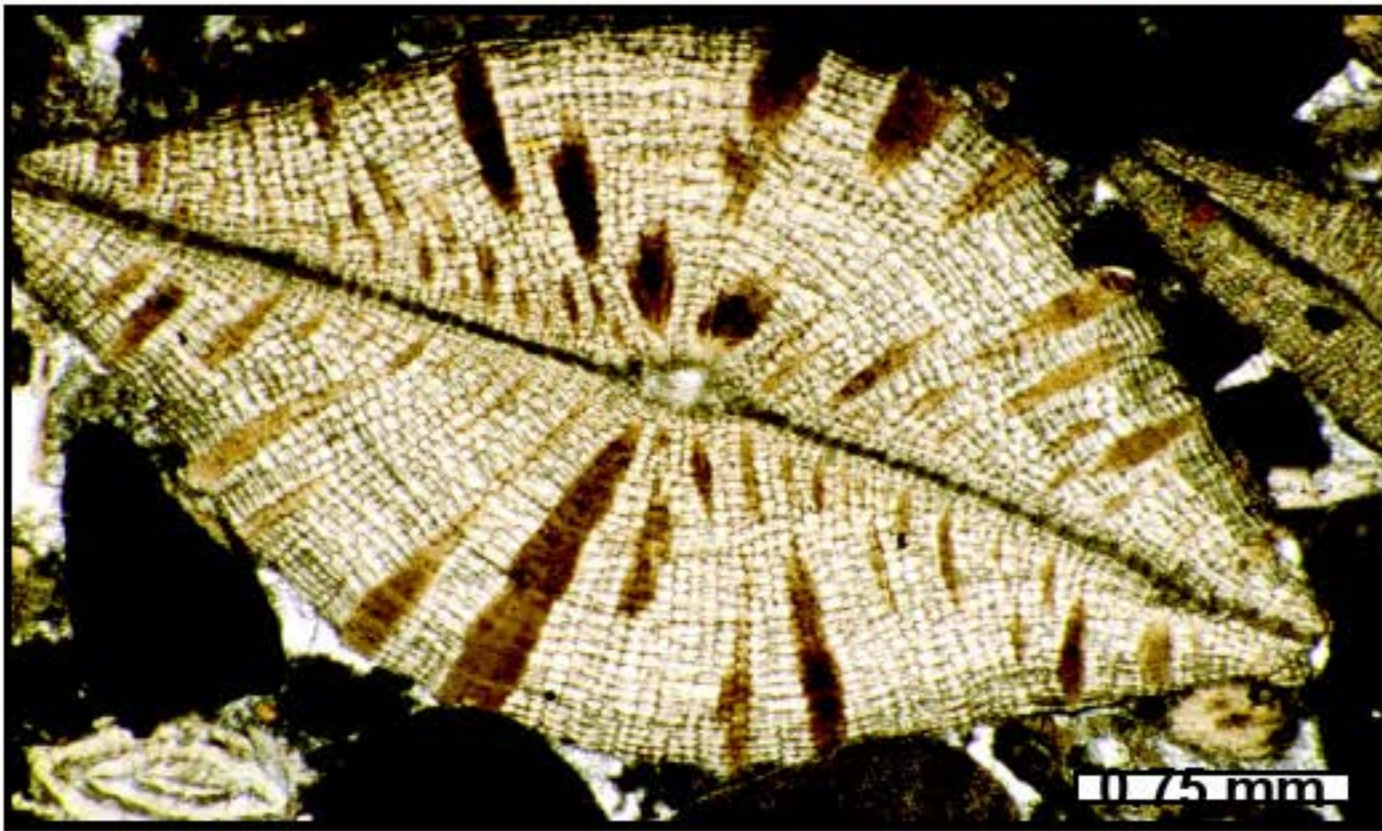
Biological sedimentary rocks (fossilized remnants of plants and/or animals, chemical precipitates)

Carbonates (limestones and dolomites) are
by far the most important

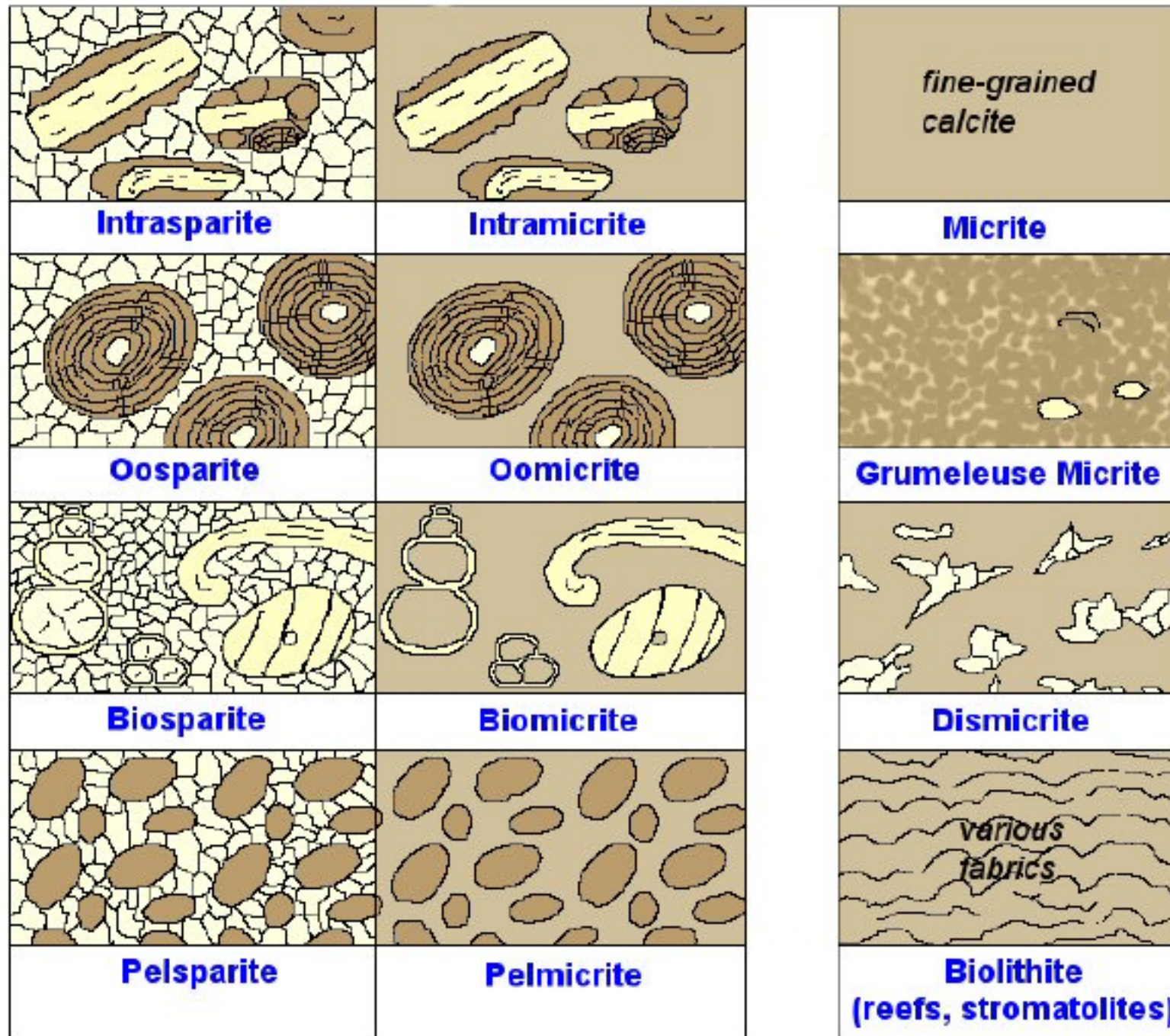


Sediment	Rock	Chemical Composition	Minerals
BIOLOGICAL Sand and mud (primarily bioclastic)	Limestone	Calcium carbonate (CaCO_3)	Calcite (aragonite)
Siliceous sediment	Chert	Silica (SiO_2)	Opal, chalcedony, quartz
Peat, organic matter	Organics	Carbon compounds; Carbon compounded with oxygen and hydrogen	(coal), (oil), (gas)

Carbonates are generated by life!



Carbonates have their special classification schemes



more will follow

Folk's Classification of Limestones - a modified version based on Folk (1962). Coarser varieties can be classified with "rudite" endings. E.g. - biosparrudite, biomicrudite. Grumeleuse micrite is clotted micrite, common in lagoonal facies. Like pelmicrites and pelsparites it may have ostracod valves. Ian West & Tonya West (c) 2005.

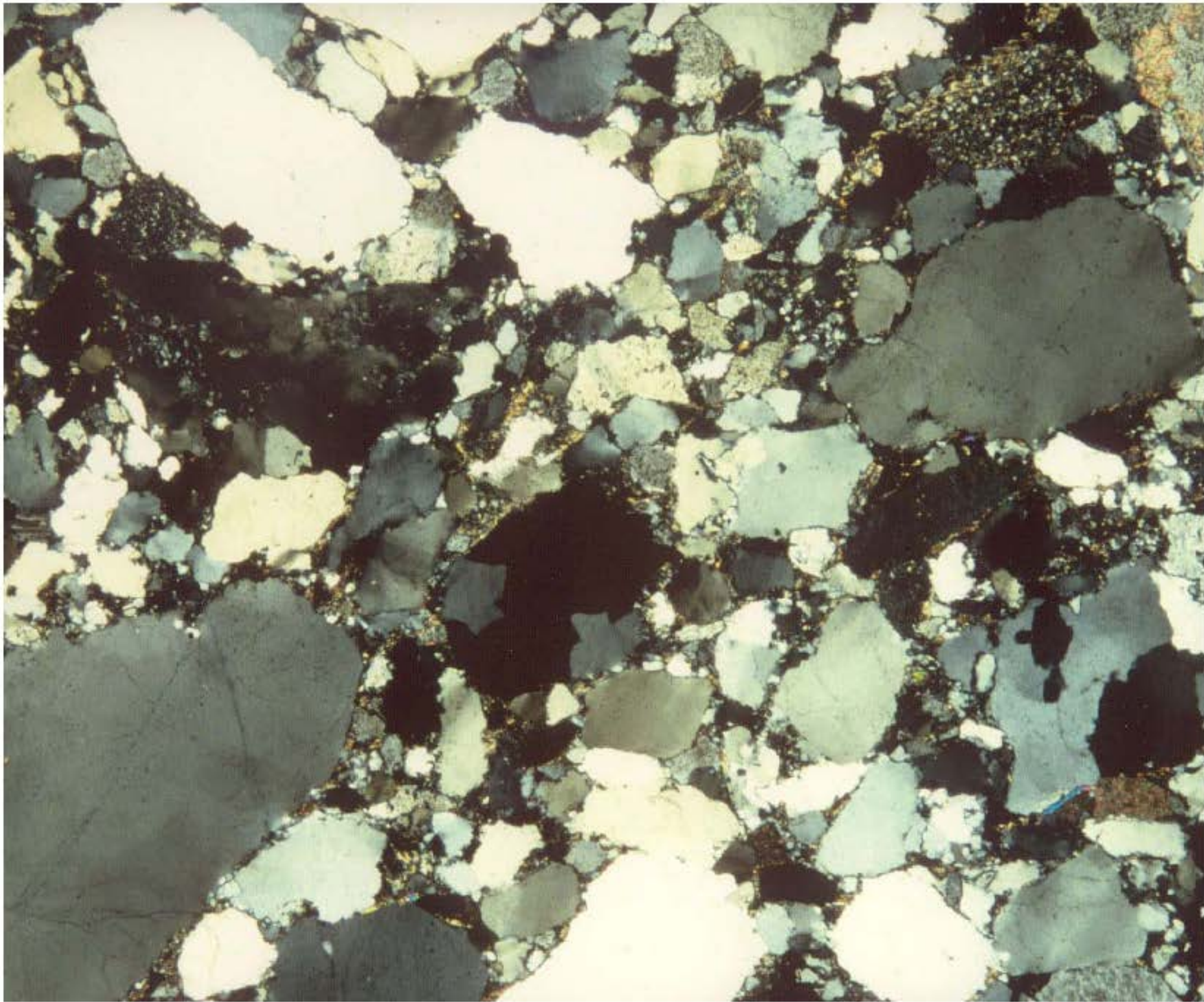
Chemical sedimentary rocks

Typical examples are evaporites, sediments formed by progressive evaporation of sea water.



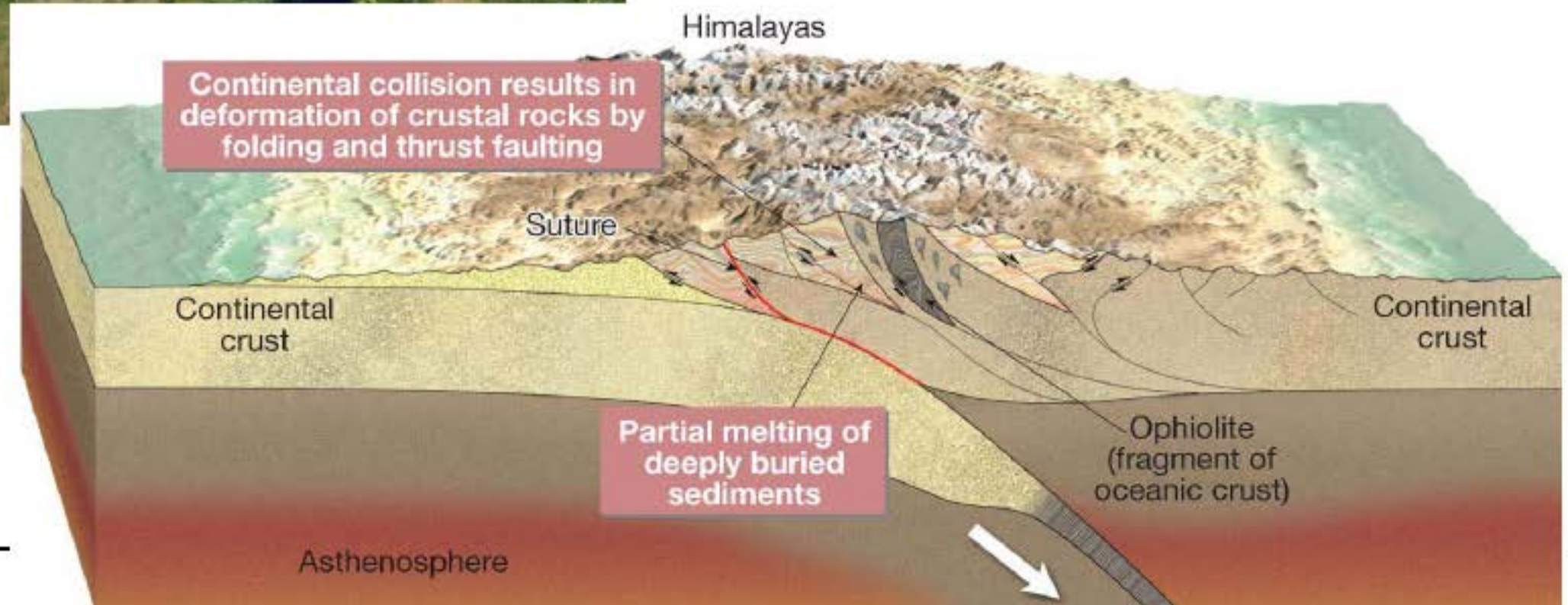
Sediment	Rock	Chemical Composition	Minerals
CHEMICAL No primary sediment (formed by diagenesis)	Dolostone	Calcium-magnesium carbonate (CaMg[CO₃]₂)	Dolomite
Iron oxide sediment	Iron formation	Iron silicate; oxide (Fe ₂ O ₃); limonite, carbonate	Hematite, siderite
Evaporite sediment	Evaporite	Sodium chloride (NaCl); calcium sulfate (CaSO ₄)	Gypsum, anhydrite halite, other salts

The formation of many “chemical” rocks is stimulated by organic activity!

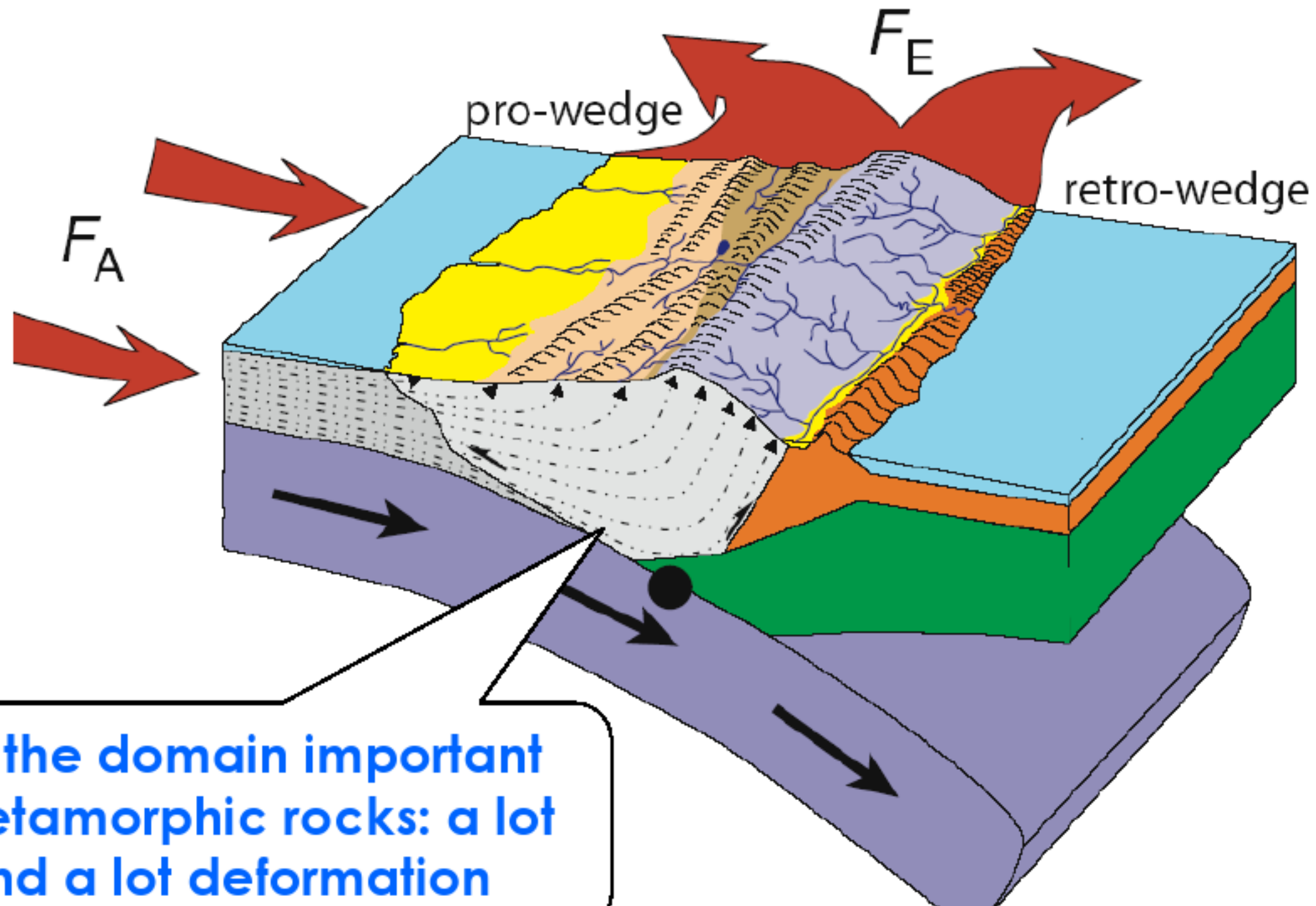


Metamorphic rocks

develop when parent (sedimentary, magmatic, metamorphic) rocks have been subjected to higher-than-normal pressure and/or temperatures

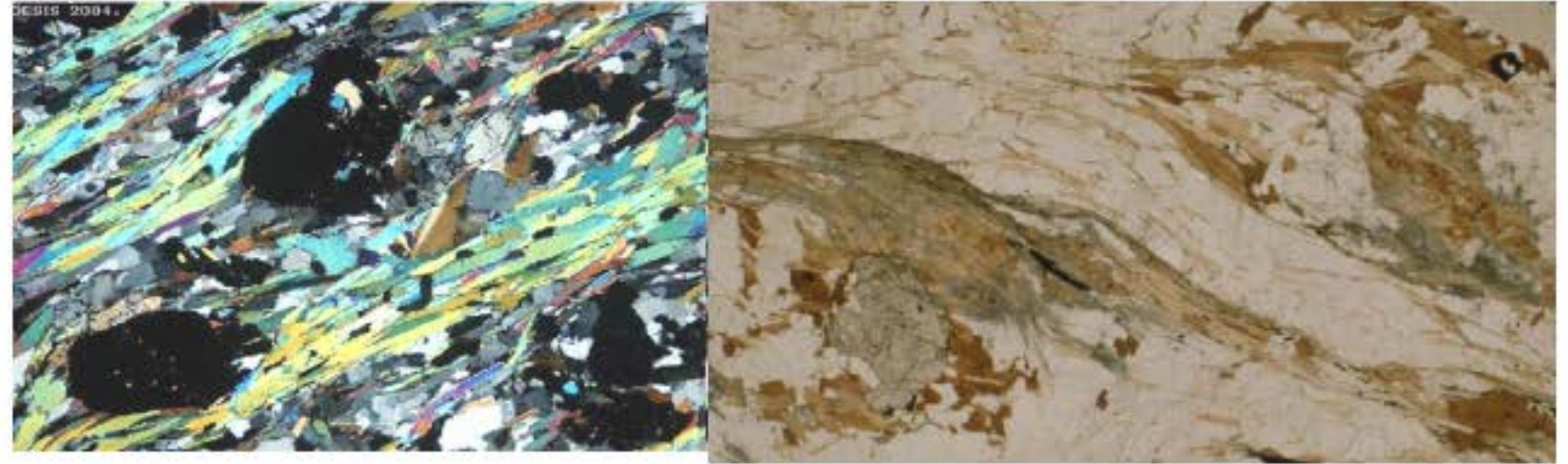


The way orogens work

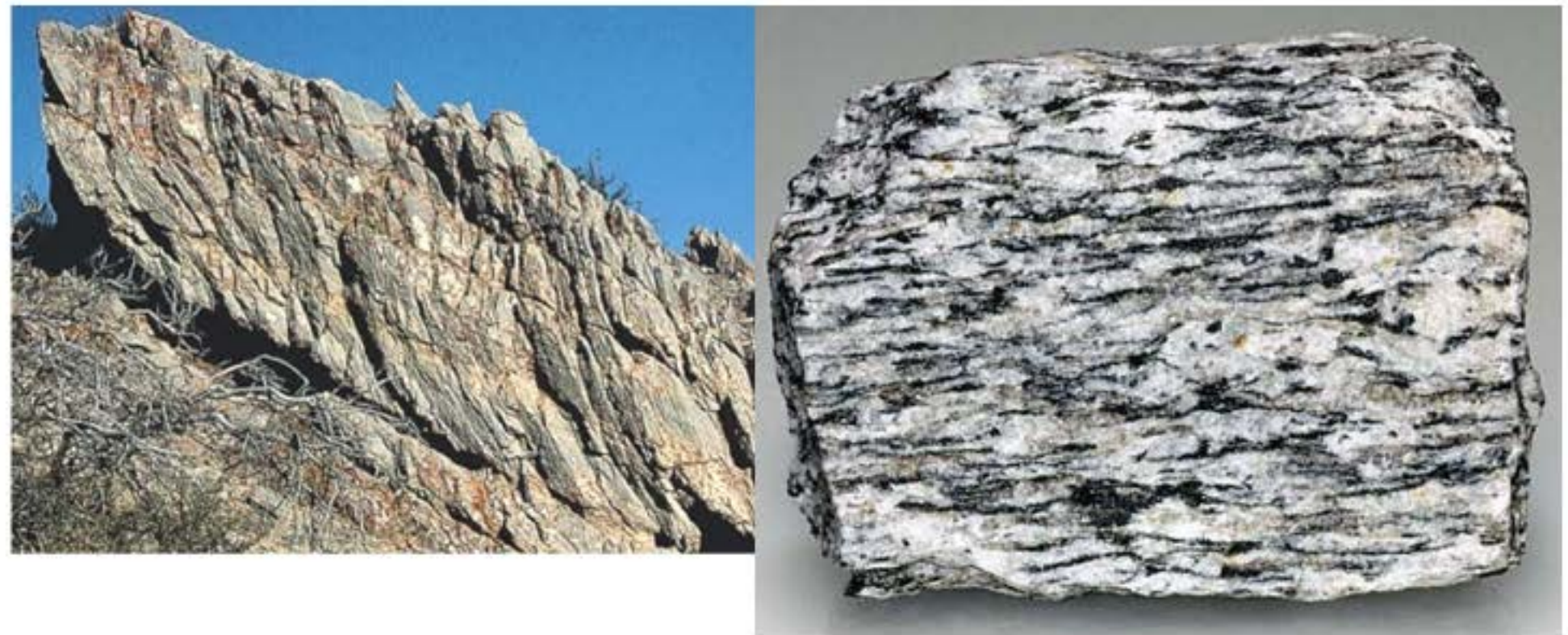


When **physical conditions** (pressure and/or temperature) change:

Minerals become unstable, and react with each other creating new minerals



The rock will be deformed producing **foliations** and **lineations** (texture) and resulting in a completely different rock

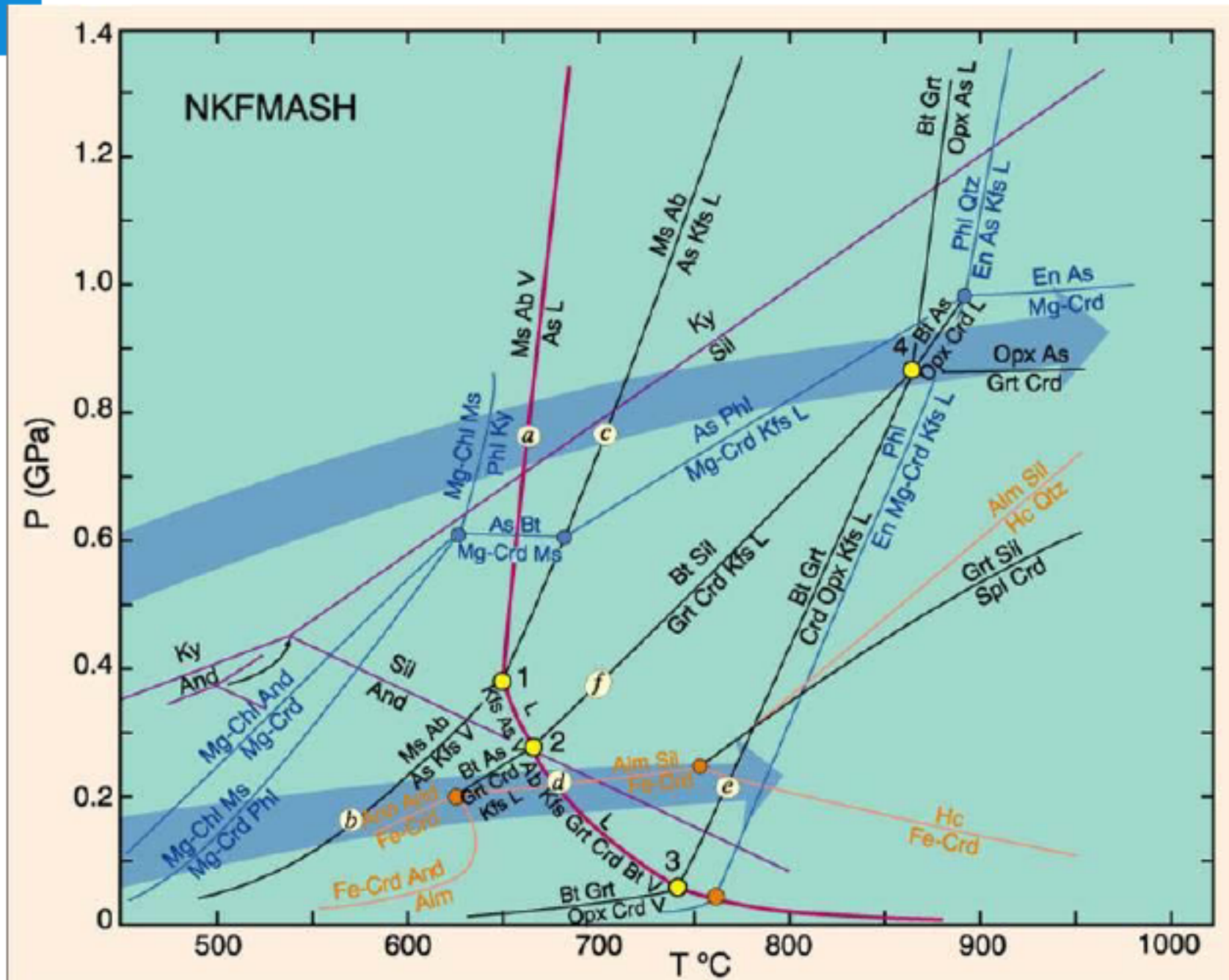


Mineralogical changes take place when rocks pass from one stability field to the other.

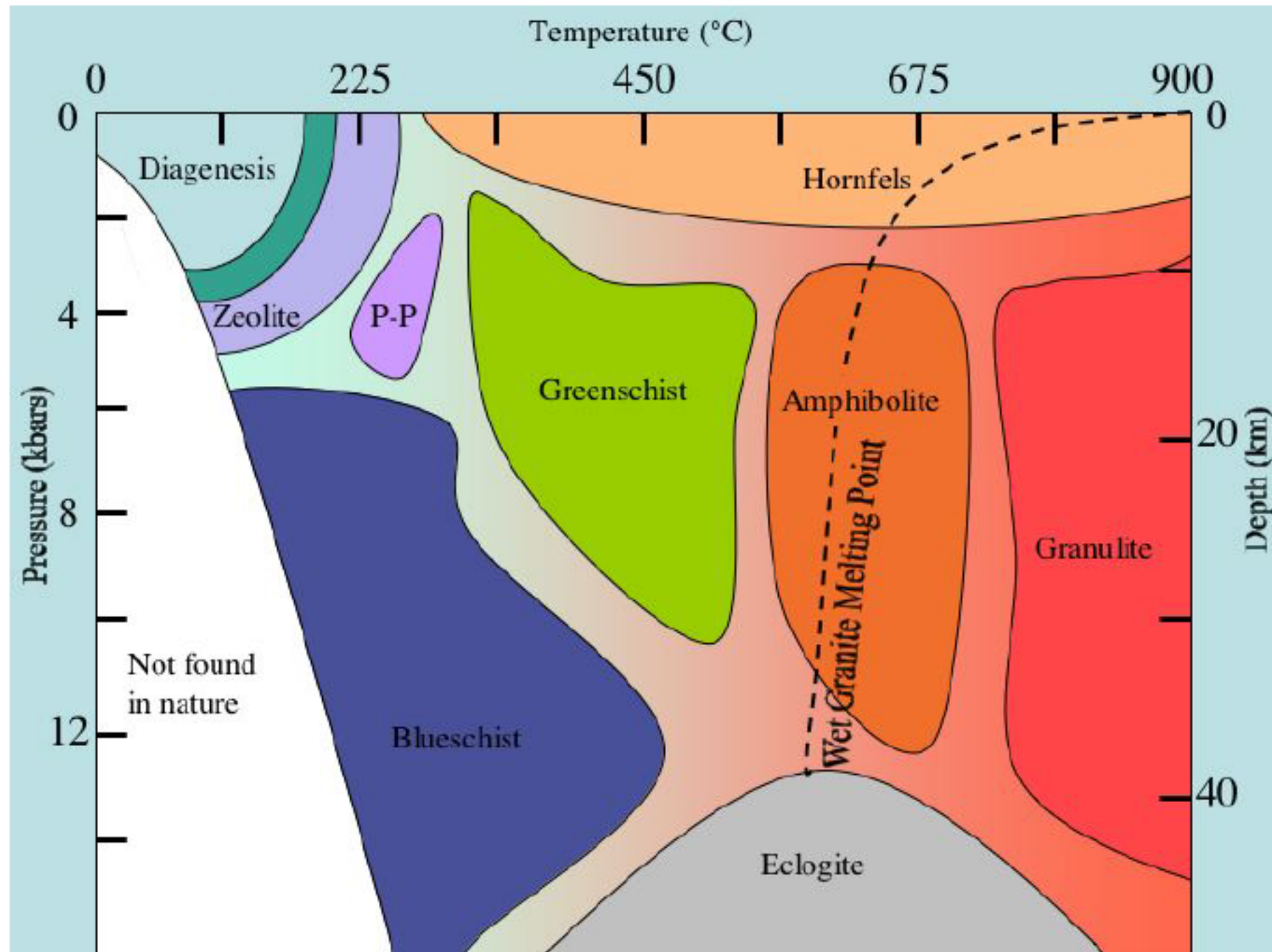
Experimental petrology provides information on the pressure, temperature conditions at which the mineral changes take place



In this way one can link the mineralogical composition of a rock to the p and T conditions of formation

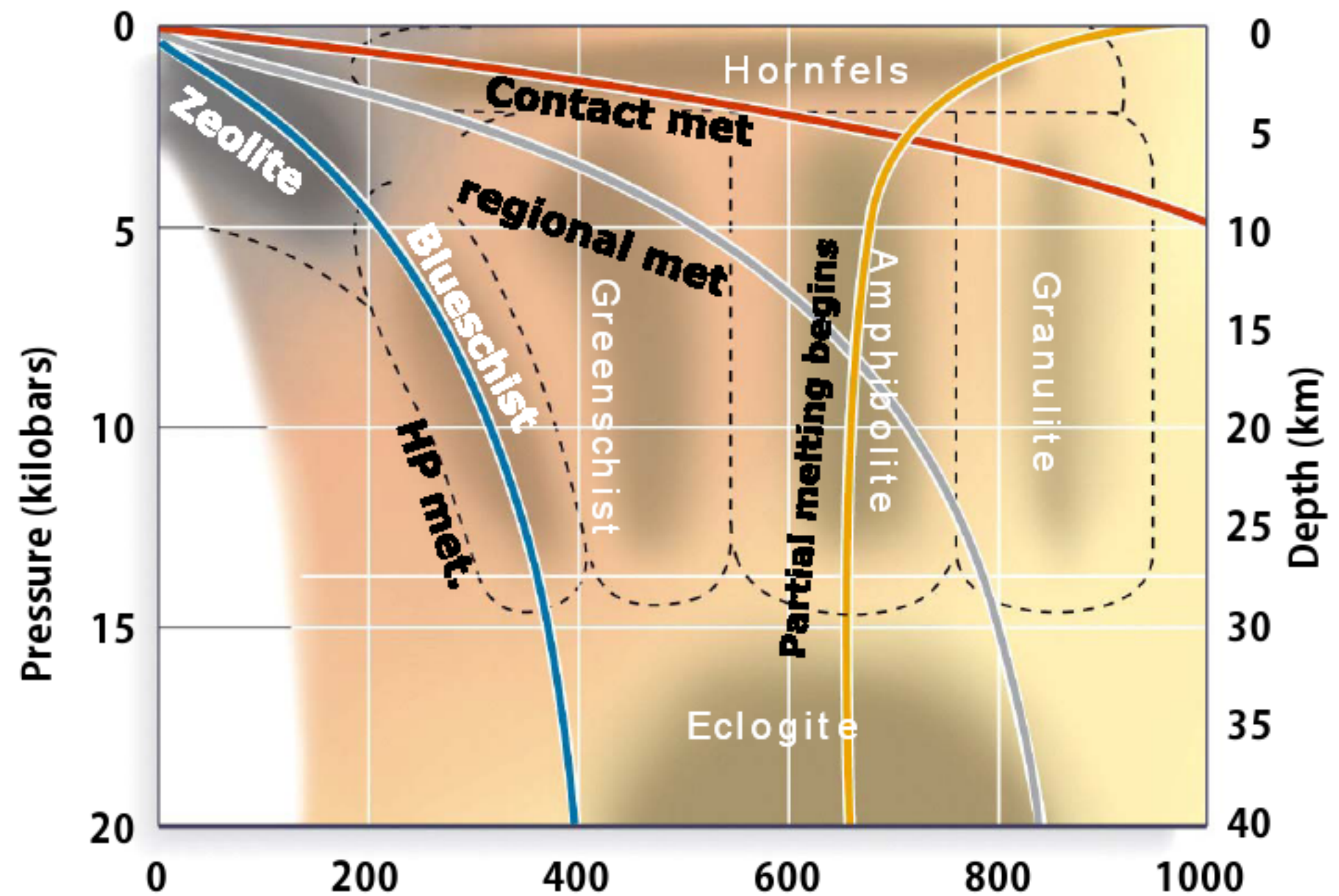


On the basis of the mineral assemblages, **metamorphic facies** are defined



Three typical gradients:

- HT/LP when temperatures are higher than what predicted by the **regional gradient**



- The regional gradient (**Barrovian metamorphism**)
- HP/LT when temperatures are lower than what predicted by the regional gradient (**high pressure metamorphism**)

Rocks following these paths differ from each other

Resulting rocks depend, obviously, also on the primary rock (**protolith**)

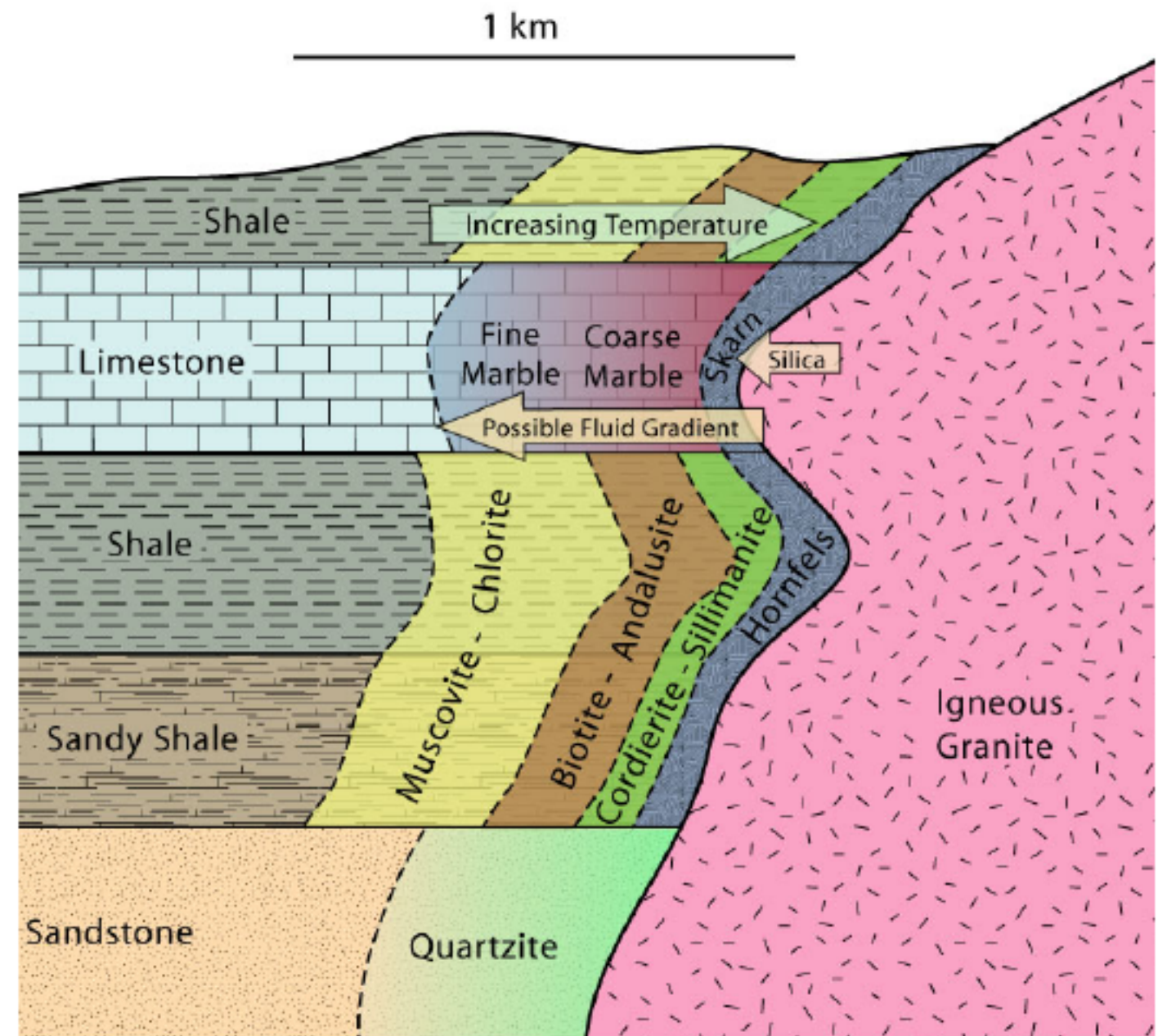
Contact metamorphism (HT/LP)

Basically associated with a magmatic intrusion

- Contact metamorphic rocks tend to preserve their primary aspect (massive or layered)
- The size of crystals increases with increasing temperatures (=towards the intrusion)
- Fluid circulation is strong and profoundly impacts the rock mineralogy (ore deposits)



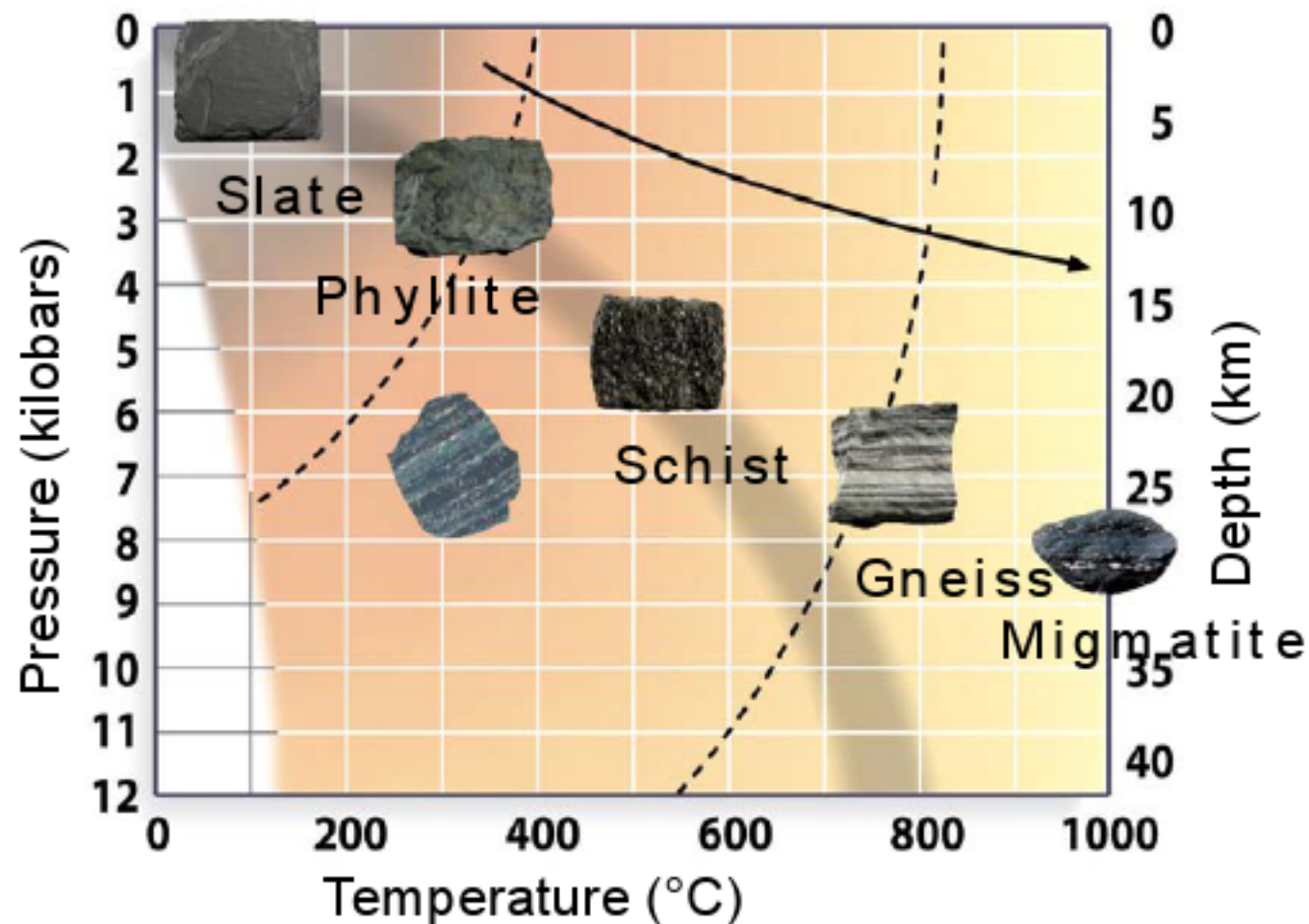
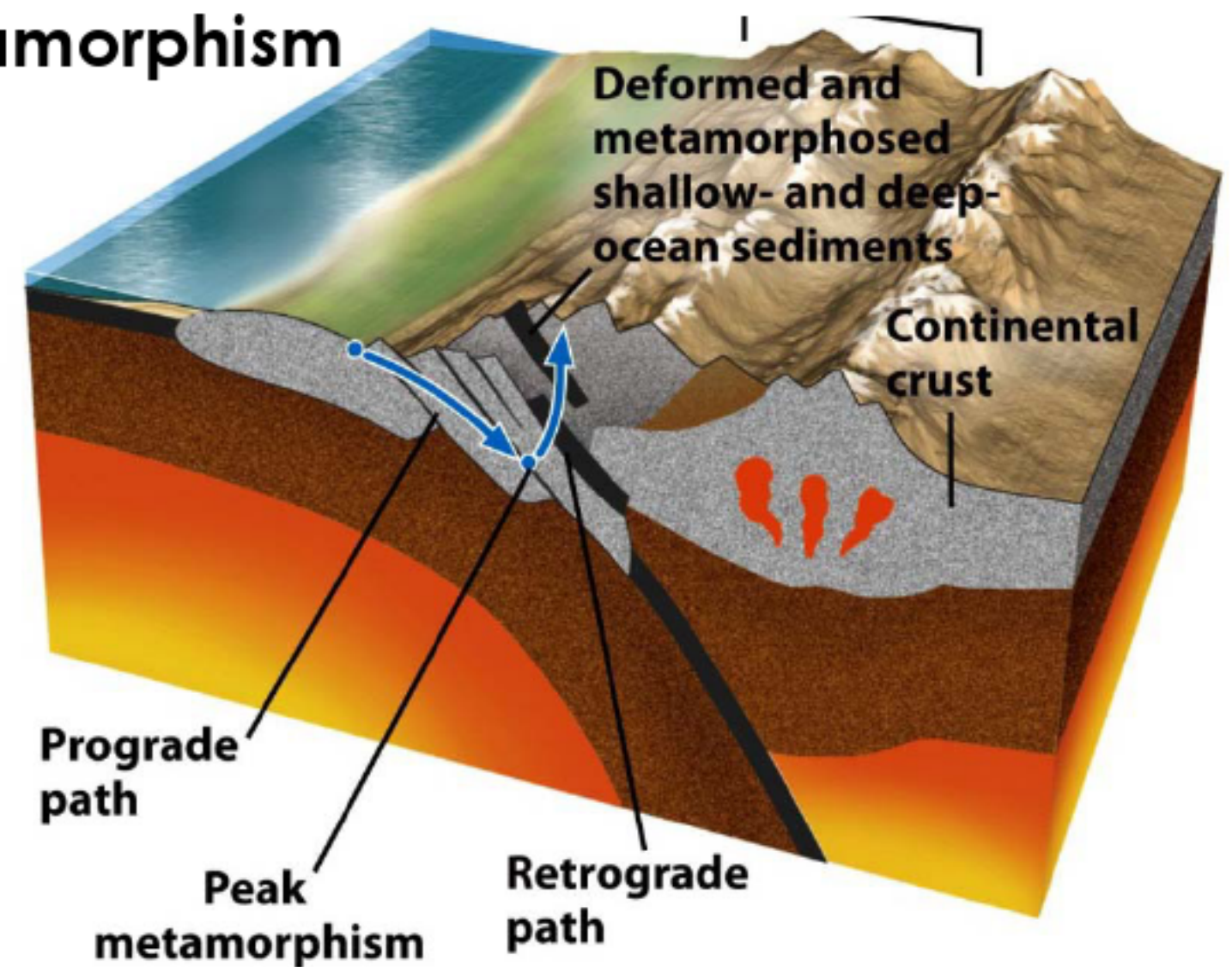
The grain-size of a marble is much larger than that of the parent rock



Regional (Barrovian type) metamorphism

Follows regional “normal” gradient.

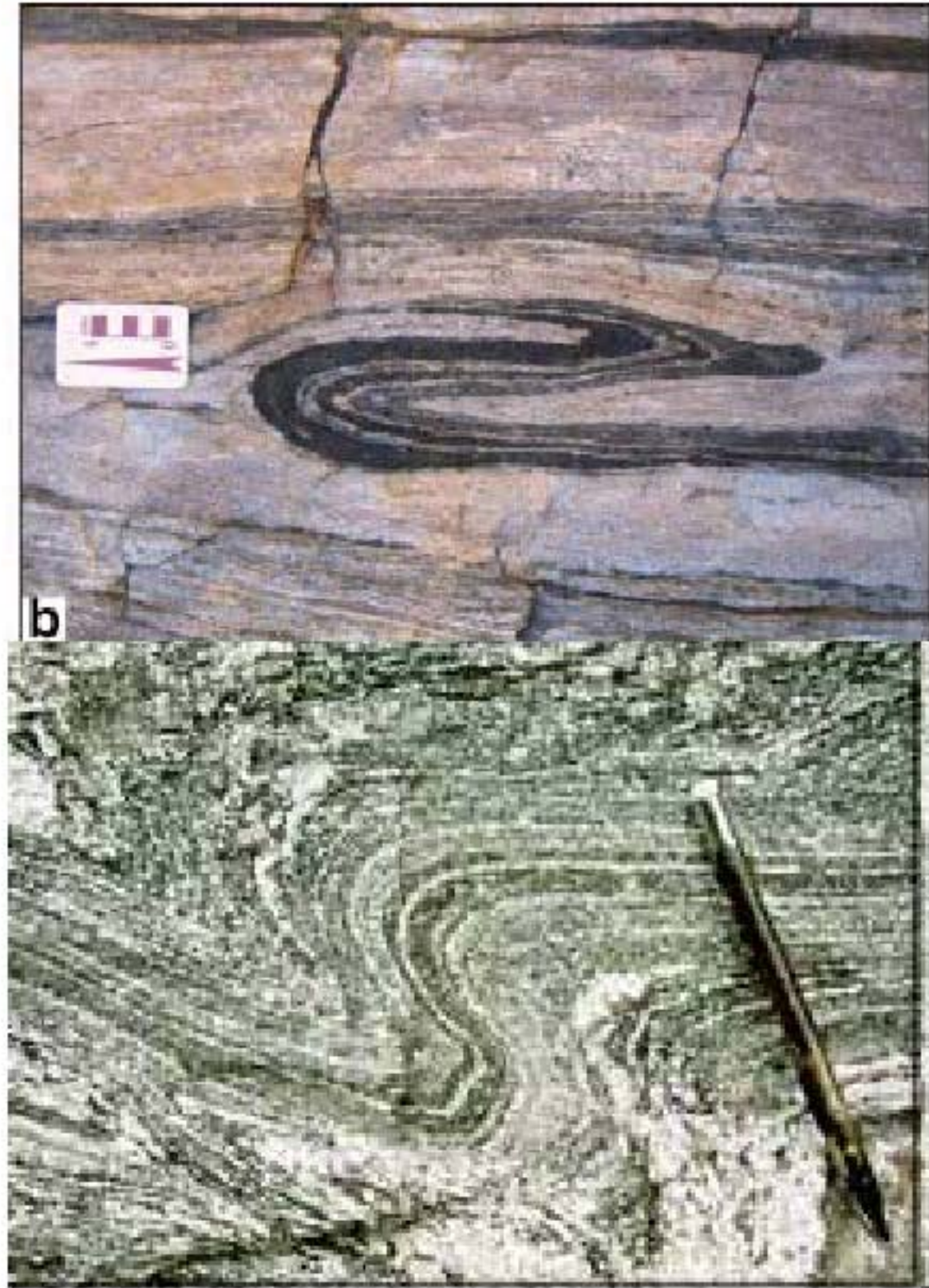
Occurs in continental convergence (collision) zones and is coupled with **strong** deformation



The normal transition for fine-grained rocks

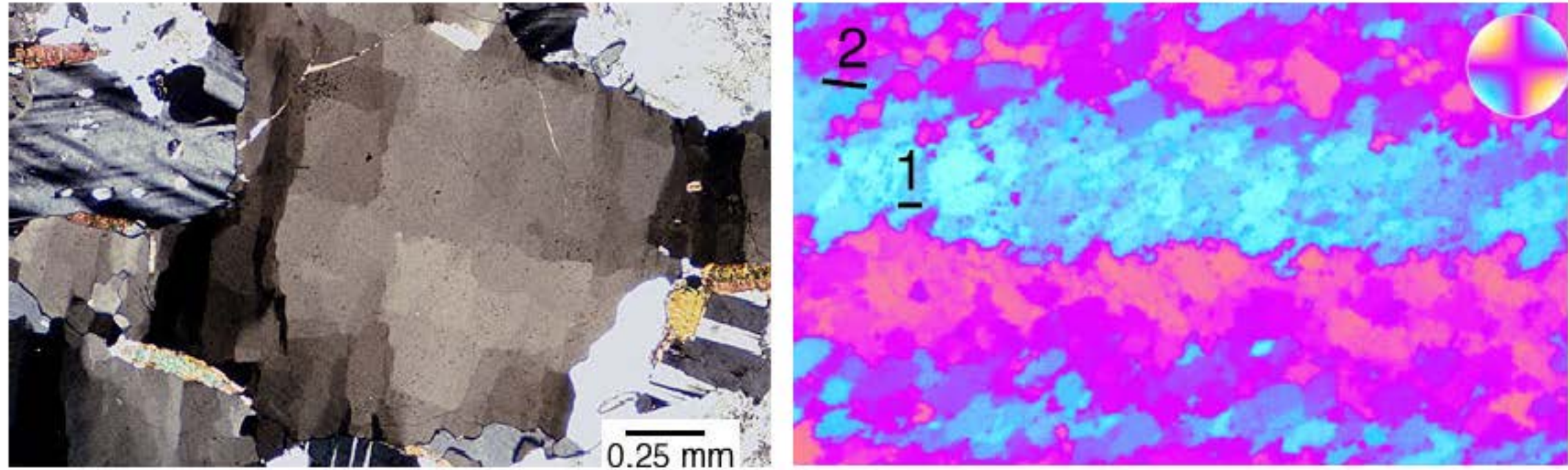
Rocks formed during regional metamorphism are characterized by

- strongly developed **foliation**
- intensive **folding**

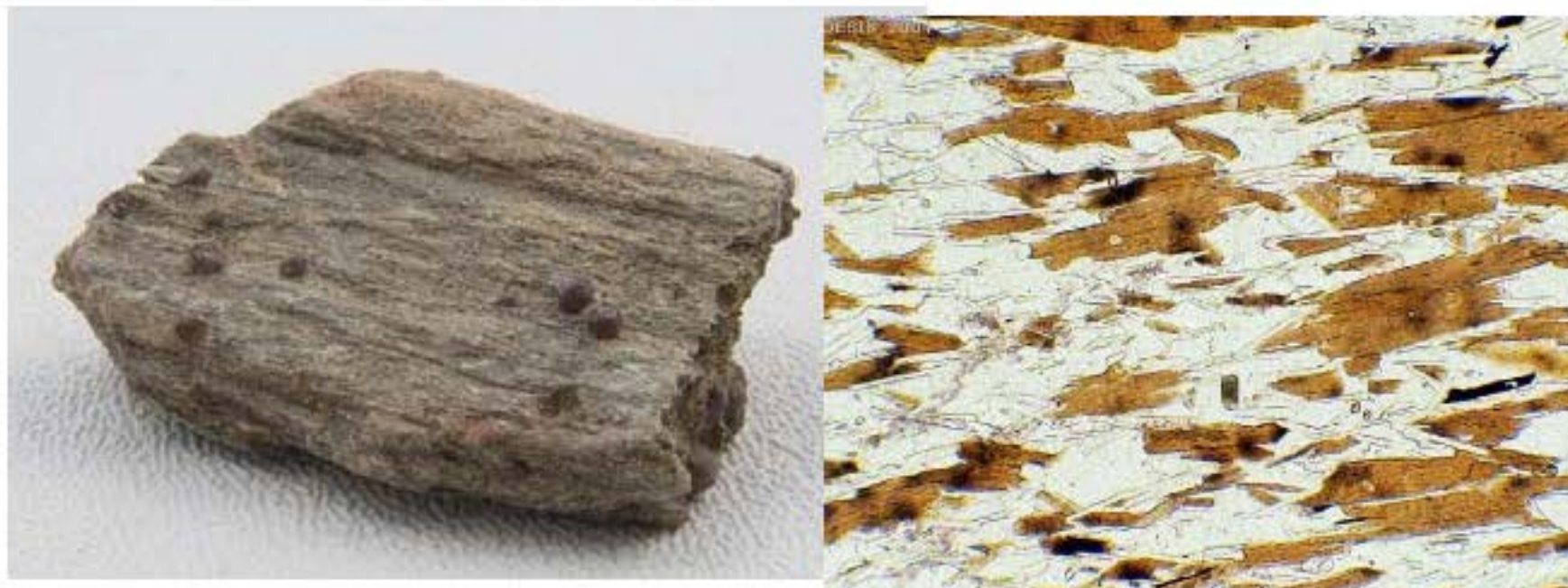


Looking inside the rock, **recrystallization** is pervasive:

Recrystallization: **solid-state** changes in grain size



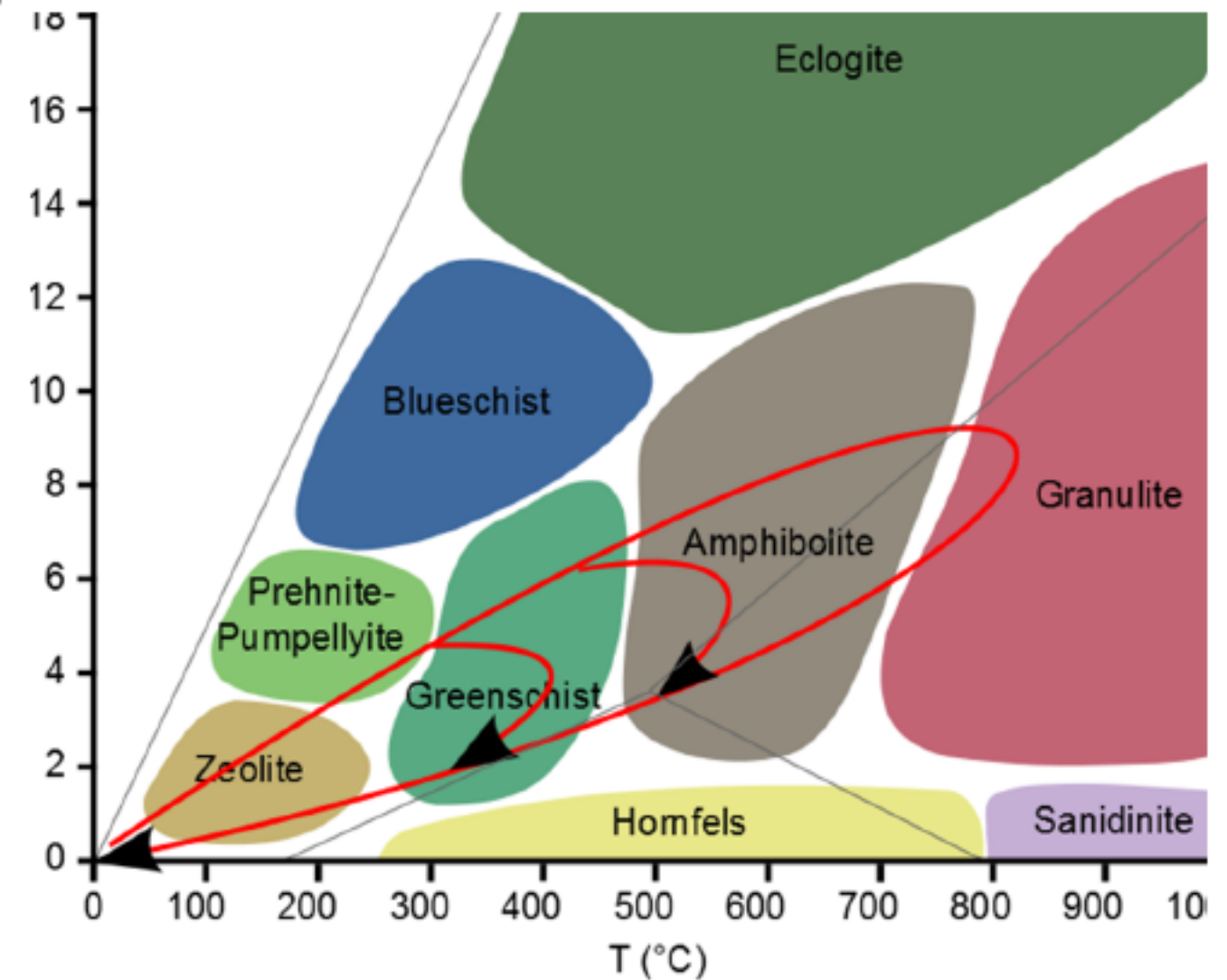
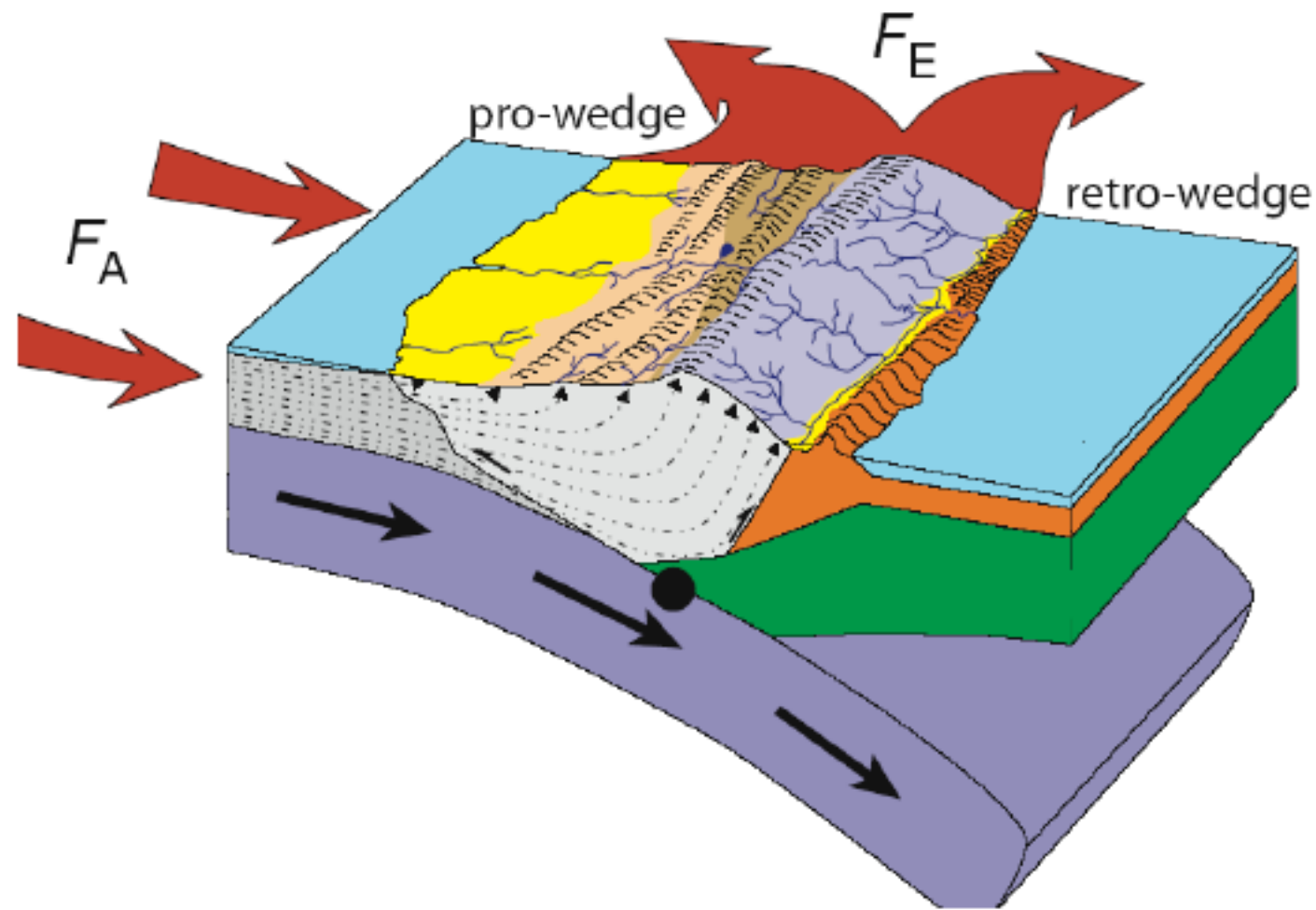
Recrystallization: **solid-state** changes mineral structure and composition



When things get very bad.. part of the rock begins to melt and a **migmatite** is formed



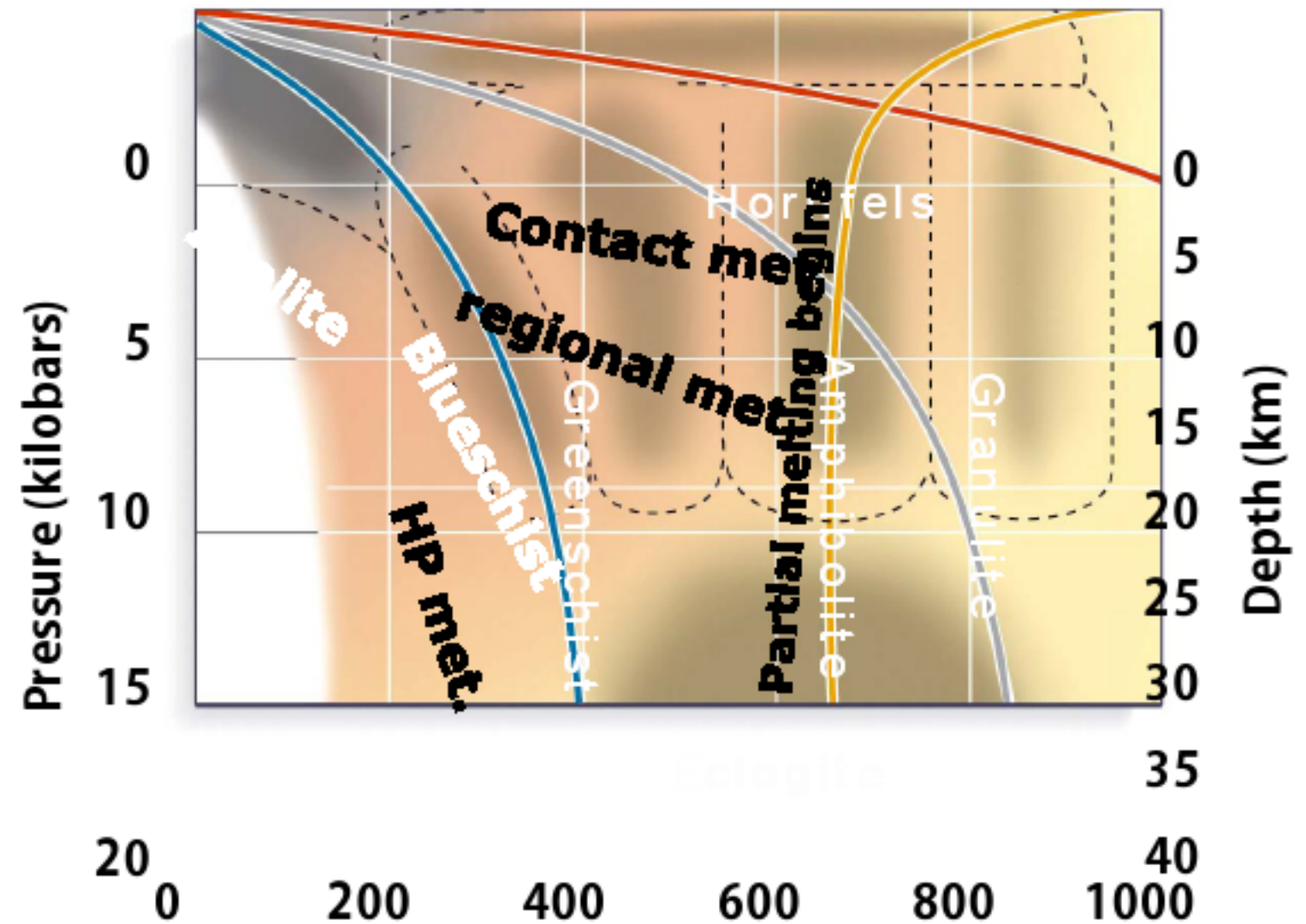
Some of the rocks deformed deep in the orogen will come back to the surface for you to collect (exhumation). This has to happen rapidly enough to prevent **re-equilibration** (also helped by the lack of water)



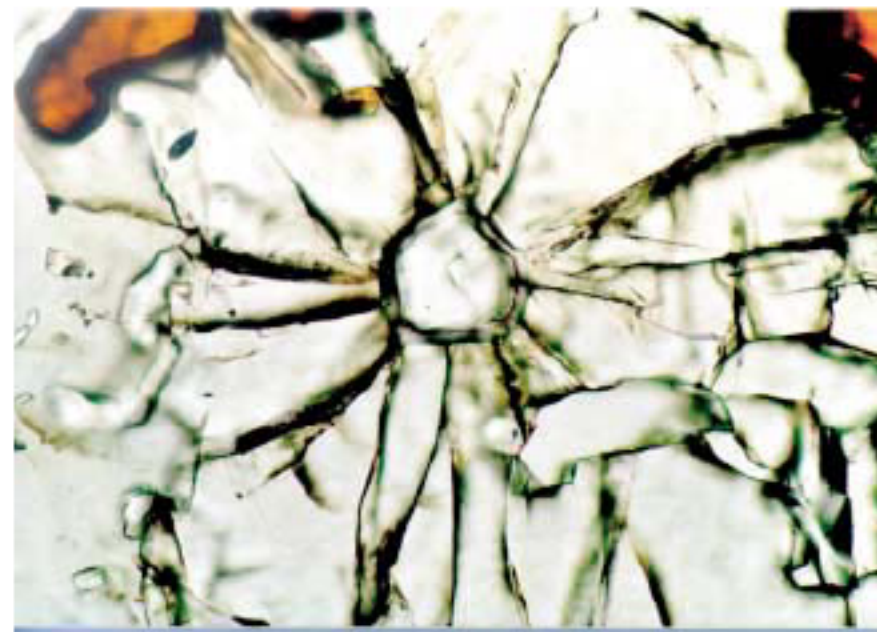
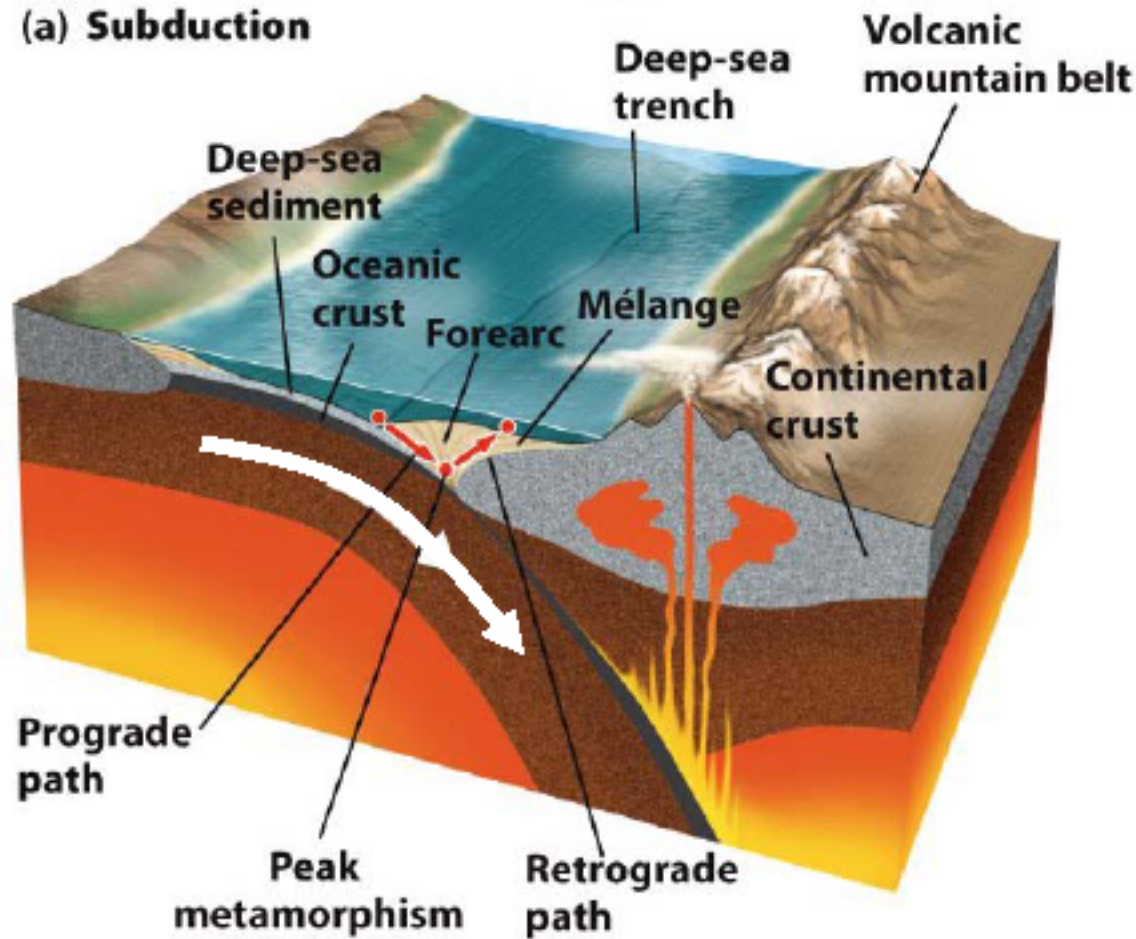
Blueschists

Form at pressures much higher than what predicted by the regional gradient

Typically form along subduction zones: the main mechanism to bring rocks at very large depths (>100km)

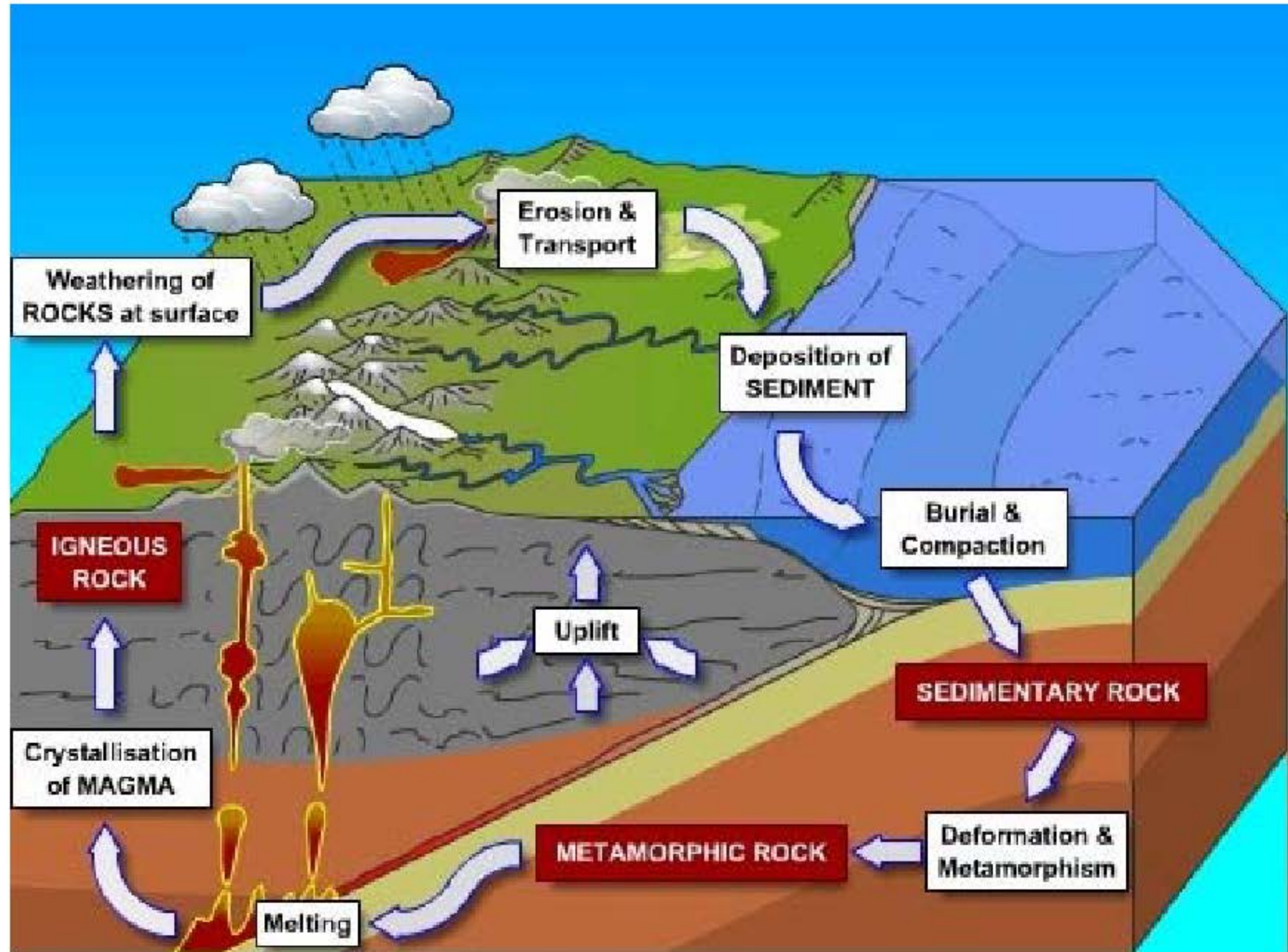


(a) Subduction



This was a diamond (formed at 100km depth) and brought back to the surface

The rock cycle



Sources of figures

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