





Università di Trieste LAUREA MAGISTRALE IN GEOSCIENZE SM62 Percorso Esplorazione Geologica

Anno accademico 2023 - 2024

Geologia Marina 953SM

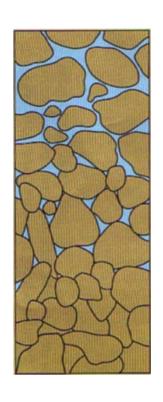
Parte IV

Modulo 4.1 Cause e modalità del movimento di fluidi nei sedimenti

Docente **A. Camerlenghi**







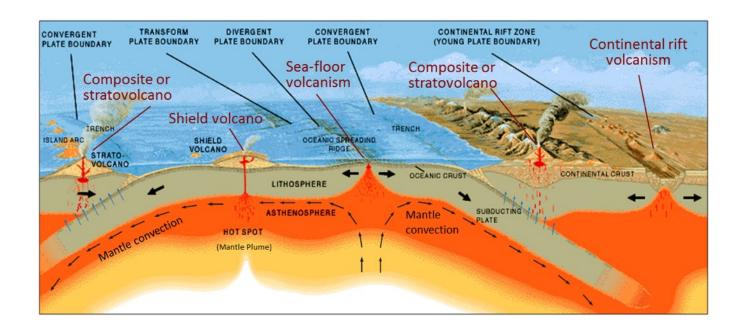
In an ideal sedimentary column where the pore pressure field is hydrostatic, the temperature distribution is controlled by conduction of heat (no advection) and there are no chemical gradients (e.g. salinity, osmosis), the pore fluids do not move. Better: they do move upwards as the porosity decreases with time, but they remain in the sedimentary column.







Net pore fluid movement from the subsurface to through the seafloor is triggered by the establishment of pore **pressure**, **thermal**, **chemical** gradients that depend on the **sedimentary**, **tectonic** and **diagenetic** histories of the sedimentary basin









During the process of consolidation, pore fluids are expelled, initially into the pores of the surrounding sediments

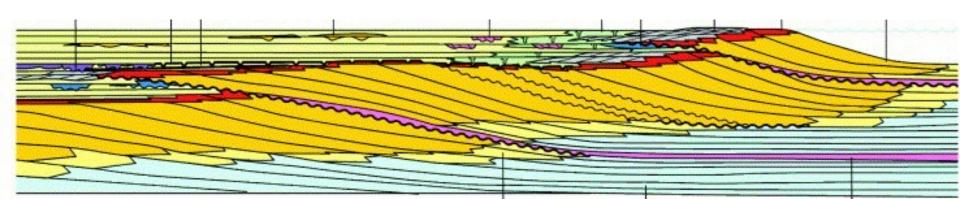
In homogeneous sedimentary columns fluids move up-wards, towards strata with higher porosity >>>> higher permeability.

In inhomogeneous sedimentary columns sands and oozes experience little compaction (they retain more fluids during compaction).

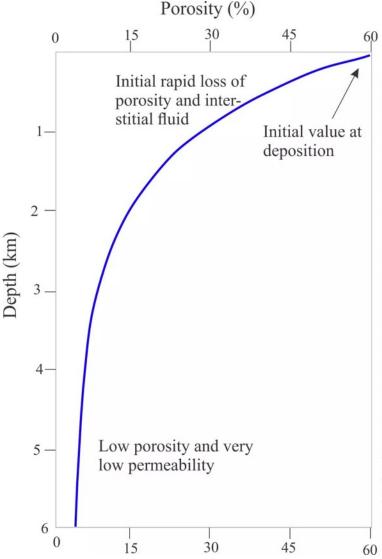
Clays undergo a much larger porosity reduction during compaction (e.g. From 80% to 40 % porosity reduction).

Fluids may move from clay-rich strata to sandy strata by moving laterally, upwards, and downwards.

Alternatively fluids may move using fractures and faults.







A typical compaction curve for shale, here represented by decreasing porosity with depth. Porosity data for this type of plot is derived mainly from wire-line (borehole) sonic and density logs.

There is rapid loss of porosity in the upper kilometer of burial. This kind of curve has great value for determining the burial history of sedimentary basins, and for predicting past episodes of fluid flow.

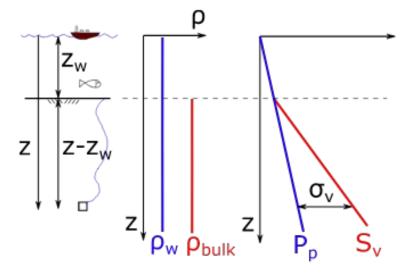
The graph is modified from P.A. Allen and J.R. Allen, Basin Analysis: Principles and Applications. 2nd Ed. Blackwell 2005, Fig. 9.3



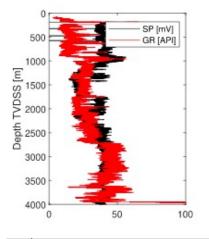


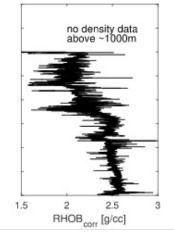


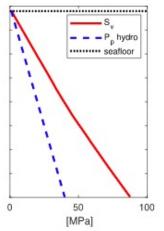
Offshore pressure and total stress gradient (constant density with depth!)



Offshore pressure and total stress gradient (real density changes with depth)







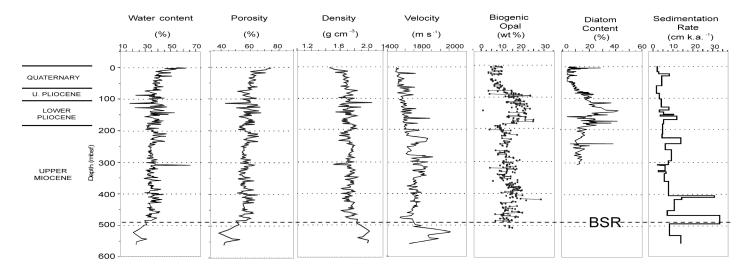




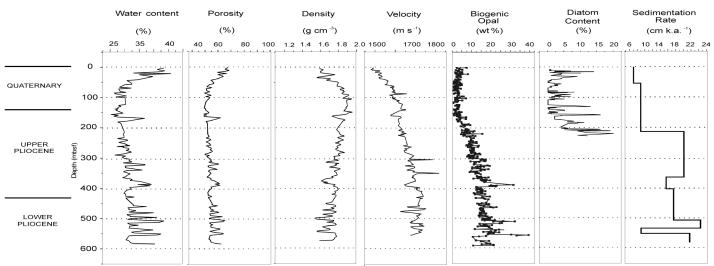


Esempio di caso reale (lo rivedremo per la diagenesis della silice biogenica)

SITE 1095



SITE 1096









Rock Porosity: a measure of its ability to hold a fluid.

Porosity (%) = (Volume of Voids / Total Volume) \times 100

If the volume of the voids is unknown, the porosity equation can be modified in order to calculate the volume of voids, and then the porosity.

Porosity (%) = $\{ (Total Volume - Volume of the Solid) / Total Volume \} x 100 \}$

In cores and drilling data, porosity is provided by the Multi-Sensor Core Logger (MSCL) (on cores) and by Neutron Porosity downhole logging tools (drilling, in situ).

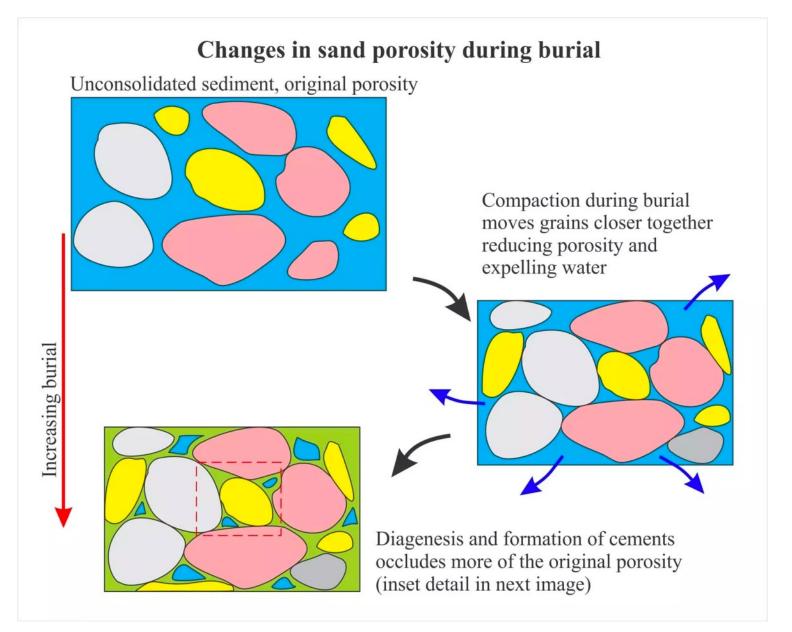
Rock Permeability: a measure of the ability to transmit fluids Measured in. a. geotechnical lab, or during consolidation tests.

The units of permeability are the darcy, D, and m 2 , where 1 D = 0.9869×10^{-12} m² . One darcy is the permeability of a sample 1 cm long with a cross-sectional area of 1 cm² , when a pressure difference of 1 dyne/cm² between the ends of the sample causes a fluid with a dynamic viscosity of 1 poise to flow at a rate of 1 cm³ /s. In geological applications the darcy is commonly too large for practical purposes, so the millidarcy (mD) is used, where 1000 mD = 1D.













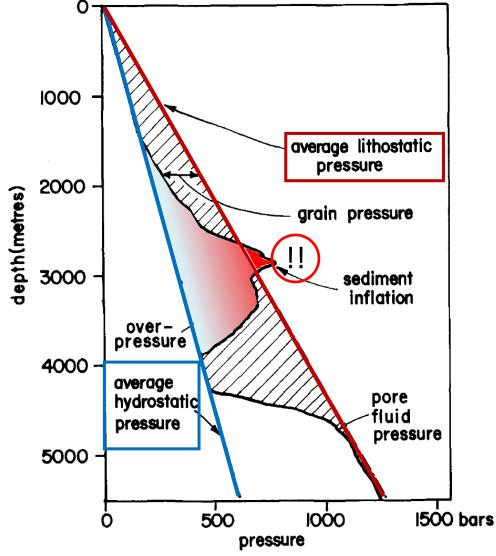


Compaction Disequilibriun

Rapid loading by a huge thickness of the same clay/silt sediment may tip the dewatering balance temporarily in favour of overpressure. In fact the dewatering process is rarely perfectly "normal".

This lack of dewatering conspires to cause the matrix stress between the grains to become "locked" as burial continues, and causes the pore fluids to be responsible for carrying the remaining overburden.

SEDIMENT COMPACTION AND INFLATION GRAIN PRESSURE AND OVERPRESSURE







Compaction Disequilibrium

The process will continue until the fluid pressure finds relief by rupturing the seal.

This rupture can occur at pressures below the overburden if the rock is brittle or even as much as 40% above the overburden if the rocks have enough tensile strength.

Since **compaction disequilibrium** is common in younger clays, a frequent result of this effect is a suite of **mud diapirs**, **mud lumps**, and **sand volcanoes**.





Pore pressure gradients

Basic concepts:

A soil (marine sediment in our case) is a compressible skeleton of particles.

- particles are not compressible
- the interstitial fluid (water, gas, or both) is compressible
- the skeleton is compressible

Volume changes (deformations) do not depend on the total stress applied. They depend on the DIFFERENCE between TOTAL STRESS and PORE FLUID PRESSURE.







$$\Delta V/V = -C_c (\Delta \sigma - \Delta u)$$

 ΔV = volume change (being V the original volume)

 $\Delta \sigma$ = change in total stress

 Δu = change in pore fluid pressure

 C_c = compressibility of the soil skeleton

A volumetric change can be obtained by an increase in $\Delta \sigma$ as well as by a decrease in Δu







Sediment consolidation state

Consolidation is the volumetric change induced by a change of total stress (sedimentary burial, tectonic burial).

When a sediment is buried, it undergoes a process of natural consolidation due to the progressive increase of the overburden (total stress).

NORMALLY CONSOLIDATED

The overburden is the maximum stress ever experienced.

OVER-CONSOLIDATED

The overburden is the less than the maximum stress ever experienced.

UNDER-CONSOLIDATED

The overburden is more than the maximum stress ever experienced.







Factors that determine excess pore water pressure and influence the consolidation state

The most common factors are:

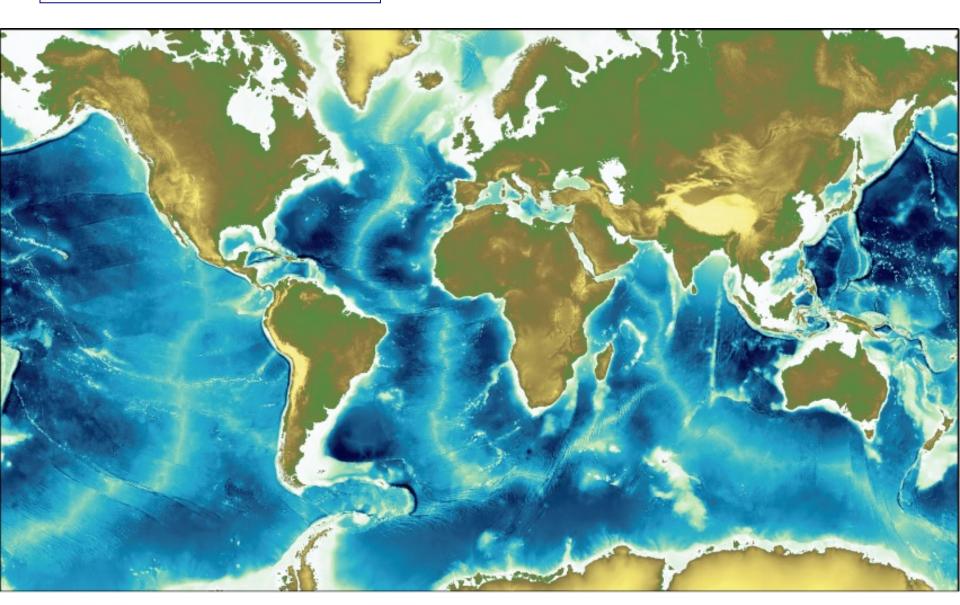
- Stress-related factors
 - Rapid sedimentation rate
 - Tectonic stress
- Fluid volume increase mechanisms
 - Temperature increase
 - Mineral transformation
 - Hydrocarbon Generation
- Fluid movement mechanisms
 - Osmosis
 - Hydraulic head
 - Hydrocarbon buoyancy

(Swarbrick and Osborne, 1998. Bryant et al., 1974; Sangrey, 1977; Arthur et al., 1980; Demaison & Moore, 1980; Bryant et al., 1981).







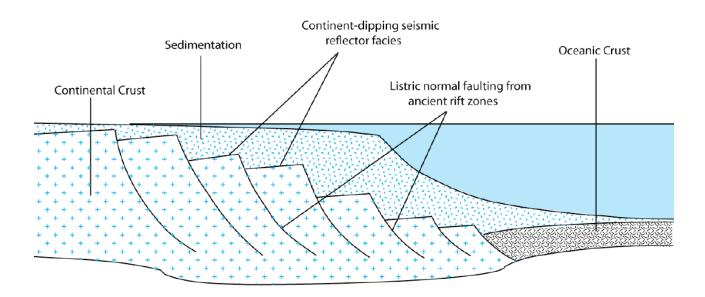








Rifted Passive Margin



Rapid sedimentation can be produced by:

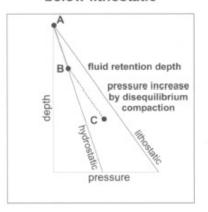
- River-dominated sedimentary systems (low and medium latitudes)
- Glacial-dominated sedimentary systems (high latitudes)



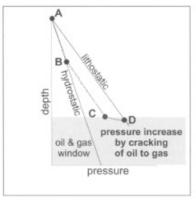




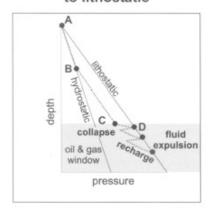
pore fluid pressure below lithostatic



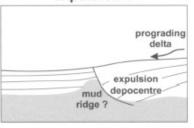
pore fluid pressure increase



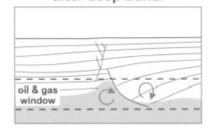
pore fluid pressure to lithostatic



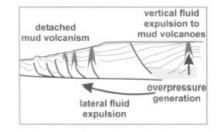
progressive compaction or mud expulsion?



pore pressure build-up in prodelta shale after deep burial



expulsion of fluid+gas+mud



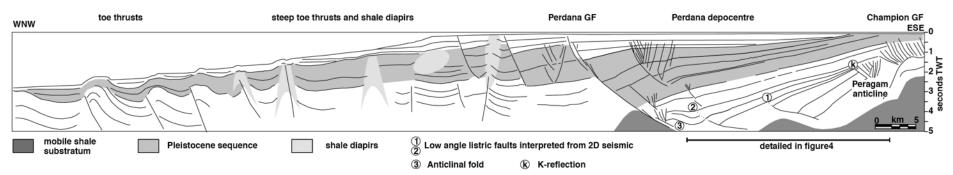
Overview of the styles of shale mobility in relation in relation with pore fluid overpressure; early syndepositional structures can be distinguished from later post-depositional injections of mobilized sediment that occur after large overpressure increase.



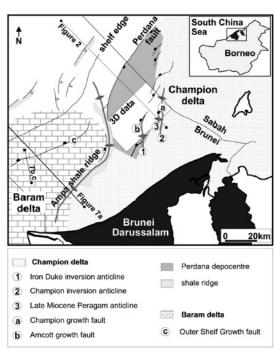




Shale tectonics, Offshore Brunei



Schematic cross-section of the Champion delta based on 2D regional seismic line.

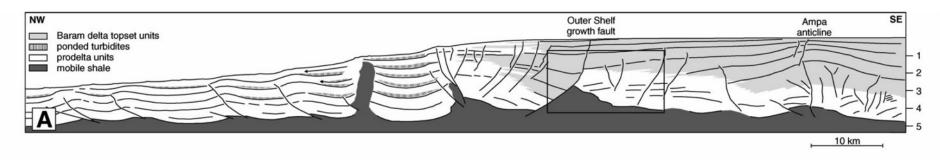


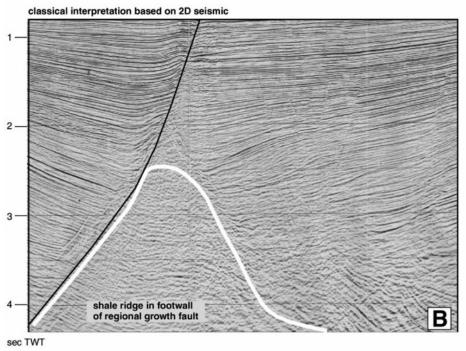


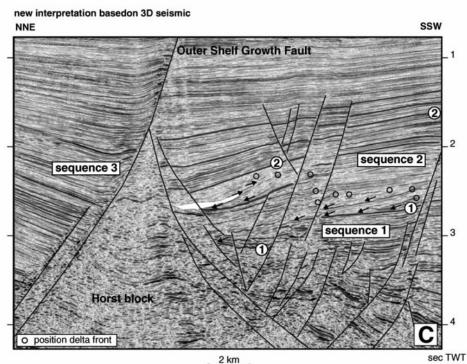




Shale tectonics, Offshore Brunei



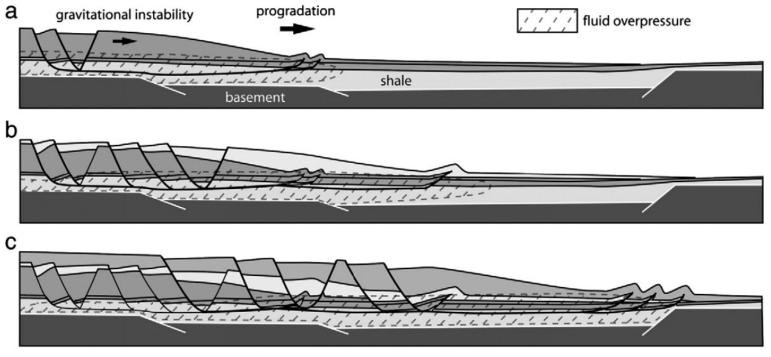












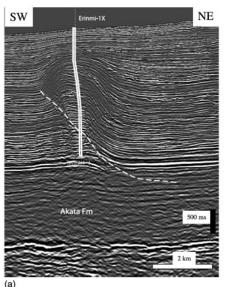
Evolutionary model of fluid overpressure and gravitational deformation in a prograding delta. Whatever the generating mechanism may be (compaction or hydrocarbon generation), the front of the overpressured domain may advance basinward as sediments prograde seaward.

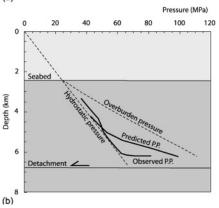


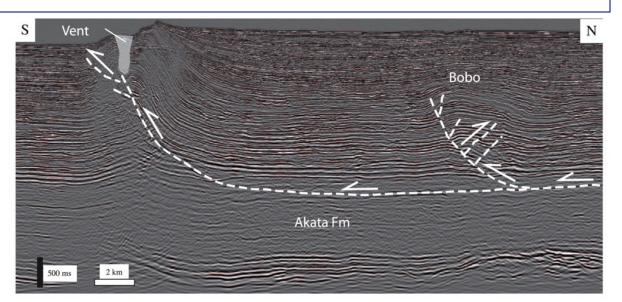




Shale tectonics in the Niger Delta





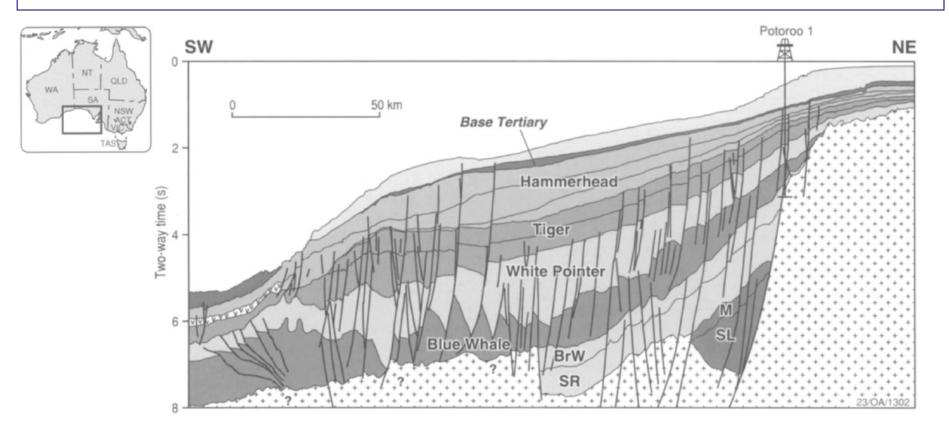










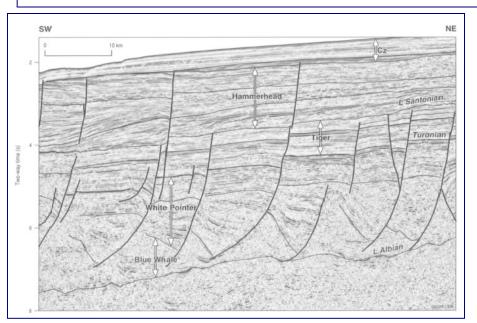


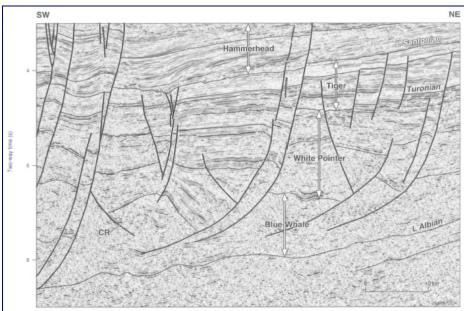
Geoseismic profile across the northern Ceduna Sub-basin. Jumssic-E~ly Cretaceous supersequences abbreviated as follows: SL Sea Lion; M Minke; SR Southern Right; BrW Bronze Whaler

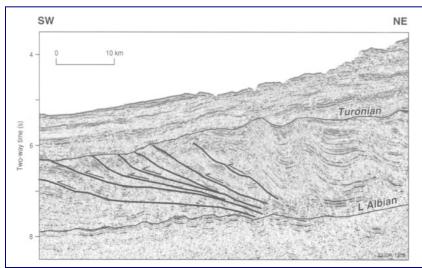














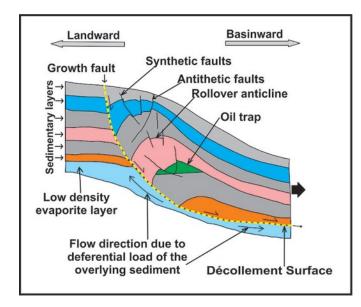




Compaction Disequilibrium

Growth Faults

Growth faults are syndepositional or syn-sedimentary extensional faults that initiate and evolve at the margins of continental plates.



The pressures can sometimes be relieved by systems of **sub-vertical faults** or by **growth fault systems**.

The high pressures in these shale masses are a major contributing factor in the formation of massive "growth faults" that cut across the delta, trapping the rollover anticlines (which often form traps for oil and gas in the hanging- wall).

The faults may also trap oil on the foot-wall side where the movement has brought sands against shales to seal them.

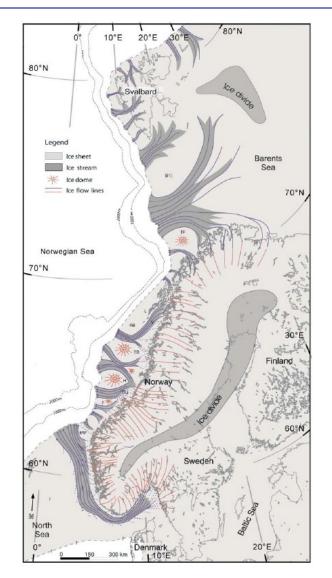


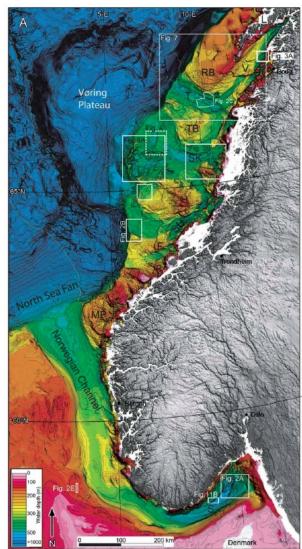




Rapid sedimentation rates in glacial environments

North Sea Fan



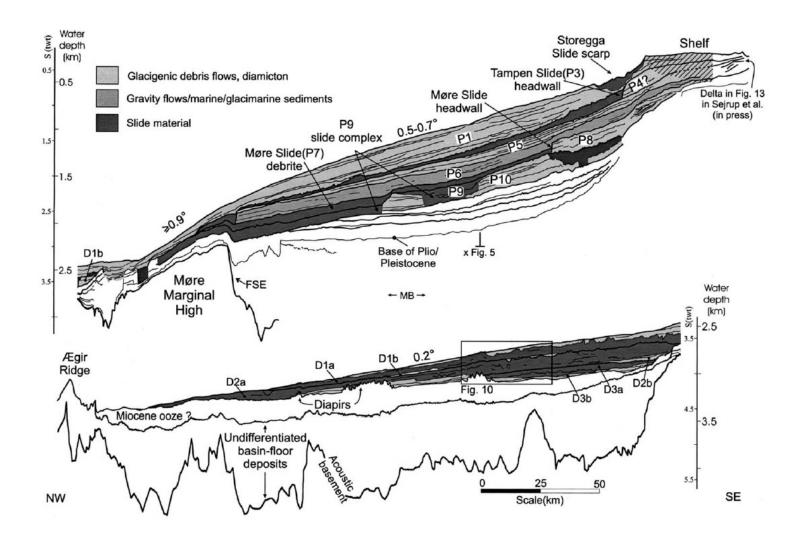








Rapid sedimentation rates in glacial environments

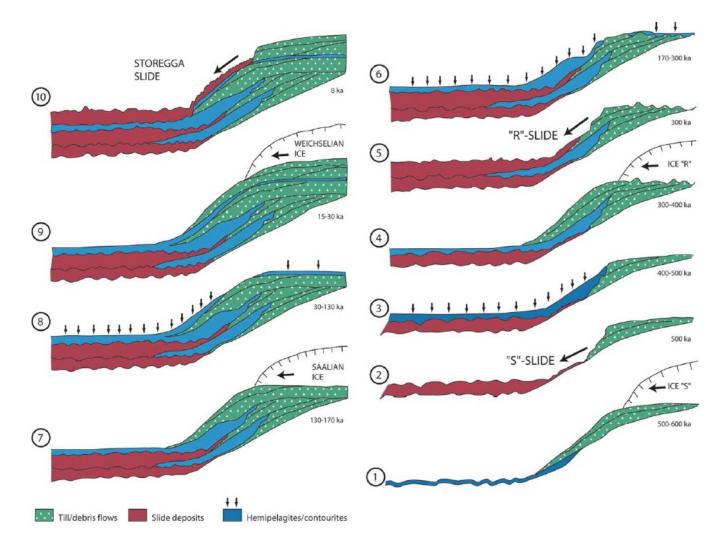








Rapid sedimentation rates in glacial environments

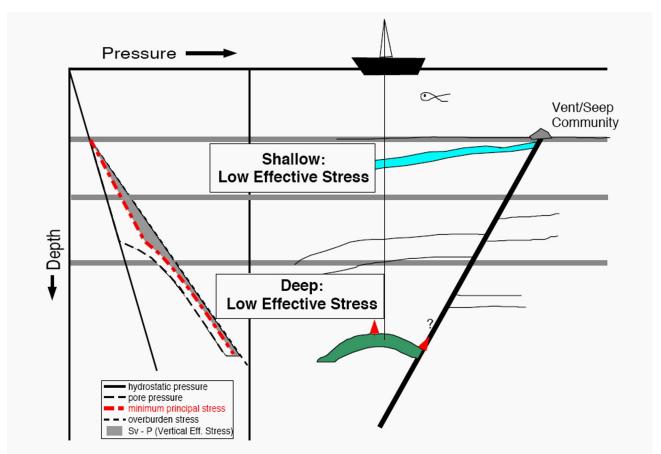








Rapid sediment loading (>1 mm a^{-1}) is documented as a source of overpressure (P^* , pressure in excess of hydrostatic) in basins around the world. When low-permeability sediments are rapidly loaded, pore fluids cannot escape, and the fluids bear some of the overlying sediment load. In this situation a pore pressure exceeding the hydrostatic pressure develops.

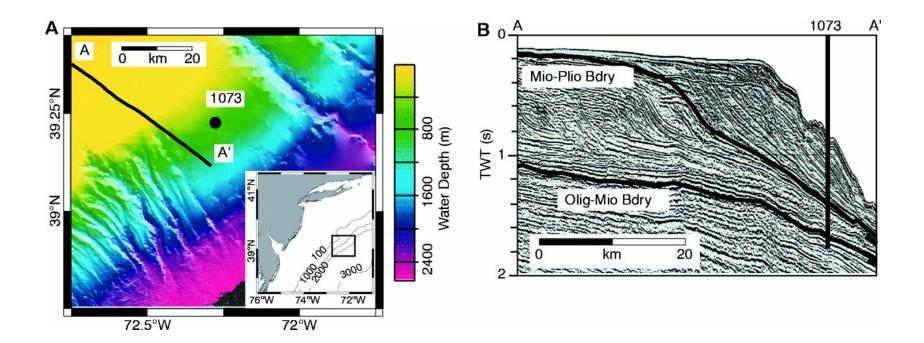








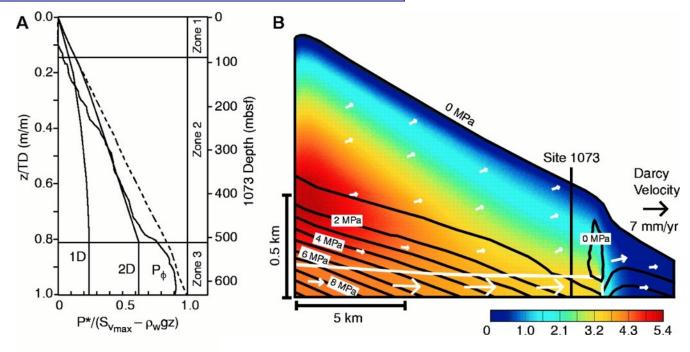
New Jersey Margin. Rapid sedimentation of clays over a high permeability sand layer

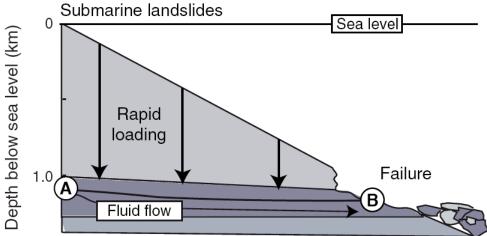








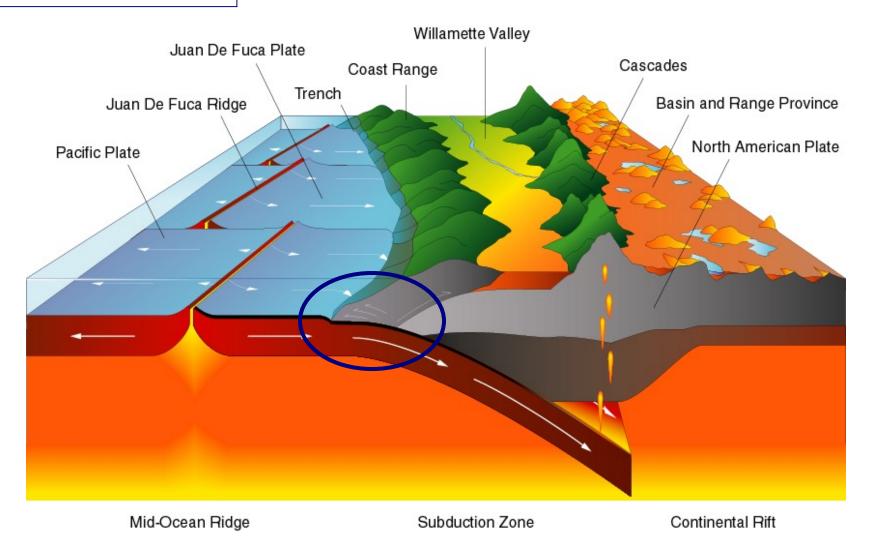








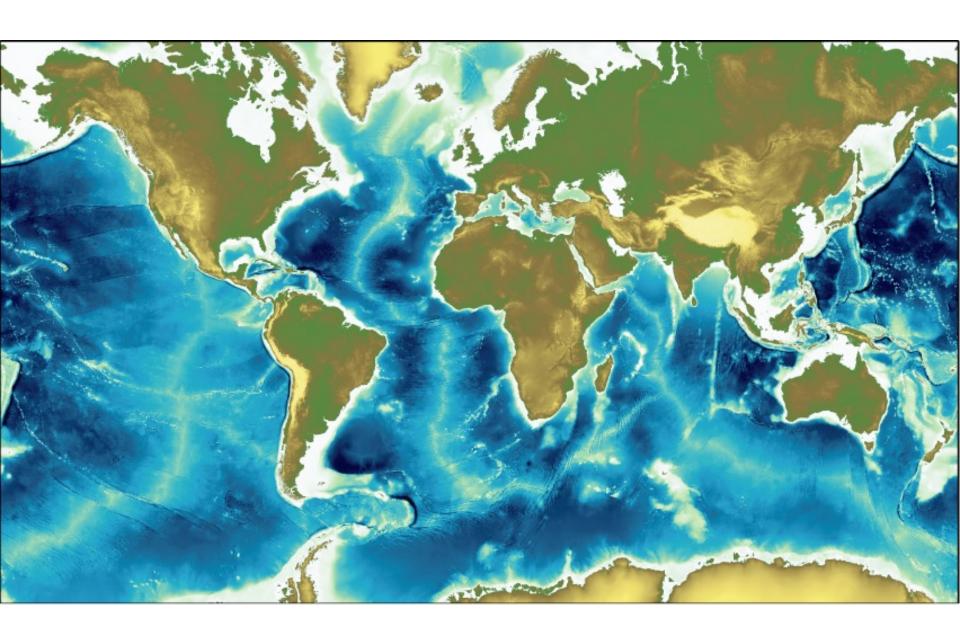




Subduction zones are most prevalent in the Pacific Ocean



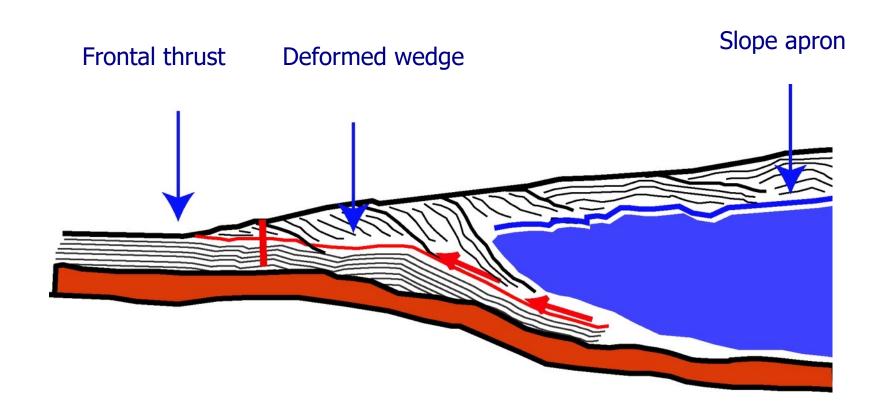


















DISPERSED FLOW

If the rate of pore fluid flow is sufficiently low that advection can be accommodated by intergranular permeability, fluids might be expected to leak out of the prism over large areas, as "dispersed flow," driven by some regional sublithostatic pressure gradient.

FOCUSSED FLOW

In contrast, observations establish that flow is often focused at vent sites, confined to fault zones, or expelled from mud volcanoes.

Fluid loss evolves from dispersed to focused flow over time. Observations indicate **episodic flow**, which may be coupled to episodic fault displacement and ultimately to the earthquake cycle.







At the deformation front the incoming sediments above the decollement are folded and/or faulted, usually into elongated ridges that rise 200-700 m above the adjacent deep-sea floor (**proto-thrust zone**)

Sequential faulting thickens the accreting sediment section and causes the deformation front to step seaward periodically, widening the lower continental slope. The **thrust faults** rise from the decollement, and controls the amount of sediment that is initially subducted.

The initial **frontal thrust fault** and others that originate after uplift and folding ("**out-of-sequence**" **thrusts**) rapidly emplace older, compacted sediments above younger, less consolidated deposits.

This tectonic burial creates overpressures that lead to fluid expulsion.

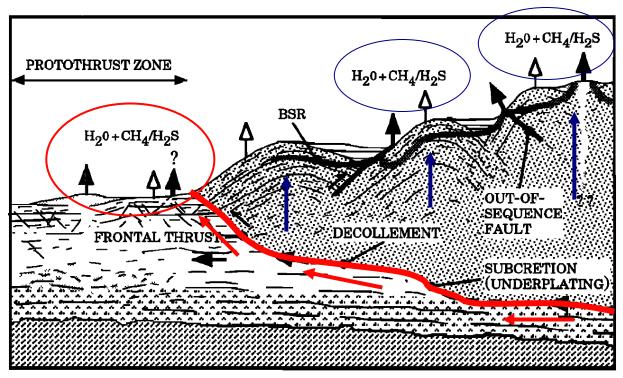






The fluids, are expelled along the decollement are primarily sourced from the underthrust sediments,

The fluids derived from compaction within the prism generally move upsection to the seafloor.



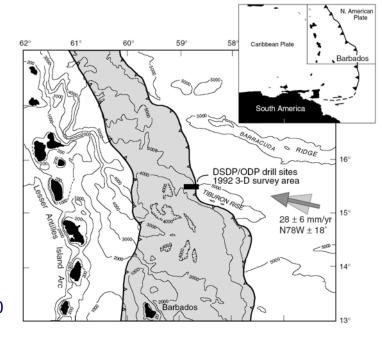
Fluid pressures at the decollement can be approximately lithostatic



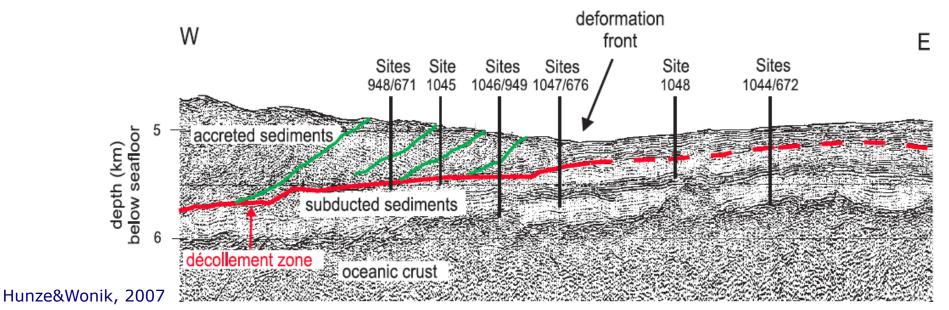




ODP Leg 171A. Barbados Accretionary Prism



Moore, 2000

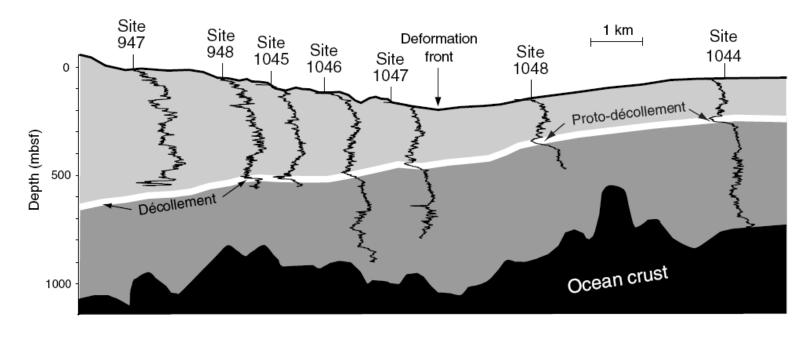


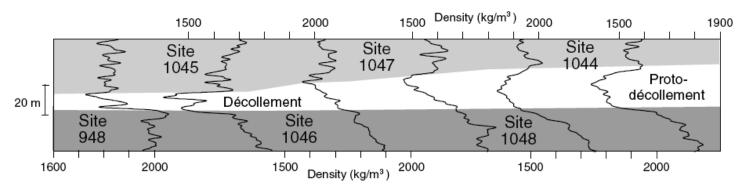






ODP Leg 171A. Barbados Accretionary Prism



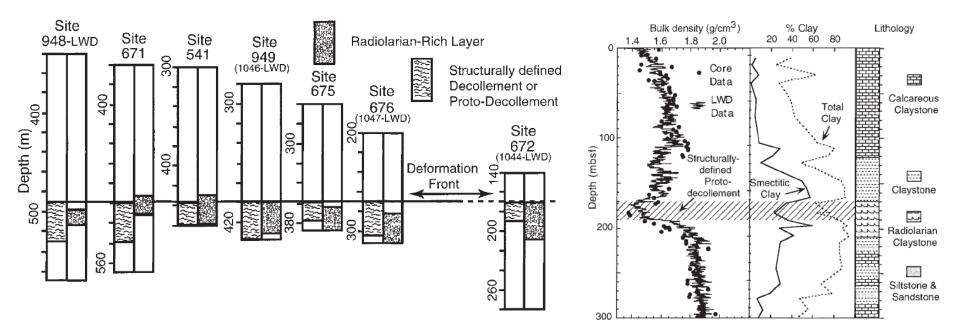








ODP Leg 171A. Barbados Accretionary Prism



The decollement zone initiates in a low-density smectitic radiolarian claystone.

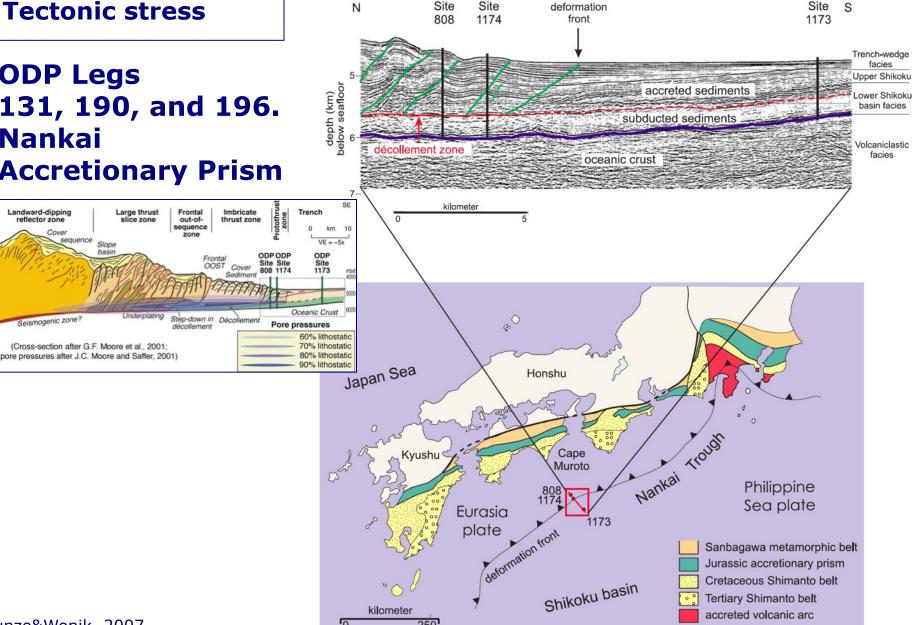
During underthrusting, the decollement zone consolidates heterogeneously due to shear-induced collapse of the radiolarian tests and clay fabric. This consolidation locally increases the pore pressure and facilitates underthrusting.







ODP Legs 131, 190, and 196. Nankai **Accretionary Prism**









Factors that determine excess pore water pressure and influence the consolidation state

The most common factors are:

- Stress-related factors
 - Rapid sedimentation rate
 - Tectonic stress
- Fluid volume increase mechanisms
 - Temperature increase
 - Mineral transformation
 - Hydrocarbon Generation
- Fluid movement mechanisms
 - Osmosis
 - Hydraulic head
 - Hydrocarbon buoyancy

(Swarbrick and Osborne, 1998. Bryant et al., 1974; Sangrey, 1977; Arthur et al., 1980; Demaison & Moore, 1980; Bryant et al., 1981).

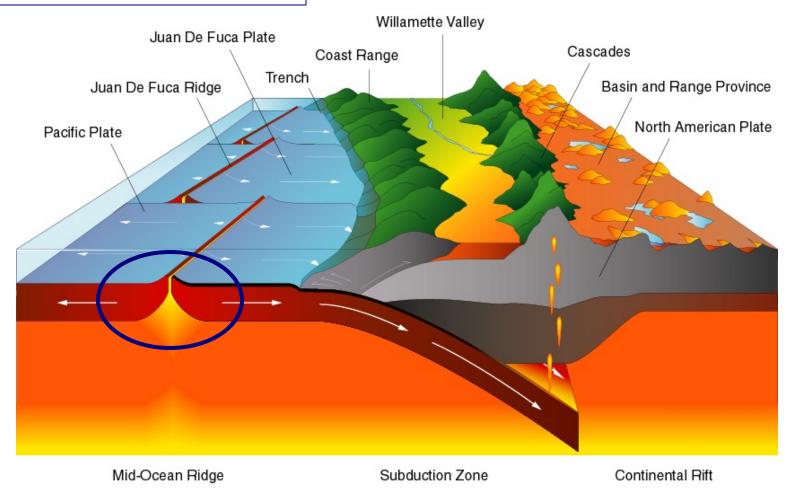






Temperature increase

Heat transfer at mid-ocean ridges



Mid ocean ridges (spreading centres) are present in all oceans. The spreading rate is fast in the Pacific, and slow in the Atlantic







Modulo 4.6 Identificatori di movimento di fluidi: la circolazione idrotermale delle dorsali oceaniche





- Smectite dehydratation
- Smectite to Illite Transformation
- Gypsum to Anhydrite Dehydratation



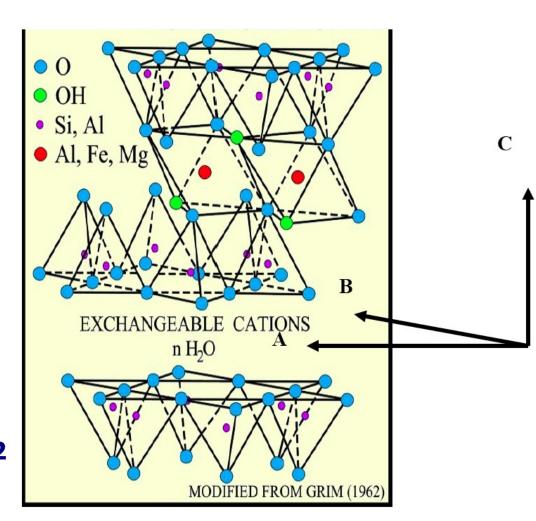




Smectite dehydratation

A significant feature of the smectite group is its very high surface area. These clay platelets are held together by a weak electromagnetic force (Van der Waal's bonds), and there is a considerable amount of area to which up to ten layers of water can bond. The result is a low density "swelling clay".

Under normal hydrostatic conditions smectite contains 2 or 3 layers of water.





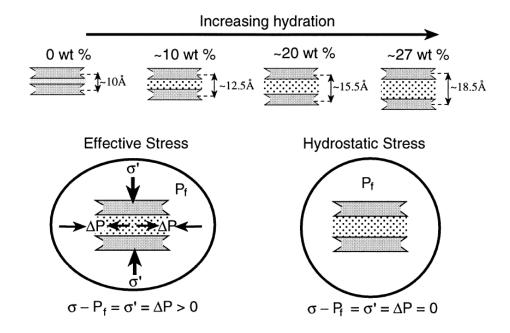




Smectite dehydratation

Smectite changes with burial. Effect of lithostatic stress.

1) Increasing pressure will drive out the loosely bound water



Smectites can partially dehydrate from 18.5 \AA to 15.4 Å hydrates when they are subjected to effective mineral framework stresses above 1.3±0.3 MPa (sort of squeezing of interlayer water out of the mineral structure)







Smectite dehydratation

Role of temperature

- 1) Under stress (burial) at temperature of about 60°C, smectite molecules will lose interlayer water except the last two layers.
- 2) With temperature of about 100°C one the penultimate layer will be displaced.
- 3) A further rise in temperature to 172° 192° C is required to drive off the last layer, which is very closely bound between the clay plates.

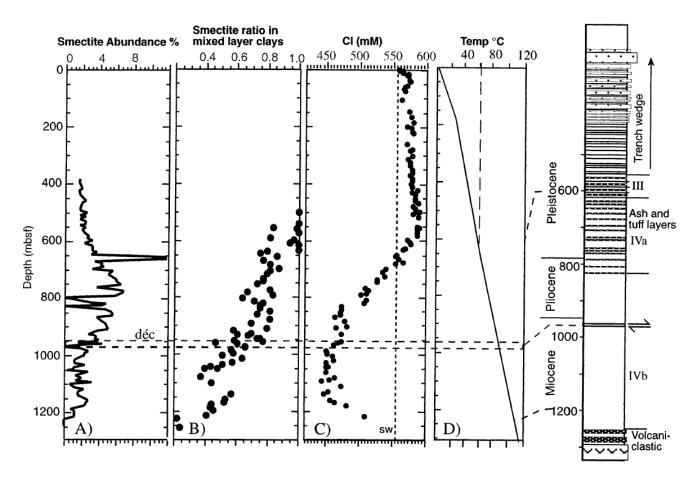
Although the initial dewatering may actually cause some overpressures, the resulting extra hydraulic pressure will also tend to inhibit further dewatering. Therefore, under most conditions the simple dewatering process will not lead to excessive overpressure (**volume increase of about 4%**), since there is a negative feedback loop at work.







Smectite dehydratation



Possibility that the observed low-Cl anomaly is a compound effect of both lateral flow and in situ smectite dehydration.



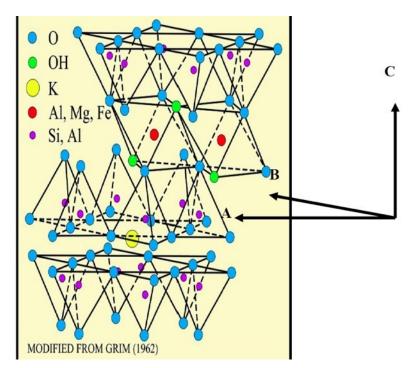




Smectite to Illite Transformation

During Burial, smectite alters chemically to produce illite

Smectite + (Al, K) - (Na, Ca, Mg, Fe, Si, water) >>>> illite







Smectite to Illite Transformation

- The reaction is kinetically controlled.
- It usually happens at temperature range of 70°C 150°C
- It often accompanies the loss of the penultimate water layer of smectites.
- It has been shown that in order for the transformation to occur Potassium **(K) ions are required** in the interstitial fluids. These come from K-Feldspars
- Illites do not have the same capacity to adsorb water as smectities.
- The **threshold temperature** for the loss of the penultimate water layer is roughly the same at which **hydrocarbons are generated**.





Smectite to Illite Transformation

The volumetric change of Smectite to Illite is unknown.

Some models say $\Delta V = + 23-25 \%$

It is a fact that overpressure is often detected in wells in correspondence with such diagenetic transformation.

Consequences during the transformation:

- Compressibility increases
- Overpressure
- Compaction disequilibrium
- Perhaps reduction of permeability by silica cementation





Gypsum to Anhydrite Dehydratation

Heating gypsum to between 100 °C and 150 °C partially dehydrates the mineral by driving off approximately 75% of the water contained in its chemical structure.

The temperature and time needed depend on ambient partial pressure of H_2O .

The reaction for the partial dehydration is:

$$CaSO_4 \cdot 2H_2O + heat >>> CaSO_4 \cdot nH_2O + nH_2O$$

Anhydrite: $CaSO_4 \cdot nH_2O$, where n is in the range 0.5 to 0.8.

The temperature of the reaction depends on the physical and geochemical environments





Gypsum to Anhydrite Dehydration

Geological factors:

- Thermal conductivity of the overlying rock
- Basal heat flow
- Porosity
- Sedimentation rate

Geochemical factors:

- Activity of pore water
- Pore fluid pressure







Hydrocarbon is an organic compound consisting entirely of hydrogen and carbon.

"Impure" hydrocarbons include bonded compounds or impurities of sulphur or nitrogen

- 1 **Saturated hydrocarbons** (**alkanes**) are the most simple of the hydrocarbon species and are composed entirely of single bonds and are saturated with hydrogen; they are the basis of petroleum fuels and are either found as linear or branched species of unlimited number. The general formula for saturated hydrocarbons is C_nH_{2n+2} .
- 2 **Unsaturated hydrocarbons** have one or more double or triple bonds between carbon atoms. Those with one double bond are called alkenes, with the formula C_nH_{2n} (assuming non-cyclic structures). Those containing triple bonds are called alkynes.
- 3 **Cycloalkanes** are hydrocarbons containing one or more carbon rings to which hydrogen atoms are attached. The general formula for a saturated hydrocarbon containing one ring is C_nH_{2n}
- 4 **Aromatic hydrocarbons**, also known as arenes which have at least one aromatic ring







Natural gases (alkanes gases)

METHANE CH₄ ETHANE C₂H₆ PROPANE C₃H₈ BUTANE C₄H₁₀

METHANE: OXYDATION OF ORGANIC MATTER BY ANAEROBIC BACTERIA (BIOGENIC)

 C_{2+} GASES: THERMAL MATURATION OF ORGANIC MATTER (THERMOGENIC)







IN NATURAL GAS THERE ARE TWO TYPES OF METHANE:

Biogenic Methane

Biogenic methane is produced as an end product of the metabolism of a diverse group of obligate **anaerobic archaea** (killed by even traces of oxygen), generally known as **methanogens**.

Methanogens live in the shallow geosphere such as **marine** and lacustrine sediments (sub-bottom depth less than 1 km, temperature less than 80°C).







In the absence of light, photosynthesis cannot occur. Respiration makes use oxygen of dissolved in water with microbial mediation.

Organic matter (preferably proteins and carbo-hydrates) is decomposed trough hydrolysis by micro-organisms, that produce sugars and aminoacids that more evolved organisms can use.

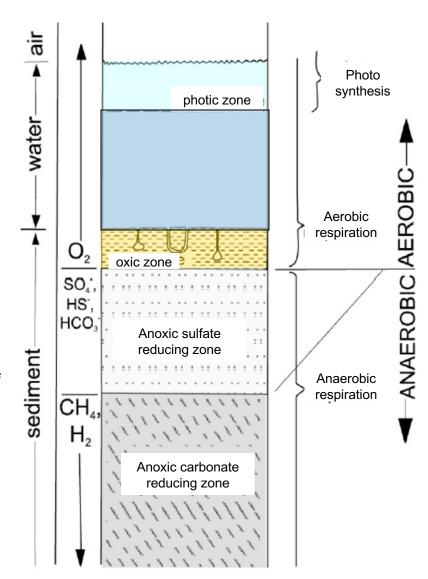
Aerobic respiration in the interstitial fluids of the uppermost sediments can cause **oxygen depletion** in pore water (usually the exchange of fluids between pores and water column is negligible due to low permeability).

Respiration then uses oxygen from Mn-Oxides, Nitrates and Iron-Oxides (**sub oxic Zone**)

Aerobic respiration is then replaced by **anaerobic fermentation**, that is the microbial reduction of the sulfate ion (SO_4) with release of S^- ions and production of H_2S , and CO_2 .

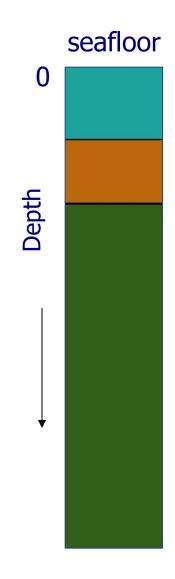
When all the Sulfate ion has been used, the Archea begin to produce methane (CH₄) from the remaining organic matter.

The biogenic production of methane continues until the temperature of 75-80°C is reached.









- 1. Aerobic (respiration): $CH_2O + O_2 = CO_2 + H_2O$
- 2. Sub-oxic zone

3. Anoxic (sulfate reducing) zone: $CH_2O + SO_4 = CO_2 + H_2S$

4. Methanogenesis: $CH_2O = CO_2 + CH_4$









			Energetically
	Oxidation by dissolved Oxygen		
	$(CH_2O)_{106}(NH_3)_{16}(H_3PO_4) + 138 O_2 \rightarrow$	-3190 kJ/mol	most favorable
	106 CO ₂ + 16 HNO ₃ + 122 H ₂ O + H ₃ PO ₄		
	Oxidation by Manganese-Oxides		
	$(CH_2O)_{106}(NH_3)_{16}(H_3PO_4) + 236 MnO_2 \rightarrow$	-3090 kJ/mol	
	236 Mn ²⁺ + 106 CO ₂ + 8 N ₂ + 366 H ₂ O + H ₃ PO ₄		•
	Oxidation by Nitrate		
	$(CH_2O)_{106}(NH_3)_{16}(H_3PO_4) + 84.4 \text{ HNO}_3 \rightarrow$	-2750 kJ/mol	
	106 CO ₂ + 42.2 N ₂ + 16 NH ₃ + 148.4 H ₂ O + H ₃ PO ₂	4	
	Oxidation by Iron-Oxides		
	$(CH_2O)_{106}(NH_3)_{16}(H_3PO_4) + 212 \text{ Fe}_2O_3 + 848 \text{ H}^+ \rightarrow$	-1410 kJ/mol	<u> </u>
	424 Fe ²⁺ + 106 CO ₂ + 16 NH ₃ + 530 H ₂ O + H ₃ PO ₄		
_	Oxidation by Sulfate		
anoxic	$(CH_2O)_{106}(NH_3)_{16}(H_3PO_4) + 53 SO_4^{2-} \rightarrow$	-380 kJ/mol	
	106 CO ₂ + 16 NH ₃ + 53 S ² + 106 H ₂ O + H ₃ PO ₄		
	Methane Fermentation		
	$(CH_{2}O)_{106}(NH_{3})_{16}(H_{3}PO_{4}) \rightarrow$	-350 kJ/mol	▼
	53 CO ₂ + 53 CH ₄ + 16 NH ₃ + H ₃ PO ₄	7	
		•	least favorable



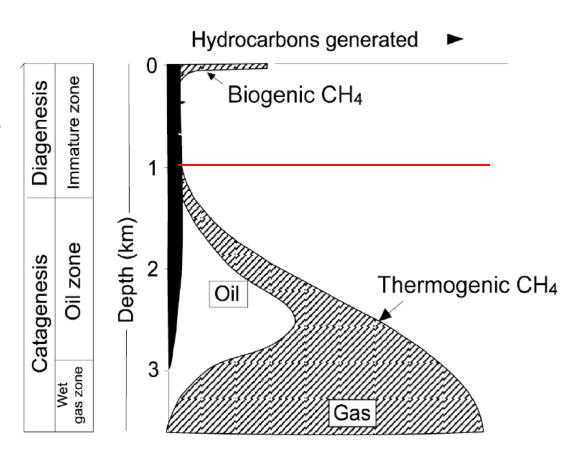




Thermogenic Methane

thermal alteration of organic matter generates methane and higher order hydrocarbons by catagenesis.

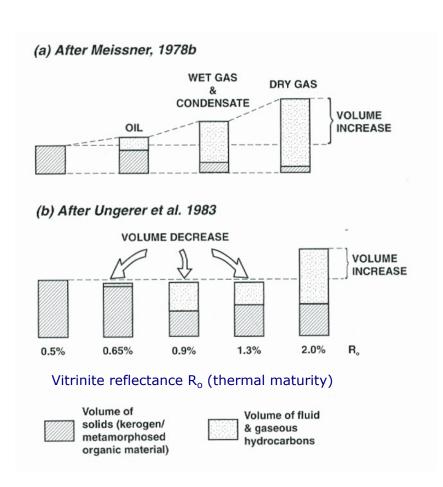
Catagenesis occurs within the temperature range of 50° to 200° C, and gases (methane to butane) are produced at rates that are proportional to temperature.



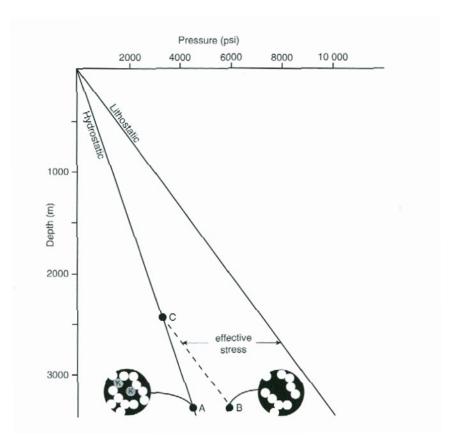
Tissot and Welte, Petroleum Formation and Occurrence, Springer-Verlag, 1992.







Estimation of volume change when Type II Kerogene matures to produce oil and gas.



Estimation of the overpressure created by disequilibrium compaction in a maturing source rock. Half of the initial Kerogene (10% by volume) is transformed into liquid products, thereby increasing the porosity from 13 to 18%. The effect is to transfer the part of the overburden supported by the original kerogene onto the pore fluid.







Factors that determine excess pore water pressure and influence the consolidation state

The most common factors are:

- Stress-related factors
 - Rapid sedimentation rate
 - Tectonic stress
- Fluid volume increase mechanisms
 - Temperature increase
 - Mineral transformation
 - Hydrocarbon Generation
- Fluid movement mechanisms
 - Osmosis
 - Hydraulic head
 - Hydrocarbon buoyancy

(Swarbrick and Osborne, 1998. Bryant et al., 1974; Sangrey, 1977; Arthur et al., 1980; Demaison & Moore, 1980; Bryant et al., 1981).





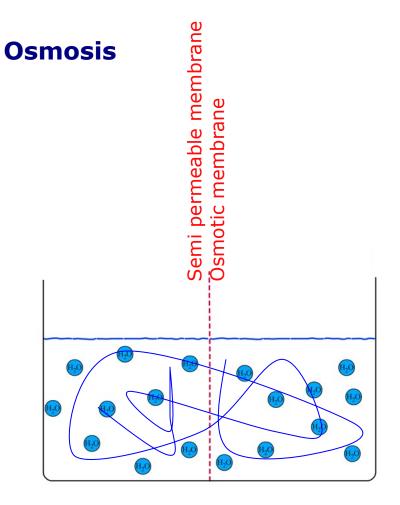
Osmosis

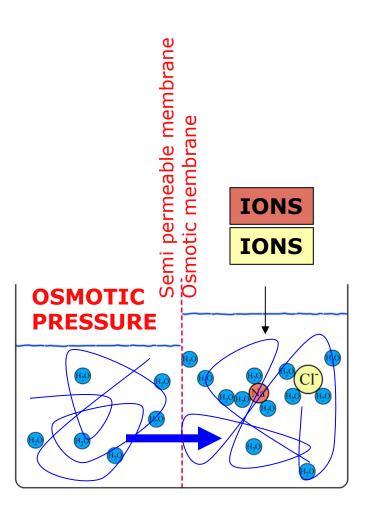
Osmosis is the diffusion of a solvent (e.g. water) through a membrane (osmotic membrane, or semi-permeable membrane) from a solution of low solute concentration to a solution with high solute concentration, up a solute concentration gradient.

A solvent moves without input of energy, across a semi permeable membrane (permeable to the solvent, but not the solute) separating two solutions of different concentrations.













Osmosis

As a consequence of pore water expulsion due to consolidation, pore water salinity often increases with subsurface depth in a sedimentary basin due to

REVERSE OSMOSIS.

Reverse osmosis occurs when a pressure head forces brines through an osmotic membrane, allowing the solvent to pass and retaining the solute.

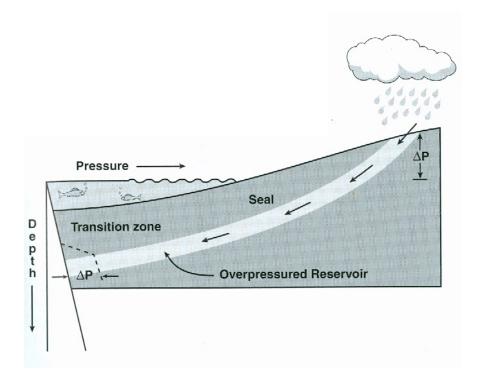
Reverse osmosis is therefore considered as a possible sink of Na⁺ and Cl⁺ ions from seawater.







Hydraulic head







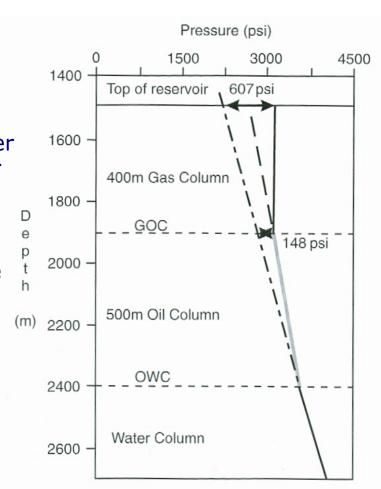


Hydrocarbon buoyancy

- All gasses and oils have lower density than water
- They have a lower pressure gradient than water
- Because overpressure is an excess pressure above hydrostatic for a given depth,

There is always a certain amount of overpressure in a reservoir.

This is a **local** overpressure







APPENDIX. PRESSURE UNITS CONVERSION

The SI unit for **pressure** is the Pascal (Pa)

1 Pa = one Newton per square metre ($N \cdot m^{-2}$ or $kg \cdot m^{-1} \cdot s^{-2}$).

Non-SI measures such as *pound* per square inch (**psi**) and *bar*

1 Pa =
$$10^{-5}$$
 bar = $145.04 \cdot 10^{-6}$ psi

$$1 \text{ bar} = 100,000 \text{ Pa} = 14.504 \text{ psi}$$

1 psi =
$$6,894.76$$
 Pa = $68.948 \cdot 10^{-3}$ Bar

$$1 \text{ bar} = 0.98692 \text{ atmospheres}$$







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