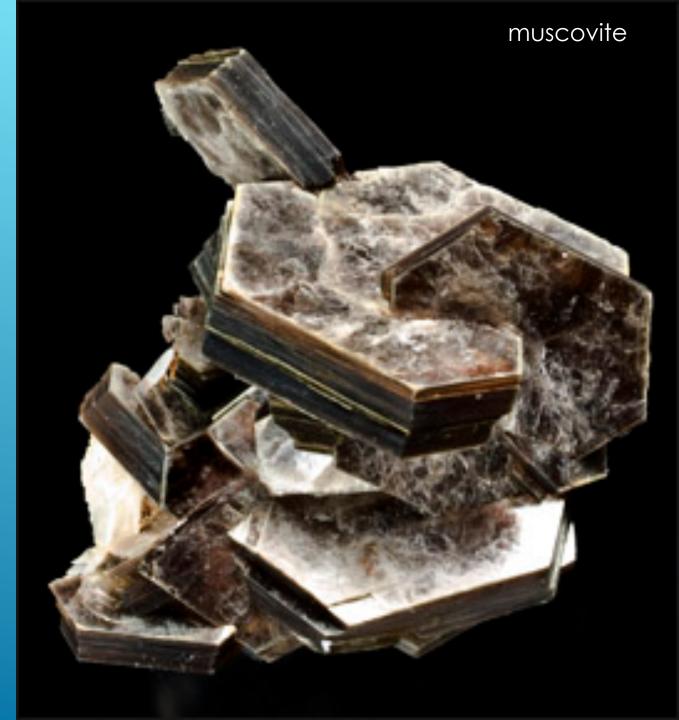
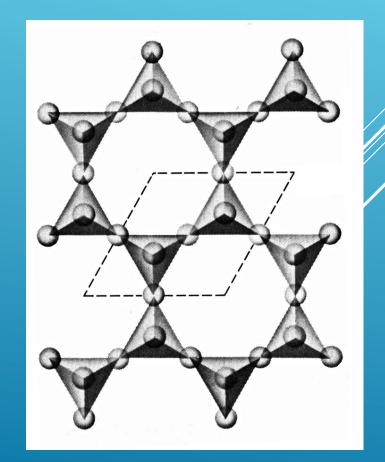
PART 4 SHEET SILICATES



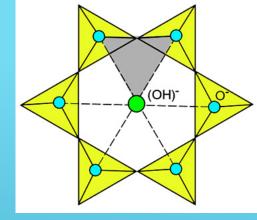
SHEET SILICATES = PHYLLOSILICATES

- > "Phyllon" = leaf
- Large group of mineral including many common minerals: muscovite, biotite, serpentine, chlorite, talc, clay minerals
- Structure: interconnected rings (6-fold) of silica tetrahedra that extend outwards infinite sheet: 3 of the 4 oxygen in the silica tetrahedra are shared with other tetrahedra \Rightarrow basic structural unit Si₂O₅²



SHEET SILCATES

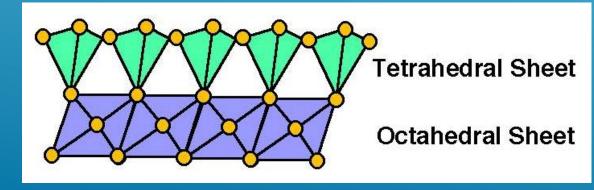
- Most sheet silicates: hydrous
- Most phyllosilicates: OH- located at the center of the rings ⇒ basic structural unit: Si₂O₅(OH)³⁻
- Groups: based on structure
- Two kind of "layers" within the "sheet":
 - I layers: tetrahedral sites: tetrahedral coordination of Si and Al
 - Olayers: octahedral sites: mostly Al and Mg, occasionally Fe
- T and O layers bounded to form sheet The space between sheets can be:
 - Vacant
 - **Filled with interlayer cations**, water or other sheets



SHEET SILCATES

 Different stacking arrangements of tetrahedral sheets and octahedral sheets, along with the type of cation that occupies the octahedral site, allow for the variety of phyllosicilicates that occur in nature.

Stacking of the sheets



Tetrahedral Sheet Octahedral Sheet Tetrahedral Sheet

SHEET SILCATES

Octahedral sheet variety:

Dioctahedral:

- > cations are trivalent (Al³⁺ or Fe³⁺)
- One every three sites is vacant
- > OH and O are bounded to 2 cations

► Trioctahedral

- Cations are divalent (Mg²⁺ or Fe²⁺)
- > All cation sites are filled
- > OH and O are bounded to 3 cations

COMMON SHEET SILCATES

► T-O Phyllosilicates Si2O5:

- Dioctahedral: kaolinite Al₂Si₂O₅(OH)₄
- Trioctahedral:Serpentine (Mg,Fe)₃Si₂O₅(OH)₄

► T-O-T phyllosilicates Si4O10

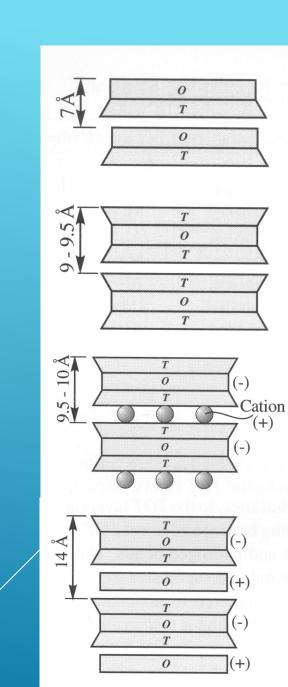
- Dioctahedral: pyrophyllite Al₂Si₄O₁₀(OH)₂
- Trioctahedral: talc (Mg,Fe)₃Si₄O₁₀(OH)₂

T-O-T + interlayer cations: the micas

- Dioctahedral: muscovite KAl₂Si₃Al₁O₁₀(OH)₂
- Trioctahedral: biotite K(Mg,Fe)₃Si₃Al₁O₁₀(OH)₂

T-O-T + interlayer octahedral sheet: chlorite

- Di-dioctahedral [AI(OH)₃]Al₂(Si₄)O₁₀(OH)₂
 Di-Trioctahedral: [AI(OH)₃](Mg,Fe)₃(Si₄)O₁₀(OH)₂



SERPENTINE GROUP (T-O STUCTURE)

- \succ Trioctahedral T-O Phyllosilicates: (Mg,Fe)₃Si₂O₅(OH)₄
- S varieties: Antigorite, Lizardite and Chrisotile
 Massive fine grains
 Fibrous
 Antigorite
 Tetrahedral Layers
 Octahedral Layers
- Where? Serpentine = hydration product of Mg-rich silicates (pyroxene, olivine)

 $2Mg_2SiO_4 + 3H_2O \iff Mg_3Si_2O_5(OH)_4 + Mg(OH)_2$

Olivine water Serpentine Brucite

- Pseudomorph after olivine and pyroxene in alterated basic and ultrabasic igneous rocks (peridotite, dunite, basalt, gabbro) - often associated with minerals magnesite (MgCO₃), chromite, and magnetite.
- Rock made up almost entirely of serpentine, it is called a serpentinite.

TALC (T-O-T STUCTURE)

- > Trioctahedral T-O-T Phyllosilicates: $(Mg,Fe)_3Si_4O_{10}(OH)_2$
- Low hardness
- Where? low grade metamorphic rocks that originated as ultrabasic to basic igneous rocks. Rock made up almost entirely of talc is called a socpsione.
 Ex.: Hydrothermal solutions concentrated during final stages of magma crystallization in batholiths or hot seawater solutions drawn down into subduction zones
- Use: in paint, plastics, rubber, various roofing compounds: as lubricant
 - in cosmetic and body powder.
 - as "whiskistone"



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MICA GROUP (T-O-T + C STUCTURE)

> White micas (dioctahedral) vs. black micas (trioctahedral)

Muscovite, Paragonite, and Margarite

Biotite and Clintonite

> White micas:

Muscovite, KAI₃Si₃O₁₀(OH)₂, and Paragonite, NaAI₃Si₃O₁₀(OH)₂: end members of the solid solution series involving K and Na but large immiscibility gap (solvus): Muscovite = 65 to 100% K vs paragonite = 80 to 100% Na

Where?

- > Al-rich medium grade metamorphic rocks (Al-rich schist)
- in siliceous, Al-rich plutonic igneous rocks (but not in volcanic rock) association with alkali feldspar, quartz, and sometimes biotite, garnet, andalusite, sillimanite, or kyanite.
- Substitution of AI by Li in the interlayer cation of muscovite: Lepidolite pink mica found in pegmatite

MICA GROUP (T-O-T + C STUCTURE)

> White micas (dioctahedral) vs. black micas (trioctahedral)

Muscovite, Paragonite, and Margarite

Biotite and Clintonite

Black micas

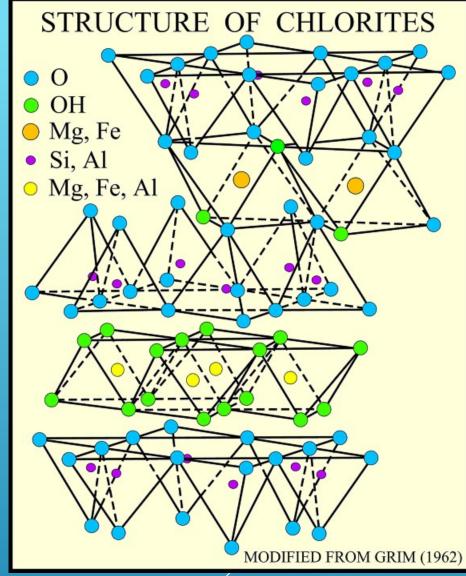
- Biotite is a solid solution between the end members Phlogopite KMg₃AlSi₃O₁₀(OH)₂ and Annite KFe₃AlSi₃O₁₀(OH)₂' (but pure Annite does not occur in nature)
- Substitutions: minor Na, Rb, Cs, and Ba may substitute for K
 minor F can substitute for OH: increase the stability at high temperature

> Where?

- > Phlogopite: in hydrous ultrabasic volcanic rocks (kimberlite), in metamorphosed dolomite
- Biotite: dacitic, rhyolitic, and trachytic volcanic rocks, granitic plutonic rocks, and a wide variety of metamorphic rocks.

CHLORITE (T-O-T + O LAYER)

- structure that consists of phlogopite T-O-T layers sandwiching brucite-like octahedral layer.
- Substitutions:
 - Mg for Fe
 - Al for (Mg, Fe) in both the octahedral sites
 - Al for Si
- Where? low grade metamorphic rocks associated with actinolite, epidote, and biotite. It also forms as an alteration product of pyroxenes, amphiboles, biotite, and garnet in igneous as well a metamorphic rocks.



- Products of chemical weathering main constituent of mudrock (mudstone, claystone, shale)
- > 40% minerals in sedimentary rocks
- Understanding their behavior: important economic use (ceramic), civil engineering (clay swelling)
- Structural classification:
 Kandites (struc. ⇔ Kaolinite) Smectites (struc. ⇔ Pyrophyllite) Illites (struc. ⇔ muscovite)
 T-O
 T-O-T
 T-O-T+c

- > **Chemical Weathering:** Minerals form in depth are not stable at the surface of Earth:
 - Lower T (-20 to 50°C)
 - Lower P (1 to few hundred atmospheres)
 - > Higher free water
 - Higher free O
 - > Stability order at the near surface conditions:

Iron oxide – Aluminium oxide, quariz, clay mineral, Muscovite, Alkali feldspar, biolite,

Amphibole, Pyroxenes, Ca-rich plagioclase, Olivine

Chemical Weathering – Main agent: water and weak acids in water

- Acid in solution: abundant free H⁺
- Most common acid: Carbonic acid: produced by reaction between rainwater and carbon dioxide in atmosphere:

H ₂ O	+ CO ₂	\rightarrow	$H_2CO_3 \rightarrow$	H+	+	HCO3-
Water	carbon dioxide		carbonic acid	hydrogen ion		bicarbonate ion

> Type of chemical weathering reactions:

► Hydralysis: H+ or OH- replaces an ion in the mineral4KAISi_3O_8 +4H^++ $2H_2O \rightarrow 4K^+$ + $AI_4Si_4O_{10}(OH)_8$ + $8SiO_2$ Orthoclasehydrogen ionwaterpotassium ionkaolinitequartz► OxideIIon: reaction of mineral with O2: change of the oxidation state (Fe²⁺ to Fe³⁺)3Fe²⁺SiO3 + $1/2O_2 \rightarrow$ Fe₃O₄+ $3SiO_2$ Pyroxeneoxygenmagnetitequartz► Dehydralion - Removal of H₂O or OH- ion from a mineral2FeO·OH →Fe₂O₃+H₂OGoethitehematitewater

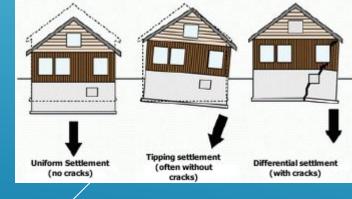
Complete dissolution (in water)

Kandites (T-O structure)

- ► Most common: kcolinite Al₂Si₂O₅(OH)₄
- > Other kandites: Anauxite, Dickite, Nacrite
- ► Where? Weathering of hydrothermal alteration of aluminosilicate minerals ⇒ feldspar-rich rocks (ex.: granitic rocks) weather to kaolinite

Smectites (T-O-T structure)

- Structure similar to pyrophyllite Al₂Si₄O₁₀(OH)₂
- > Possible substitution of AI with Mg or Fe \Rightarrow di- and trioctahedral
- Possibility to absorb water between the TOT sheets: cause volume of the mineral to increase in contact with water : expanding clays
- Most common: Montmorillinite (½Ca,Na)(Al,Mg,Fe)₄(Si,Al)₈O₂₀(OH)₄·nH₂O = main constituent of Bentonite, derived by weathering of volcanic ash.



Illites (T-O-T+c structure)

- \succ K_yAl₄(Si_{8-y},Al_y)O₂₀(OH)₄ usually with 1 < y < 1.5, but always with y < 2.
- Possible substitution of K by Ca or Mg (to preserve balance)
- > The most common in soils.
- Not an expanding clay
- Where? formed from weathering of K and Al-rich rocks under high pH conditions of minerals like muscovite and feldspar. Main constituent of ancient mudrocks and shales.

Mixed layer clays

- = change from one type to another through a stacking sequence
- > Common, regular and ordered or irregular and unordered
- Identification? To thin/small to be recognize in hand samples or petrographic microscope
 ⇒ use of the X-ray technique