

Soil Mineralogy

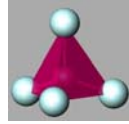
- Primary minerals—
 - inherited from parent material
 - igneous and metamorphic
 - sand and silt fraction (mostly)
- Secondary minerals—
 - formed under “normal” temperature and pressure
 - formed in the soil environment or inherited from sedimentary parent material
 - clay fraction (mostly)

Pauling Rules

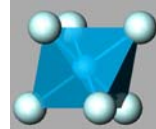
- 1. A coordinating polyhedron of anions is found about each cation, the cation-anion distance being determined by the radius sum and the coordination number of the cation by the radius ratio.
- 2. In a stable coordination structure, the total strength of the valency bonds that reach an anion from all the neighboring cations is equal to the charge of the anion.

Coordination Environment

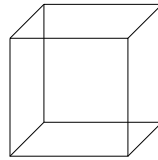
■ Tetrahedral



■ Octahedral



■ Cubic



■ Dodecahedral

Radius ratio

- Tetrahedral (4): 0.225-0.414
 - Si^{4+} , Al^{3+}
- Octahedral (6): 0.414-0.732
 - Fe^{3+} , Mg^{2+} , Ti^{4+} , Fe^{2+} , Mn^{2+} , Na^{+} , Ca^{2+}
- $\text{Si}^{4+} = 0.042 \text{ nm}$, $\text{O}^{2-} = 0.140 \text{ nm}$
 - ratio = 0.300
- $\text{Al}^{3+} = 0.051 \text{ nm}$, ratio = 0.364
- $\text{Mg}^{2+} = 0.066$, ratio = 0.471

More Pauling Rules

- 3. The existence of edges, and particularly faces, common to two anion polyhedra in a coordinated structure decreases its stability; this effect is large for cations with high valency and small coordination number.
- 4. In a crystal containing different cations, those of high valency and small coordination number tend not to share polyhedron elements with each other.
- 5. The number of essentially different kinds of constituents in a crystal tends to be small.

What does this mean?

- Sharing edges or faces brings the cations closer together and causes repulsion
- Cations such as P^{5+} will not form minerals that share any O (in nature).
- The most stable Si minerals share all the corner O but no edges or faces (quartz).
- Different cations force different geometry and result in less stable minerals.

Building Blocks



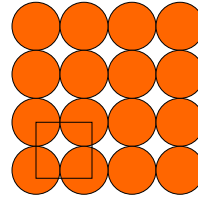
- Si tetrahedra and Al octahedra are the basic building blocks of the clay minerals.
- <http://virtual-museum.soils.wisc.edu/>

Types of silicate minerals

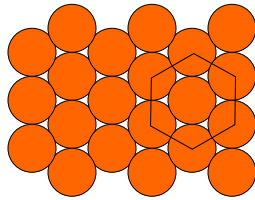
- The different minerals are a results of differences in the sharing of O and the presence or absence of different cations.
- Range from no sharing (nesosilicates) to sharing of each O with 2 Si (tectosilicates)
- Zeolites are an interesting form of tectosilicates.

Basic Structural Concepts

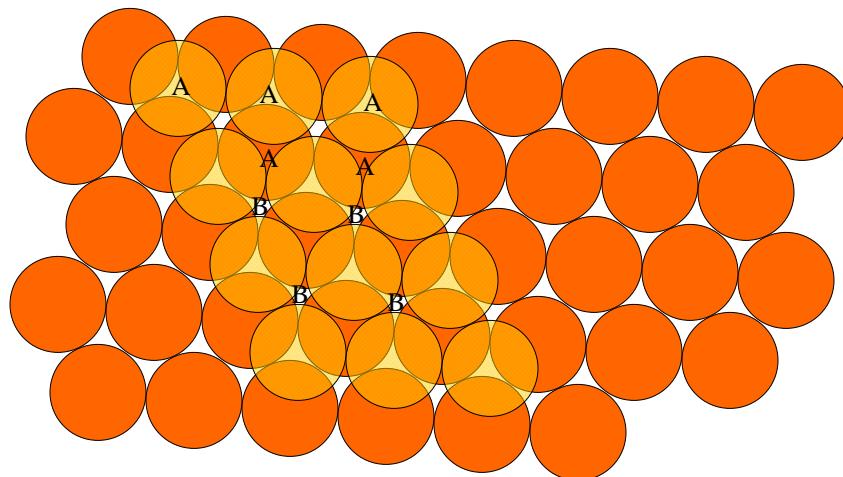
■ Cubic Close Packing (CCP).



■ Hexagonal Close Packing (HCP).



HCP

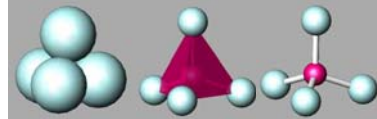


HCP

- Two types of voids:

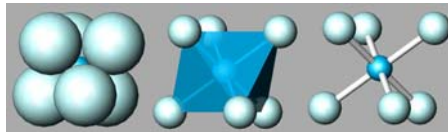
- A - void constrained by 4 spheres

- Tetrahedral coordination

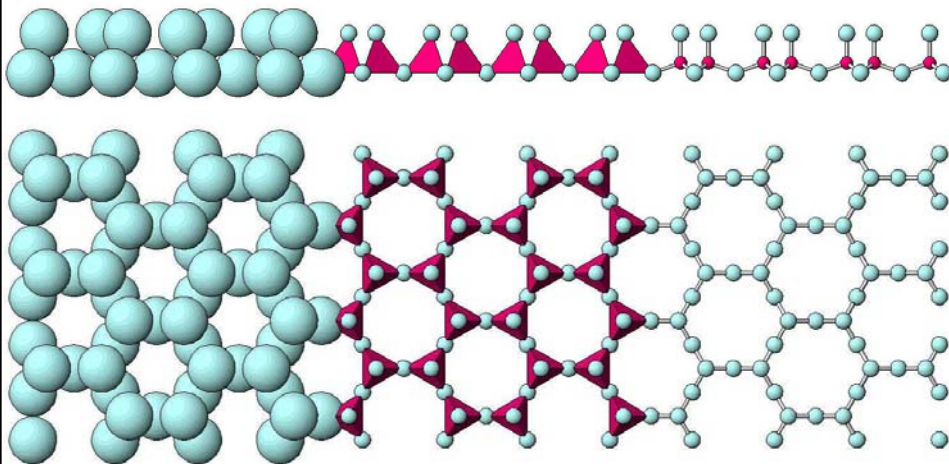


- B - void constrained by 6 spheres

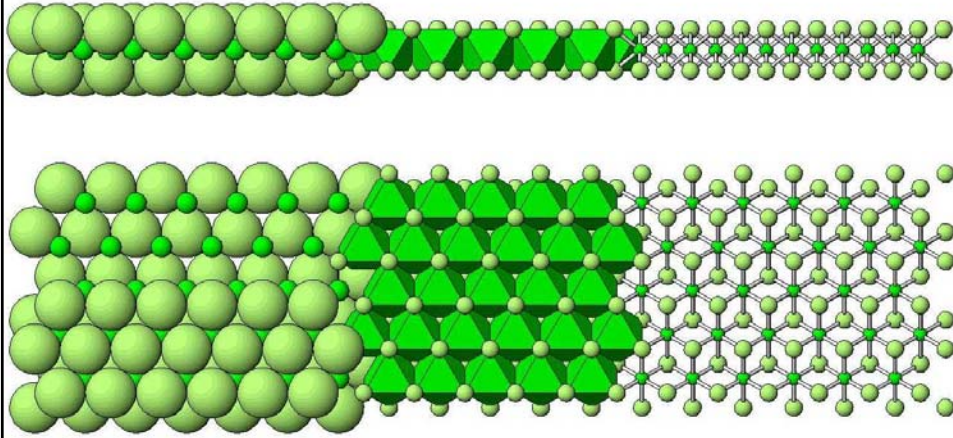
- Octahedral coordination



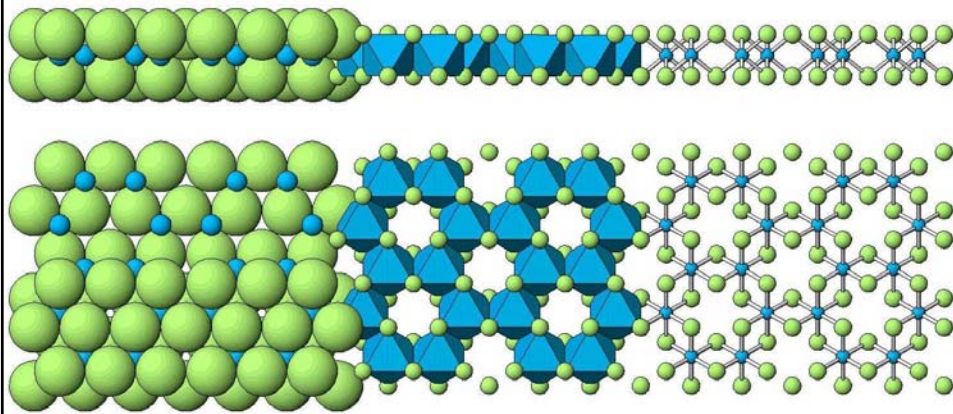
Tetrahedral Sheet



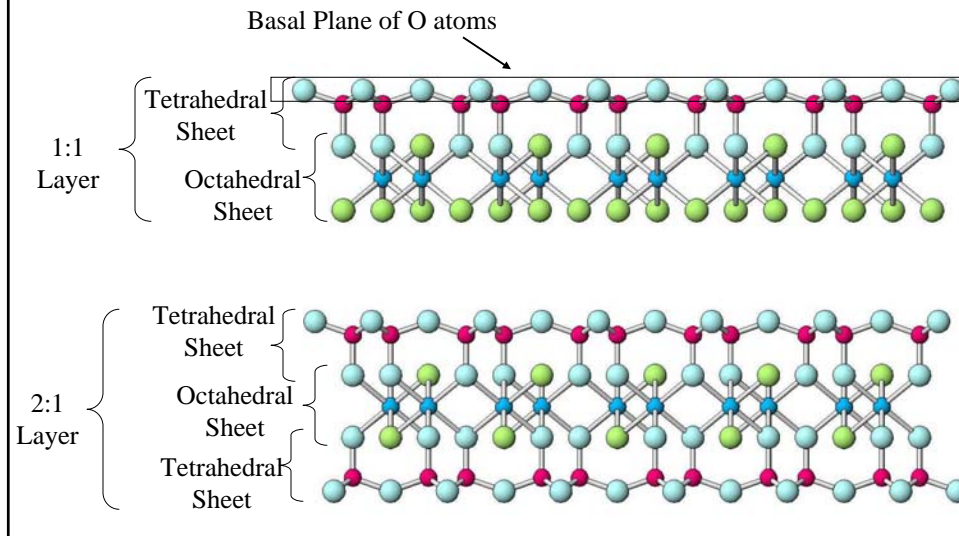
Trioctahedral Sheet



Dioctahedral Sheet



Terminology



Classification of Clay Minerals

- Layer Silicates (phyllosilicates)
 - Number of octahedral and tetrahedral sheets (1:1, 2:1, 2:1:1).
 - Type of octahedral sheet (dioctahedral or trioctahedral).
 - Type, extent, and location of isomorphous substitution
- Oxides, hydroxides, and oxyhydroxides (Fe, Mn, Al)