

Liquids and Solids

Relative Magnitudes of Forces

The types of bonding forces vary in their strength as measured by average bond energy.

Strongest
Weakest



Covalent bonds (400 kcal/mol)

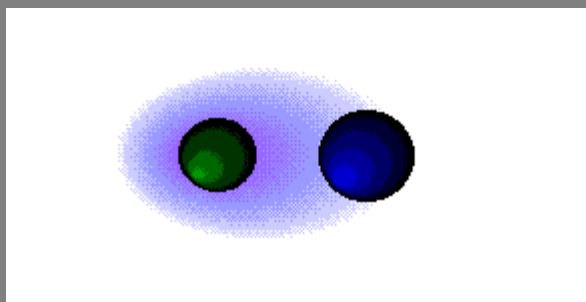
Hydrogen bonding (12-16 kcal/mol)

Dipole-dipole interactions (2-0.5 kcal/mol)

London forces (less than 1 kcal/mol)

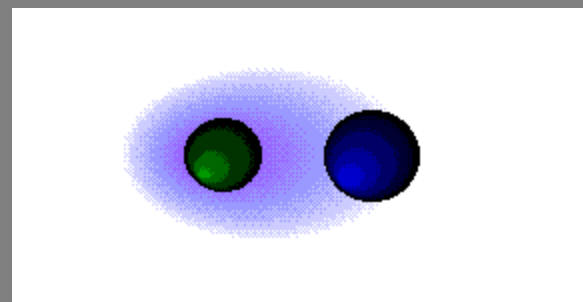
Dipole-Dipole Forces

Attraction between the positive end of one molecule and the negative end of another. Strong attractions, but not as strong as hydrogen bonding. These are only found in polar molecules.



δ^- charge

δ^+ charge

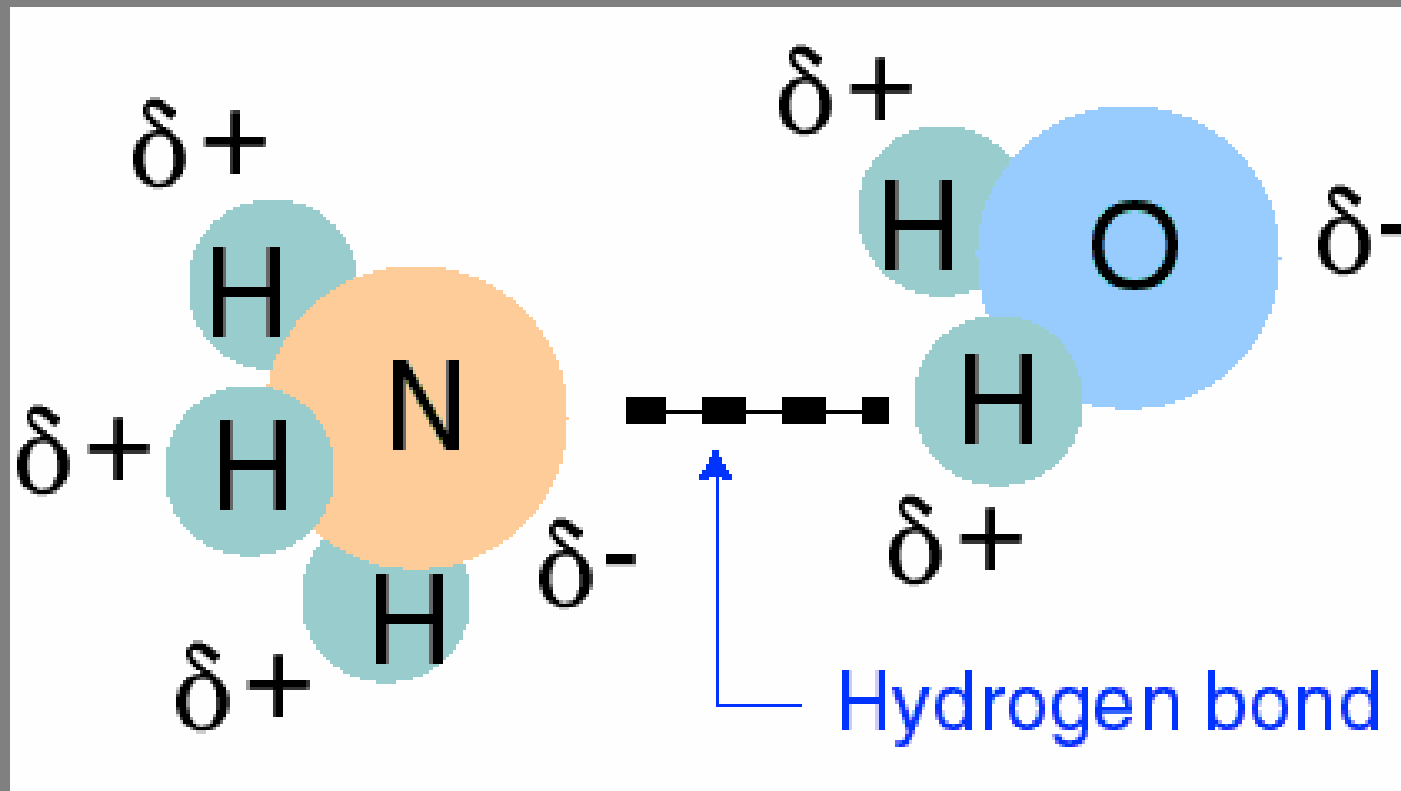


δ^- charge

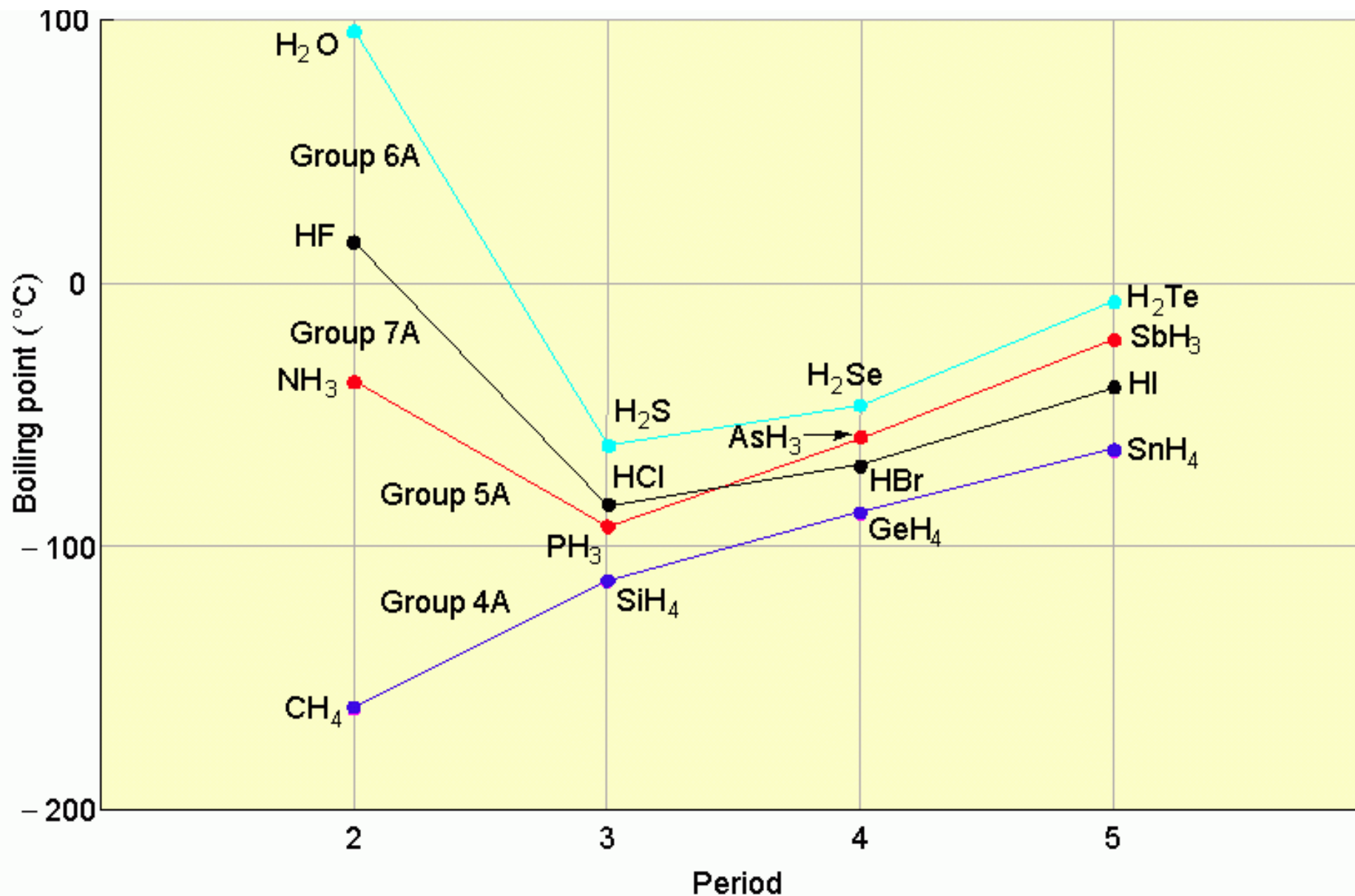
δ^+ charge

Hydrogen Bonding

Bonding between hydrogen and more electronegative neighboring atoms such as oxygen and nitrogen



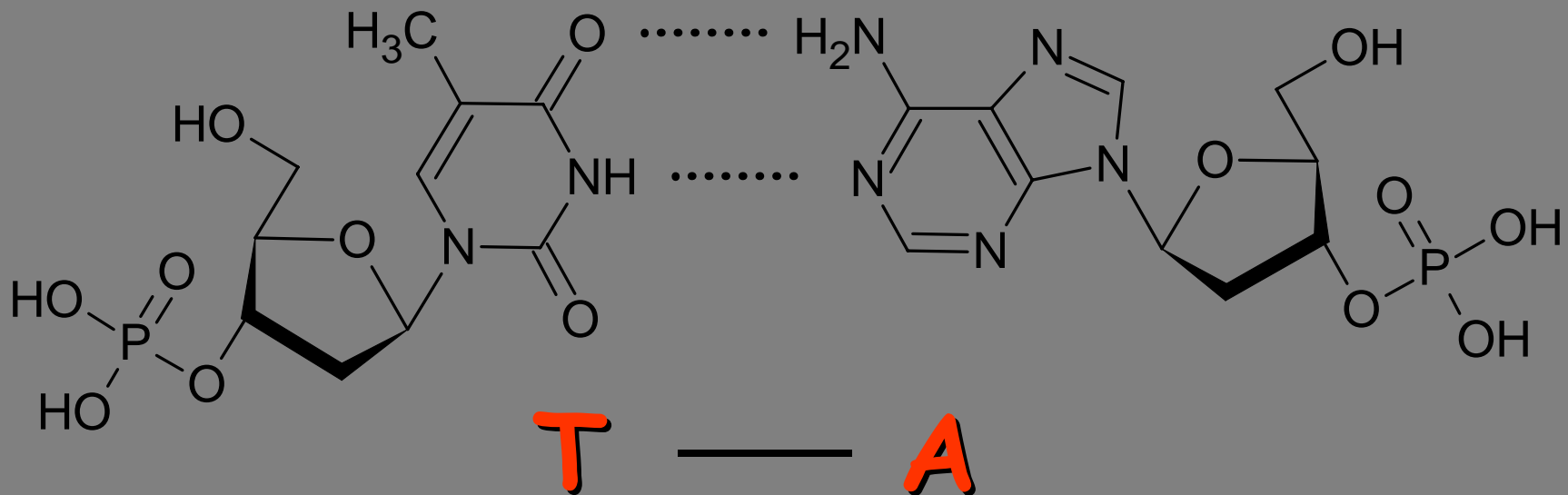
Hydrogen bonding between ammonia and water



Boiling point as a measure of intermolecular attractive forces

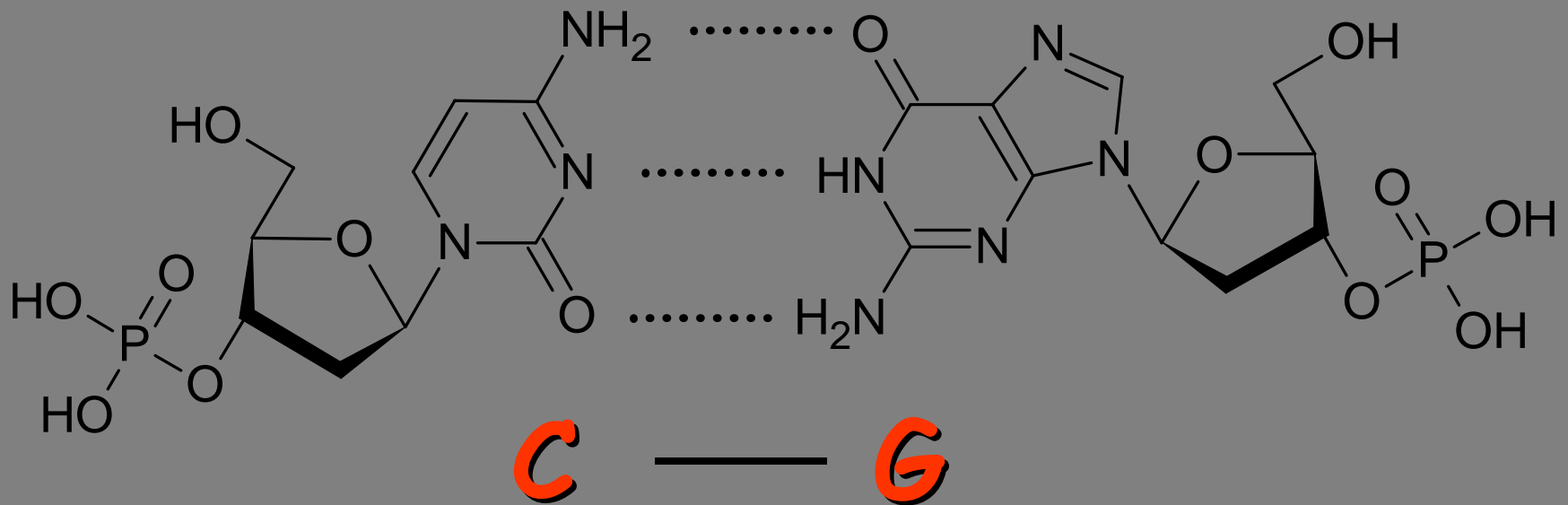
Hydrogen Bonding in DNA

Thymine hydrogen bonds to Adenine



Hydrogen Bonding in DNA

Cytosine hydrogen bonds to Guanine



Factors making Hydrogen Bonding

Special strong dipole-dipole attraction

1. The small size of the elements F,N,O
 2. The relatively high electronegativity of them.
- This allows for the creation of strong dipole forces of attraction between the molecules. The effect of which can be seen on the boiling point of substances that contain hydrogen bonding.

London Dispersion Forces



Fritz London
1900-1954

The temporary separations of charge that lead to the London force attractions are what attract one **nonpolar** molecule to its neighbors.

London forces increase with the size of the molecules.

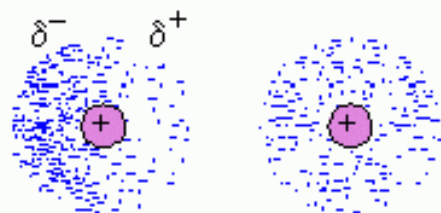


London Dispersion Forces



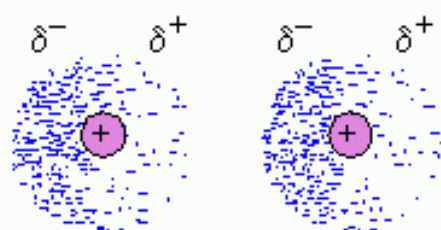
Atom A Atom B

No polarization

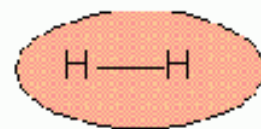


Atom A Atom B

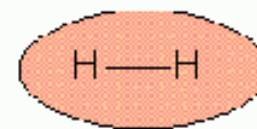
Instantaneous dipole on atom A
induces a dipole on atom B



Atom A Atom B

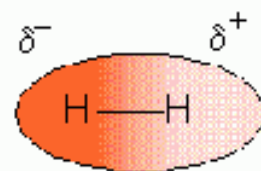


Molecule A

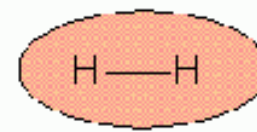


Molecule B

No polarization

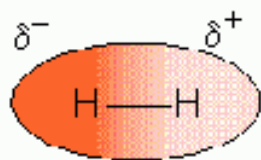


Molecule A

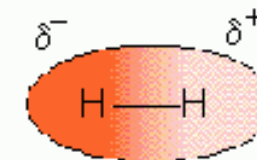


Molecule B

Instantaneous dipole on molecule A
induces a dipole on molecule B

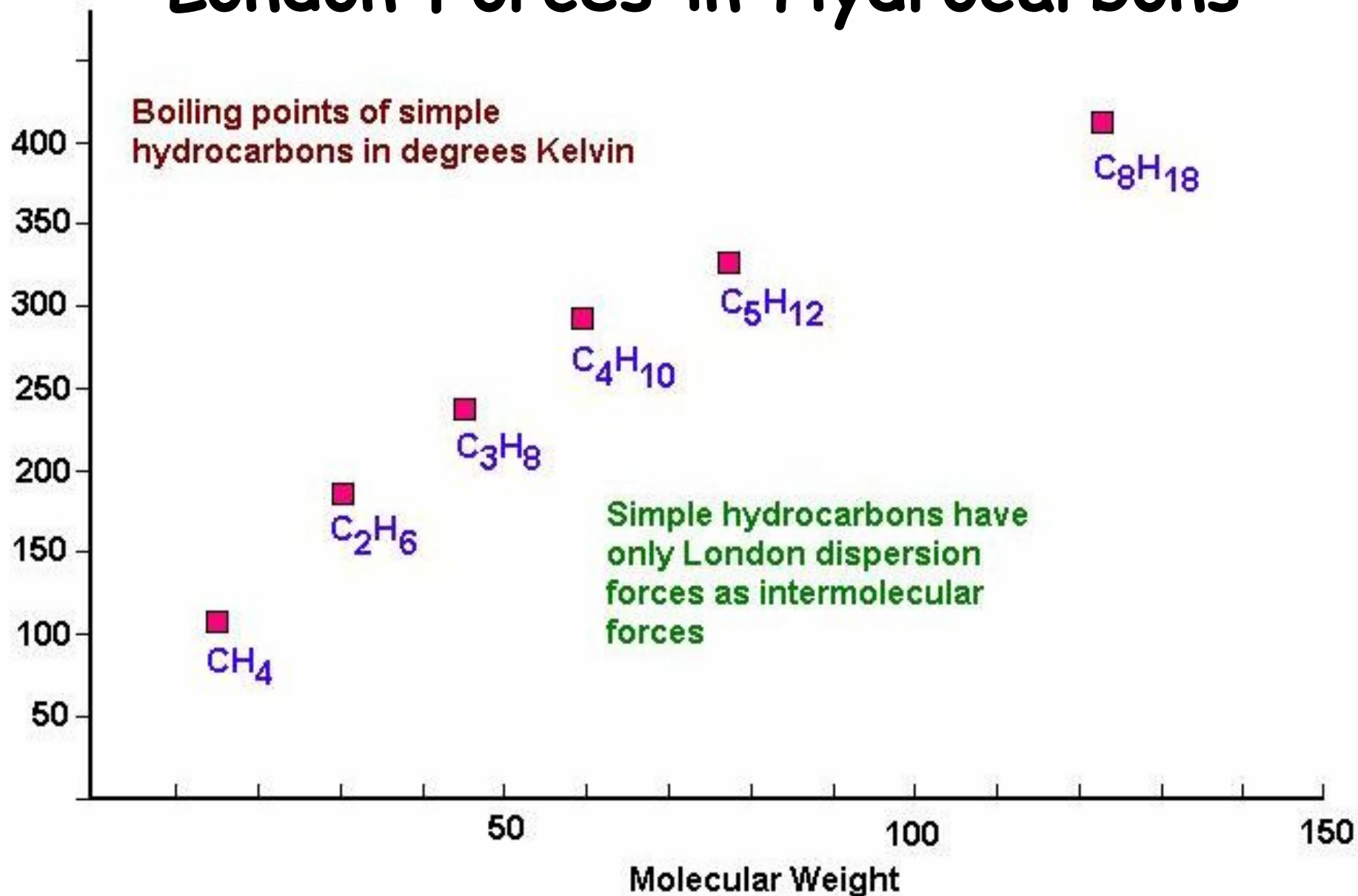


Molecule A



Molecule B

London Forces in Hydrocarbons

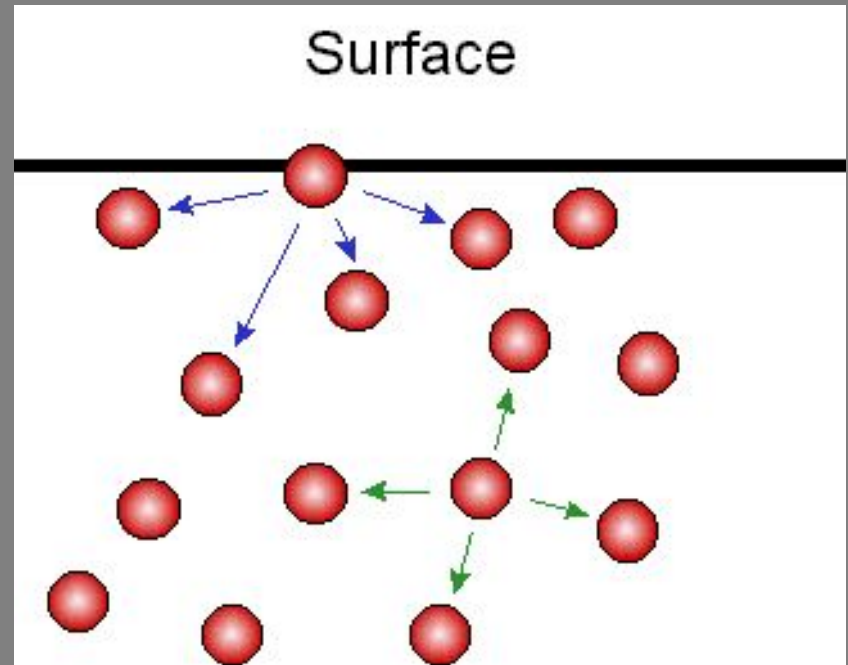


Summary of intermolecular forces

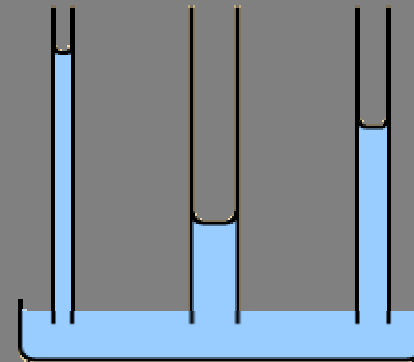
Hydrogen bonding	Dipole-dipole	LDF (London Dispersion Forces)
Occurs between molecules that are polar and have H-F, H-O or H-N bonds.	Occurs in polar molecules that do not have hydrogen bonding . Weaker than H-bond	Occurs in non-polar molecules . The weakest interaction.
Caused by attraction of highly electronegative and small F,O,N with hydrogen.	Caused by attraction of oppositely charged poles of molecules	Caused by attraction of atoms when their electrons are unequally distributed around the molecule for an instant of time.
Strongest	strong	Weak - gets stronger with more electrons. ex. He-weak Ar-Stronger

Some Properties of a Liquid

- ❖ **Surface Tension:** The resistance to an increase in its surface area (polar molecules, liquid metals).



- ❖ **Capillary Action:** Spontaneous rising of a liquid in a narrow tube.



Some Properties of a Liquid

❖ **Viscosity:** Resistance to flow

❖ **High viscosity** is an indication of strong intermolecular forces



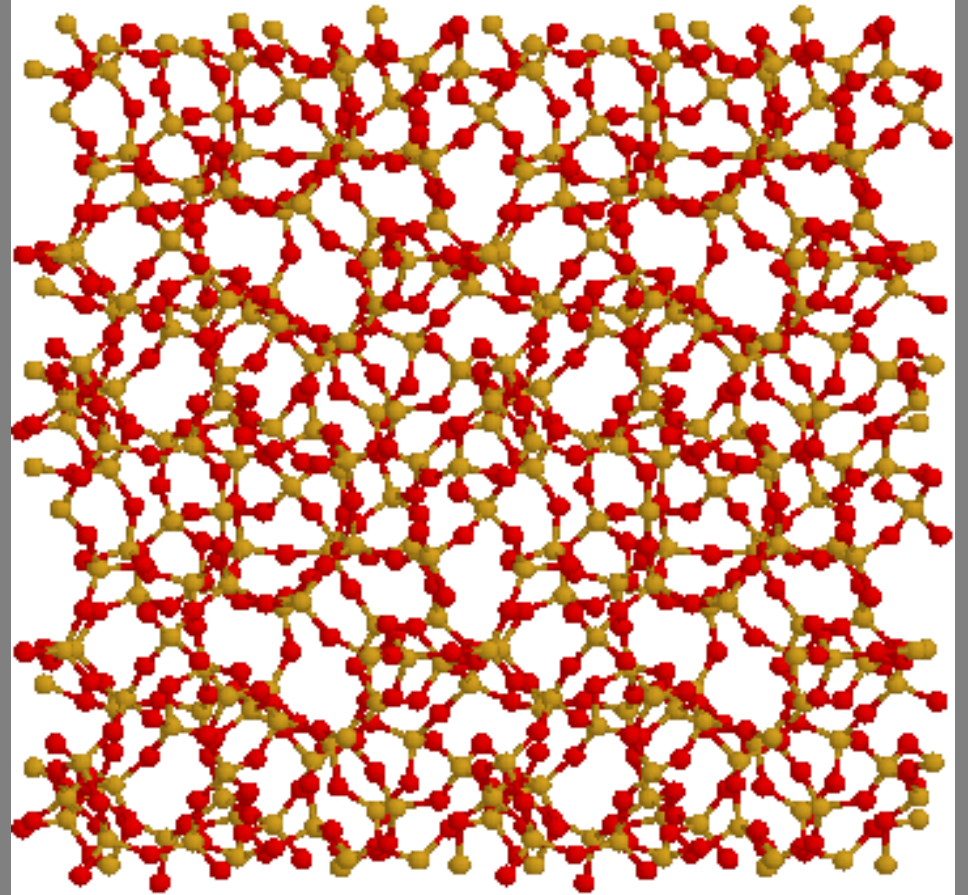
Types of Solids

- ❖ **Crystalline Solids:** highly regular arrangement of their components



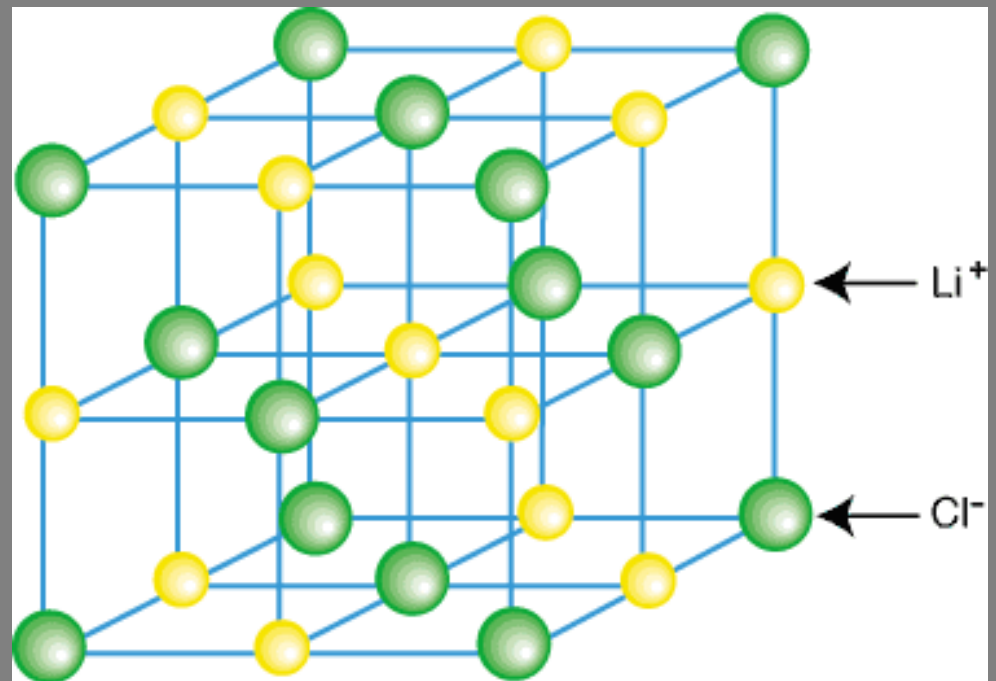
Types of Solids

- ❖ **Amorphous solids:** considerable disorder in their structures (glass).



Representation of Components in a Crystalline Solid

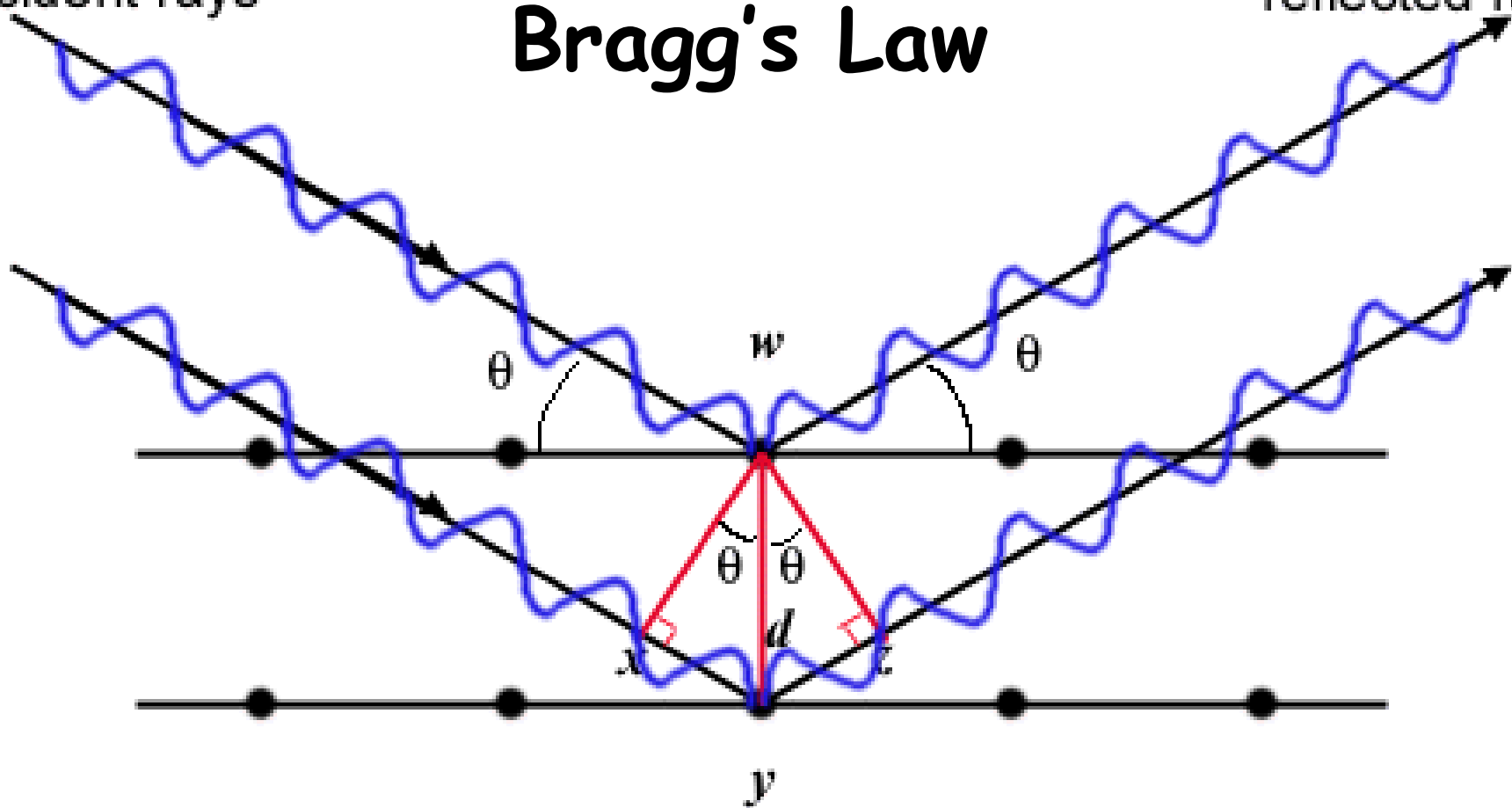
Lattice: A 3-dimensional system of points designating the centers of components (atoms, ions, or molecules) that make up the substance.



incident rays

Bragg's Law

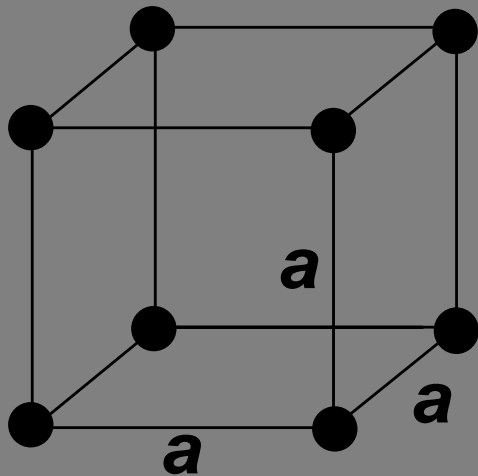
reflected rays



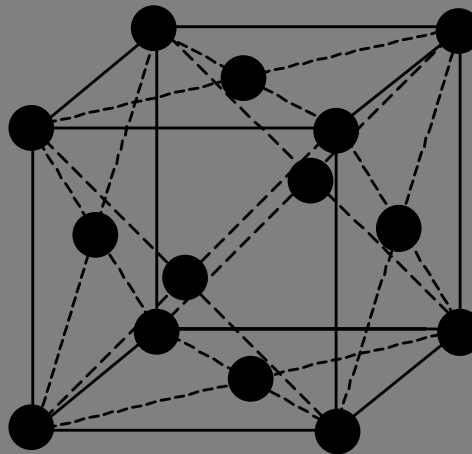
$$xy + yz = n\lambda \quad \text{and} \quad xy + yz = 2d \sin\theta$$

$$\therefore n\lambda = 2d \sin\theta$$

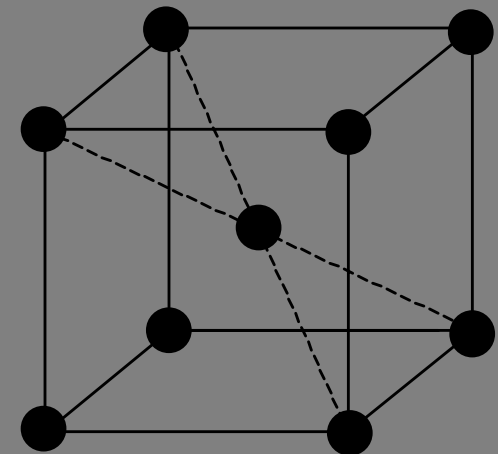
Crystal Structures - Cubic



Simple

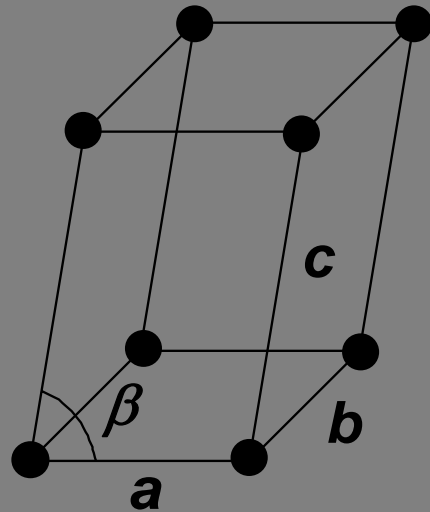


Face-Centered

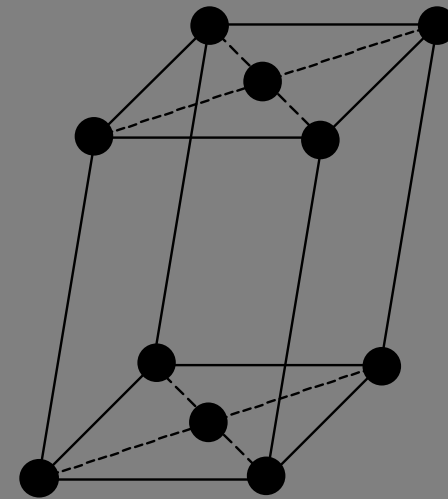


Body-Centered

Crystal Structures - Monoclinic

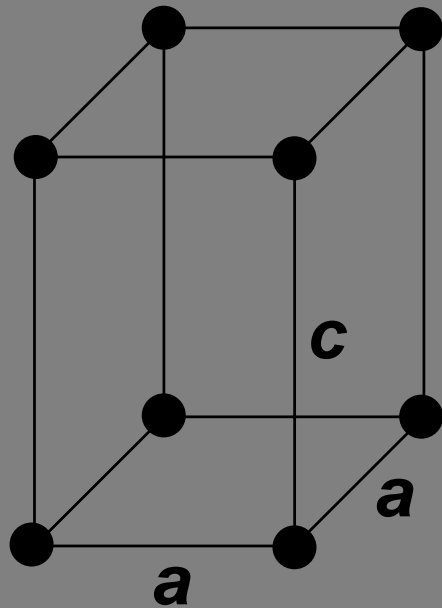


Simple

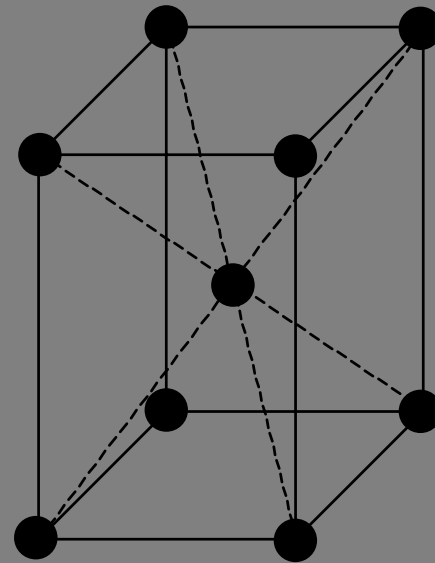


End Face-Centered

Crystal Structures - Tetragonal

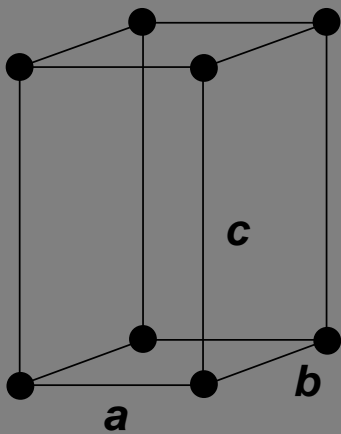


Simple

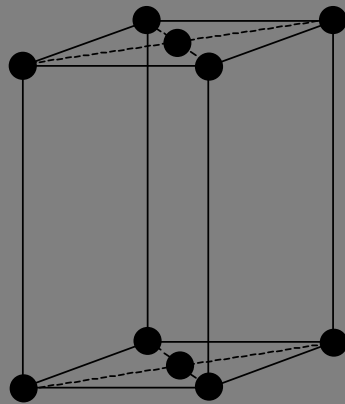


Body-Centered

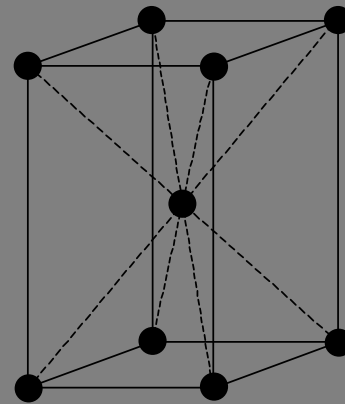
Crystal Structures - Orthorhombic



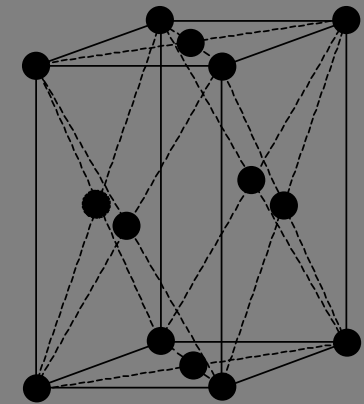
Simple



*End
Face-Centered*

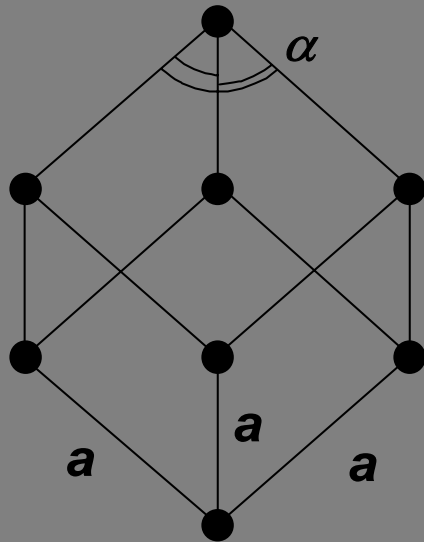


*Body
Centered*

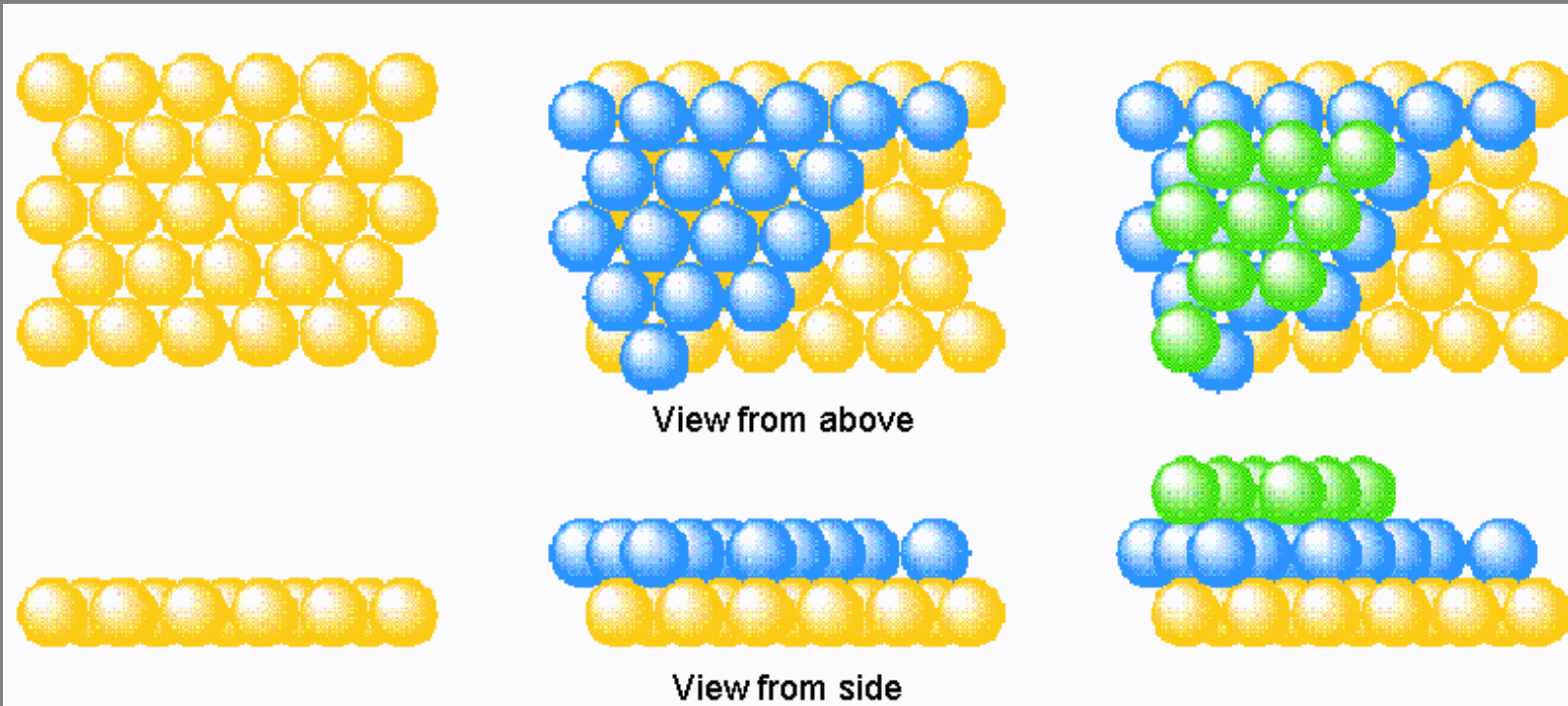


*Face
Centered*

Crystal Structures - Other Shapes

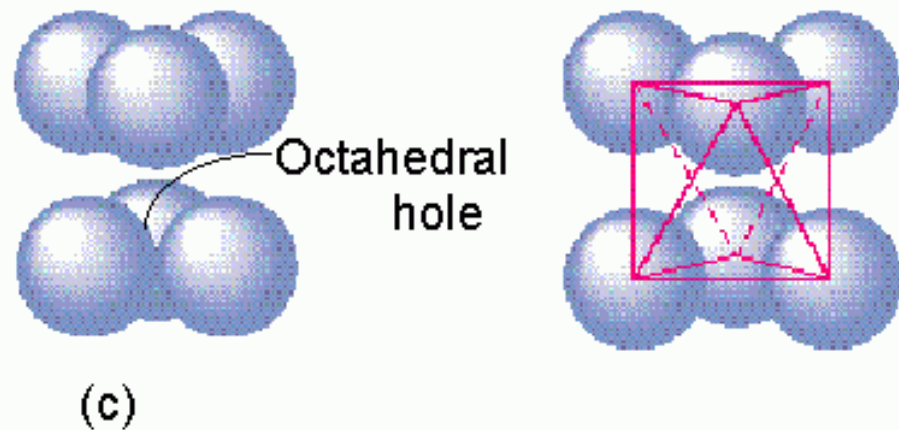
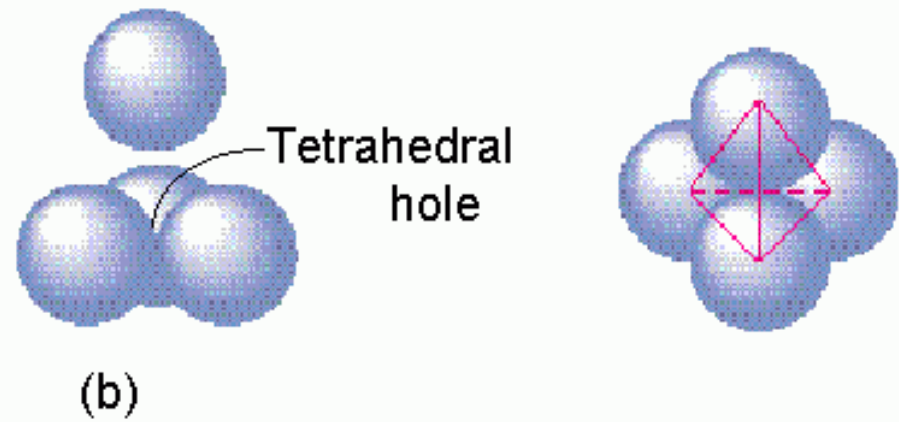
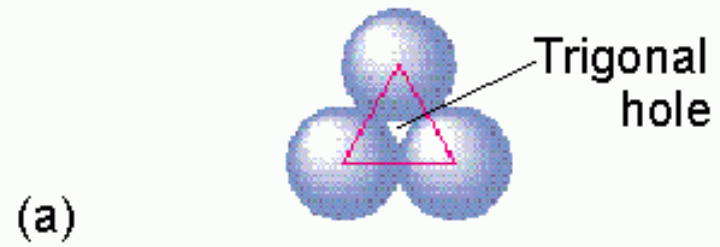


Packing in Metals



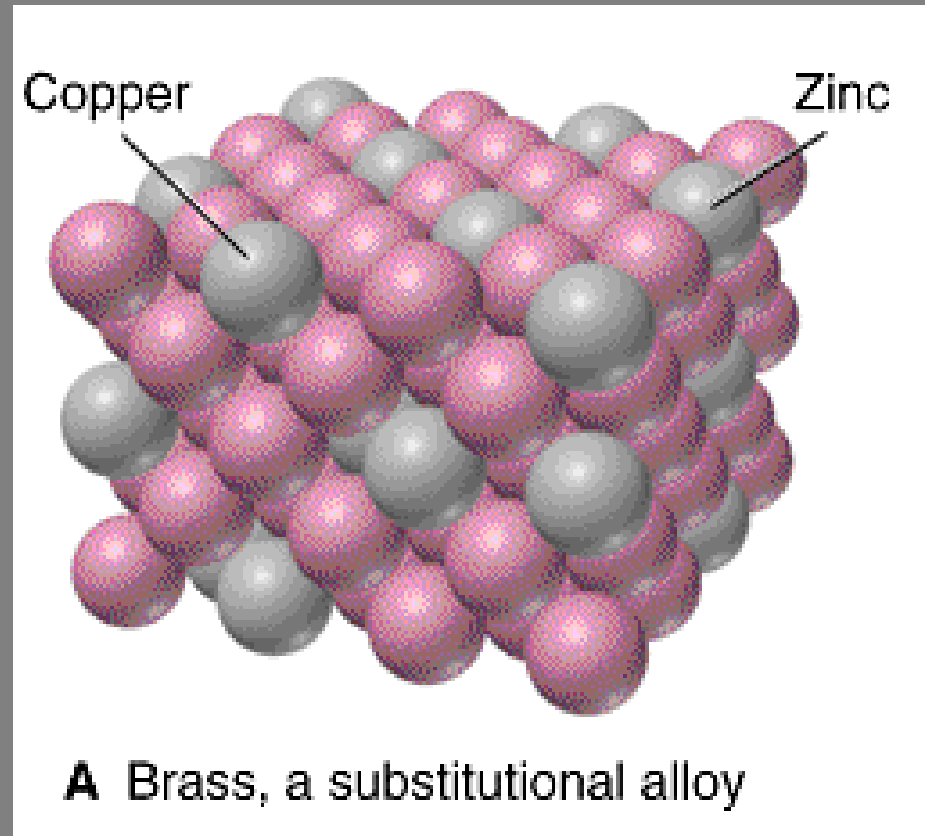
Model: Packing uniform, hard spheres to best use available space. This is called **closest packing**. Each atom has 12 nearest neighbors.

Closest Packing Holes



Metal Alloys

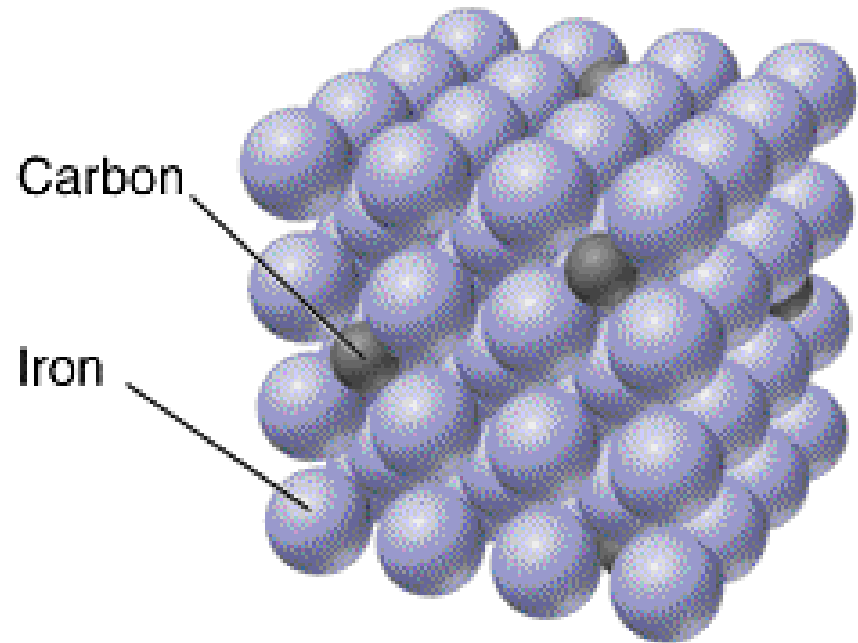
- ❖ **Substitutional Alloy:**
some metal atoms replaced by others of similar size.
 - brass = Cu/Zn



Metal Alloys

(continued)

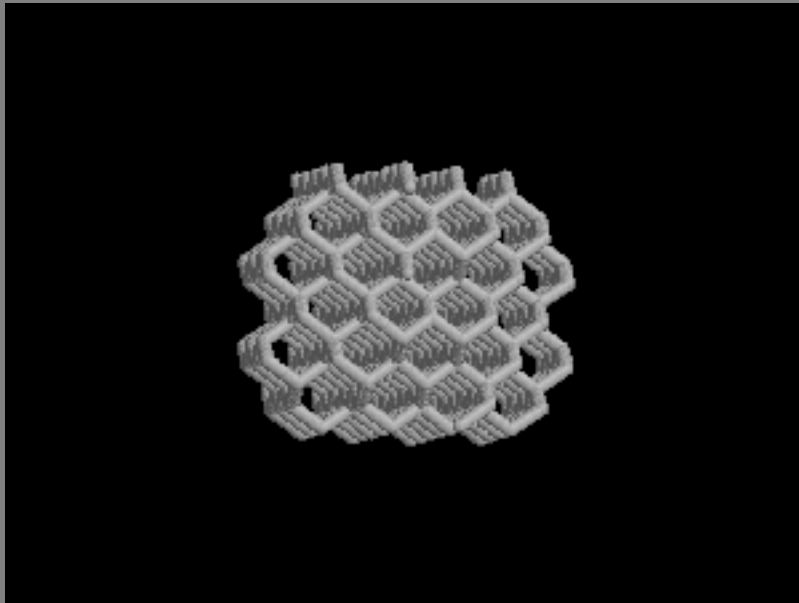
- ❖ **Interstitial Alloy:**
Interstices (holes) in closest packed metal structure are occupied by small atoms.
- steel = iron + carbon



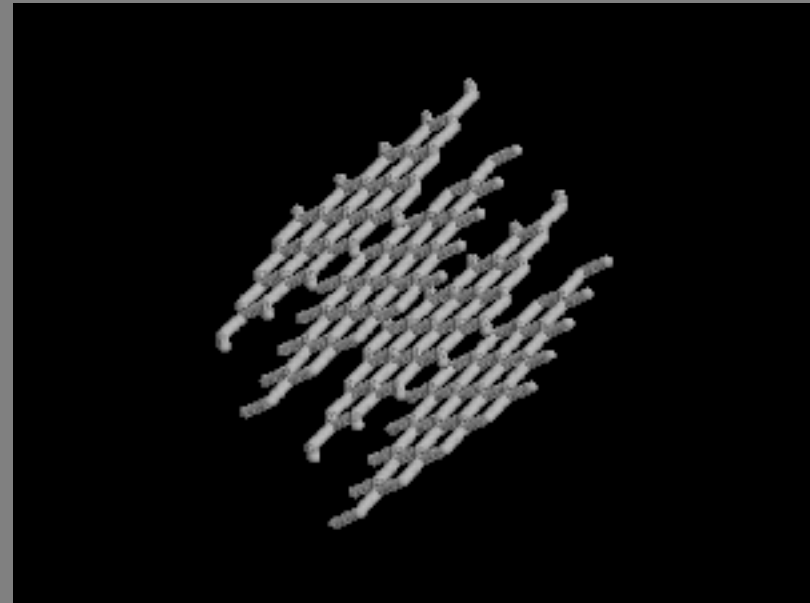
B Carbon steel, an interstitial alloy

Network Atomic Solids

Some covalently bonded substances DO NOT form discrete molecules.



Diamond, a network of covalently bonded carbon atoms

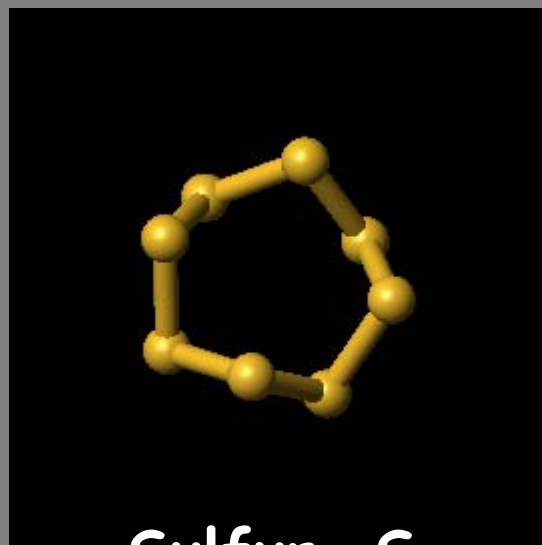


Graphite, a network of covalently bonded carbon atoms

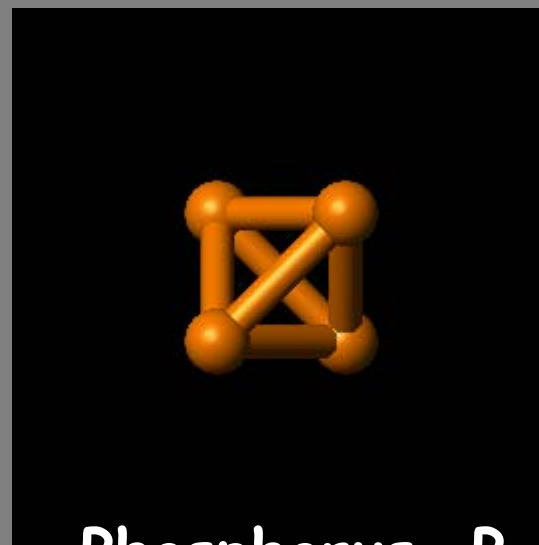
Molecular Solids

Strong covalent forces within molecules

Weak covalent forces between molecules



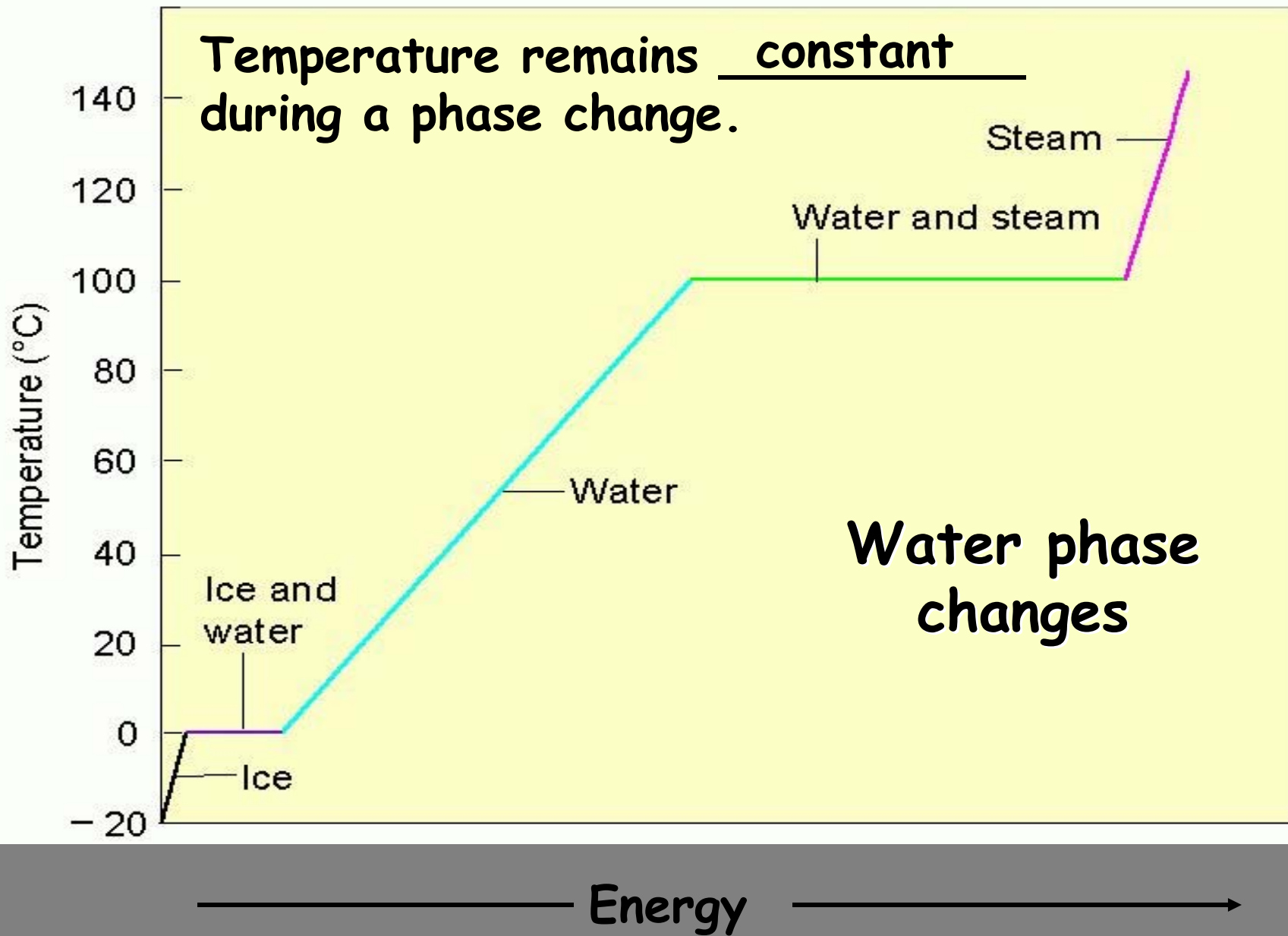
Sulfur, S_8



Phosphorus, P_4

Equilibrium Vapor Pressure

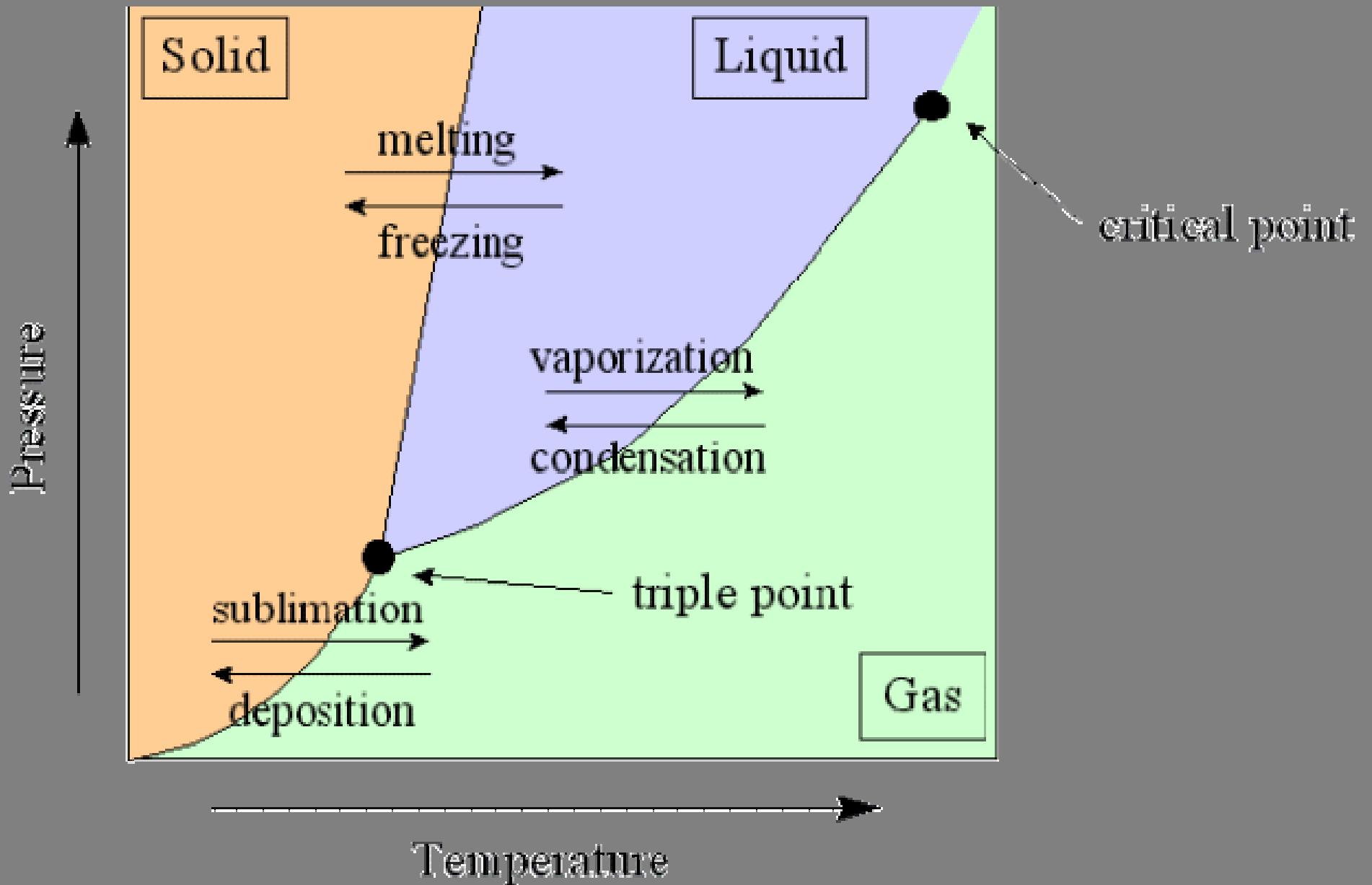
- The pressure of the vapor present **at equilibrium**.
- Determined principally by the size of the intermolecular forces in the liquid.
- Increases significantly with temperature.
- **Volatile liquids** have high vapor pressures.



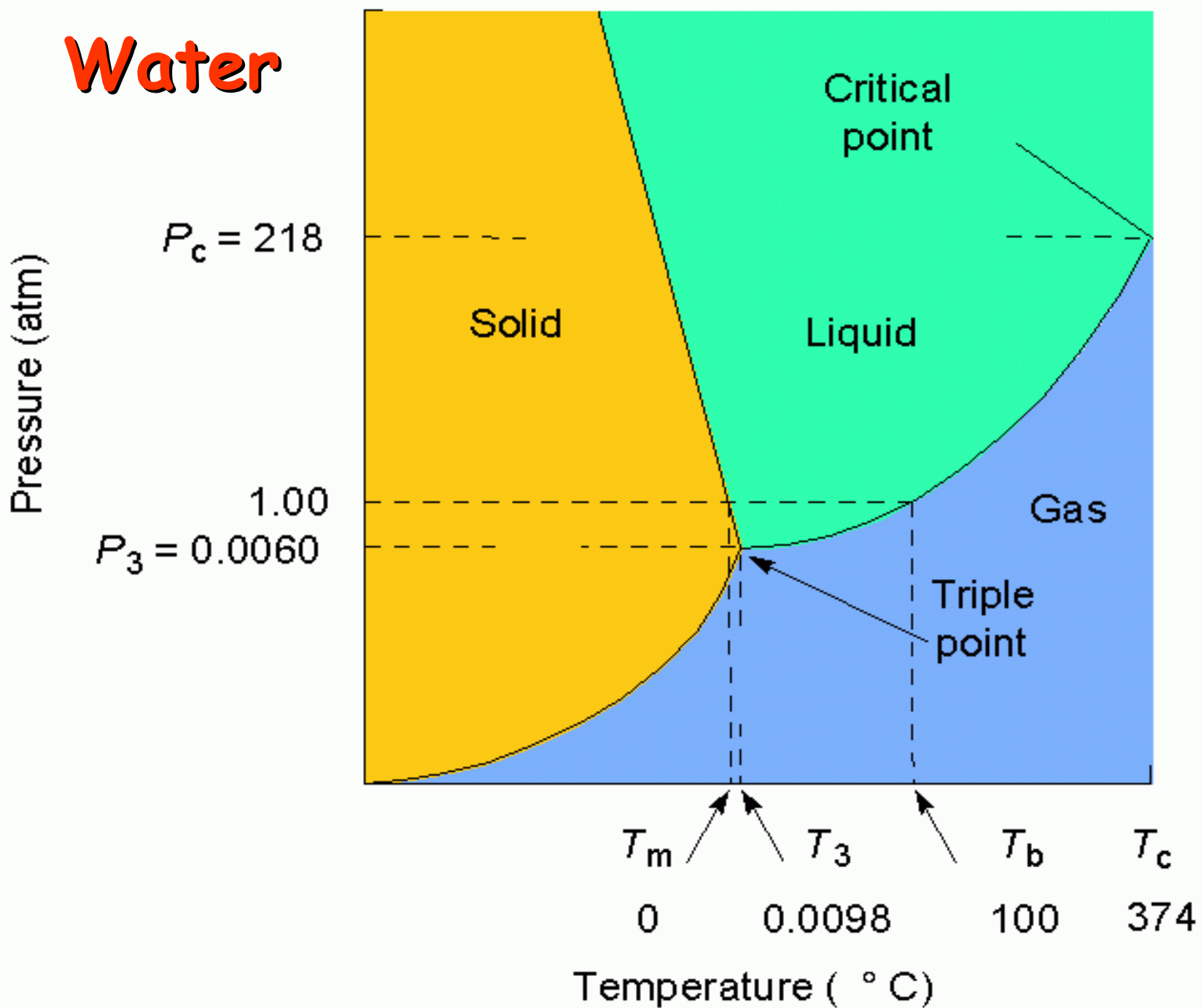
Phase Diagram

- Represents phases as a function of temperature and pressure.
- Critical temperature: temperature above which the vapor can not be liquefied.
- Critical pressure: pressure required to liquefy AT the critical temperature.
- Critical point: critical temperature and pressure (for water, $T_c = 374^\circ\text{C}$ and 218 atm).

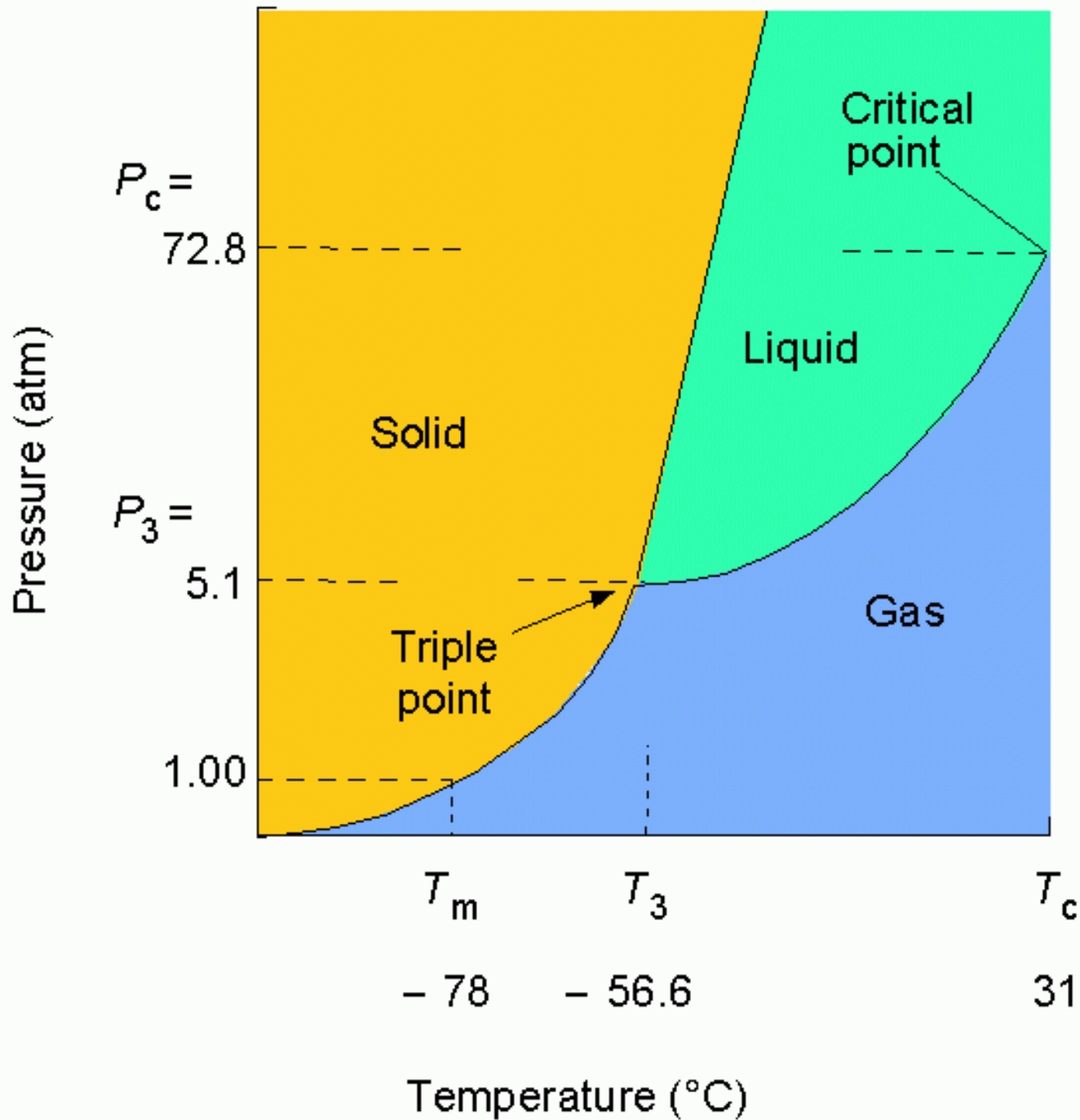
Phase changes by Name

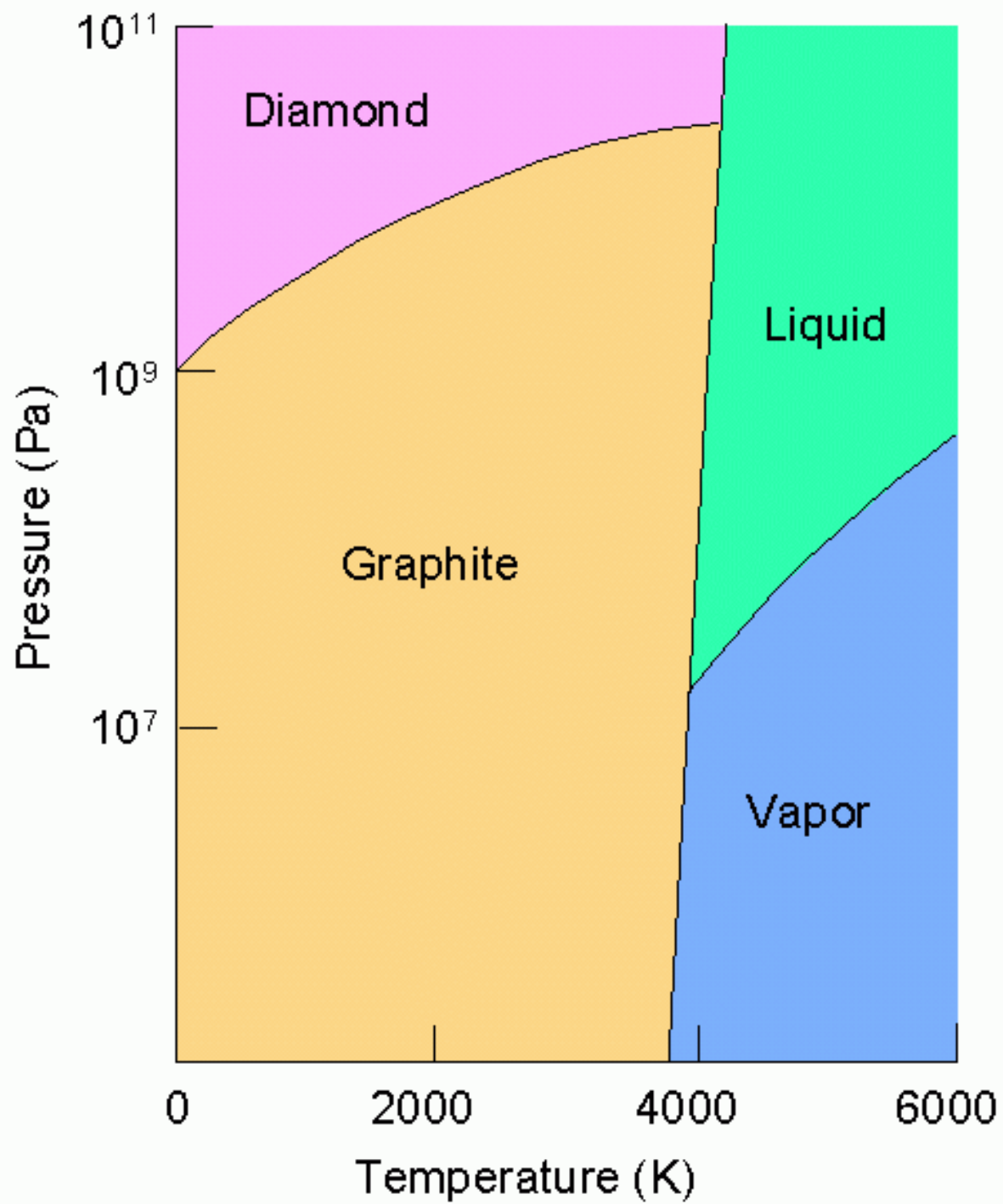


Water



Carbon dioxide





Carbon

Sulfur

