Inspired by Paul Groves

- 1. Int<u>RA</u>molecular = forces holding atoms together...the "bonds"
  - Broken during chemical reactions
- 2. Int<u>ER</u>molecular = forces holding molecules near other molecules
  - Broken during phase changes, pulling molecules away from each other but not pulling the bonded atoms apart! We do not break int<u>RA</u>molecular bonds during a phase change.
- 3. We will be classifying substances into:
  - Covalent (molecular)
    - London dispersion forces
    - o Dipole-Dipole
    - Hydrogen bond
  - Bulk Solids
    - o Metallic
      - "sea of electrons," "free flowing electrons," "mobile carriers" etc
      - Electrons detached and flow freely throughout the material
    - $\circ$  lonic
    - Network covalent
      - Covalent bonded substance that has so many intER and intRa bonds that it is much stronger than expected
      - Common network covalents:
        - Carbon as Diamond and Carbon as Graphite
        - WC, SiC
        - Si, SiO<sub>2</sub>, SiO<sub>4</sub>
        - (any others you come across!)
    - Some new to you IMFs:
      - o Ion-Dipole
        - Interaction between an ion and a polar molecule
      - o Ion Induced Dipole
        - When an ion causes a dipole to form in a non-polar molecule
      - Dipole Induced Dipole
        - When a polar molecule causes a dipole to form in a non-polar molecule

## A BLUFFER'S GUIDE

- **4.** To determine what type of IMF is present
  - Determine if it is ionic, covalent or metallic
    - If covalent determine if it is:
      - Non-polar = London Forces
        - Temporary dipole due to the unequal distribution of electrons.
        - Everything has London Forces!
        - Weak compared to other forces BUT if you have enough cumulative forces it can result in higher properties than expected.
        - Increases with increased
          "polarizability of electron cloud" and surface area.
        - Polar = Dipole-Dipole
          - Permanent polarity due to the structure of the molecule
          - Increases with increased difference in electro negativities between atoms.
          - Bond dipoles do not cancel out, resulting in a "net dipole" for the molecule
        - Polar AND "H-NOF:" = H-bonding
          - Stronger than a typical dipoledipole interaction.
          - Must meet the criteria of a hydrogen bonded to a highly electronegative element (N, O, or F) with a lone pair.
          - Important examples: DNA, protein folding, water
- 5. As the amount of IMFs increases
  - Properties increase
    - Boiling point increases
    - Melting point increases
    - Viscosity increases
    - Surface tension increases
  - Deviates more from "ideal" behavior
- 6. Miscibility is based on IMFs
  - Polar things mix with polar things
  - Nonpolar things mix with nonpolar things

- 7. Vapor Pressure
  - Vapor pressure is based on the dynamic between the liquid and the vapor above the liquid's surface, when measured in a closed container.



- Temperature is the only variable that changes the vapor pressure, not the quantity of liquid, the pressure above the liquid, or the volume of the container. Vapor-Liquid dynamics are an equilibrium situation. Example:
  - If you increase the volume of the container, the pressure inside will decrease.
  - When the pressure decreases, there are fewer vapor molecules per volume so the rate of condensation will slow down.
  - So for a period of time the rate of vaporization will be higher than the rate of condensation.
  - This will increase the amount of vapor in the container.
  - Eventually enough vapor is made so the rate of vaporization and rate of condensation are equal again.
  - Equilibrium is restablished!
  - Vapor pressure is now back to what it was before! Changing the container volume did not affect the vapor pressure value.
- The lower the IMFs, the higher the vapor pressure above the liquid will be. The liquid is more "volatile"
  - The molecules are not being held into the liquid phase as much. More are in the vapor phase
- The higher the IMFs, the lower the vapor pressure above the liquid will be. The liquid is less "volatile"
  - The molecules are being held in the liquid phase! Less are in the vapor phase.



- Boiling occurs when the vapor pressure matches the pressure above the container.
  - You can make things boil by either heating the temperature so the vapor pressure increases, OR by lowering the pressure above the liquid so that it matches the vapor pressure.
  - "Normal Boiling Point" is the temperature at which a liquid will boil when at sea level, 1atm.



- $s \leftrightarrow l = melting$ , freezing
- $I \leftrightarrow g = vaporizing$ , condensing
- During heating/cooling, there is a change to the kinetic energy (temperature is changing)
- During phase changes, there is a change to the potential energy (particles are changing position, closer or further apart, NO temperature change!)
- To calculate energy change involved:
  - During a heating/cooling segment  $\Delta H = mC\Delta T$
  - $\circ \quad \mbox{During a phase change} \\ \Delta H_{fus} = m L_{fus}$ 
    - $\Delta H_{vap} = mL_{vap}$

Properties Indicate Strength of Intermolecular Forces (IMF's)

IMF	BP	FP	$H_{vap}$	$H_{\text{fus}}$	VP
IMF	BP	FP	$\mathbf{H}_{\mathrm{vap}}$	H <sub>fus</sub>	VP

8. Phase Changes