

Chapter 11 HW pp. 494 - 501: Set 1- 8, 10, 12, 16, 19, 20; Set 2- 36, 40, 48, 52, 54, 56;  
Set 3- 74, 75, 78, 82, 85, 93, 96; Set 4- 110, 118, 125, 126, 133, 137, 140.

Intermolecular Attractive Forces and Liquids and Solids

A. \_\_\_\_\_ Attractive Forces

1. Johannes \_\_\_\_\_

a. Tried to explain why something condenses. As the molecules \_\_\_\_\_ down in a gas being \_\_\_\_\_ something must \_\_\_\_\_ these molecules together to change \_\_\_\_\_. \_\_\_\_\_ forces. The higher the temperature at which the gas condenses the stronger the intermolecular forces. There are two basic types

b. \_\_\_\_\_ forces ( \_\_\_\_\_ interaction or \_\_\_\_\_ forces) {Named for Fritz London}

(1) This type exists between \_\_\_\_\_ molecules. The \_\_\_\_\_ the molecule ( number of electrons) the \_\_\_\_\_ the attraction. This is the \_\_\_\_\_ van der Waals force. This can also be viewed as a momentary dipole. Since the electrons are constantly moving about the molecule, it is logical to assume at a particular time most of the electrons will be clustered about \_\_\_\_\_ end of a molecule thus creating a \_\_\_\_\_ distribution of charge or a dipole. This creates attractive forces between two molecules, momentarily.

c. \_\_\_\_\_ - \_\_\_\_\_ attractive force.

(1) This only exists between \_\_\_\_\_ (polar molecules with two distinct regions of opposite charge). This is a \_\_\_\_\_ force than dispersion. The slightly positive end of a molecule is attracted to the slightly negative end of another.

(2) A special type of Dipole - Dipole attraction is a \_\_\_\_\_ bond. This is the \_\_\_\_\_ type of dipole - dipole attraction since the hydrogen only has one valence electron and its proton is easily exposed. This exposed proton attracts other negatively charged regions of other dipoles. This has 5% of the strength of a covalent bond. It is \_\_\_\_\_ a true chemical bond but is important in determining properties.

B. Liquids - \_\_\_\_\_ common phase of matter in universe because liquid phase of any substance can exist only within a \_\_\_\_\_ range of \_\_\_\_\_ and \_\_\_\_\_.

1. General description - \_\_\_\_\_ shape, \_\_\_\_\_ volume

2. KMT description (Review this in chapter 5, page 197)

a. consists of \_\_\_\_\_

b. substance is a liquid because of the \_\_\_\_\_ and \_\_\_\_\_ of particles not the \_\_\_\_\_ of particles

- c. particles are in \_\_\_\_\_ motion
- d. particles are \_\_\_\_\_ together and lower in \_\_\_\_\_ and \_\_\_\_\_ than gas particles thus the attractive forces must be \_\_\_\_\_ than gas and limits motion
- e. kinetic energy is \_\_\_\_\_ enough so the particles are not bound together as in solids.

3. Properties

- a. \_\_\_\_\_ - ability to flow since particle are not bound about a fixed position, the  $E_K$  overcomes attractive forces.
- b. \_\_\_\_\_ - attractive forces are strong enough to hold particles together on one surface
- c. \_\_\_\_\_ - 1000 times greater than corresponding gas; particles are more closely packed due to attractive forces.
- d. \_\_\_\_\_ - particles are closely packed and can evenly distribute pressure in all directions
- e. \_\_\_\_\_ - Definition - when one substance is added to another so that, after mixing, only one phase is observed, the substance has been dissolved. Liquid is the most common dissolving agent. Why ? Motion of particles and attractive forces they exert on solute.
- f. \_\_\_\_\_ - any liquid will gradually diffuse throughout any other liquid in which it can be dissolved. Due to constant random motion of particles. Occurs slower than gases since forward moving particles of closely packed liquid will soon strike another particle, changing its path and attractive forces impede movement. Increases with temperature since avg.  $E_K$  increases thus the particles are moving faster.
- g. \_\_\_\_\_ - common to all liquids to different degrees; attractive forces tend to pull adjacent parts of liquid's surface together, thus making the surface less penetrable by solid bodies; acts to minimize free surface area - as a result, liquid droplets tend to be spherical shape which is the least possible surface area.
- h. \_\_\_\_\_ - tendency for liquid to rise up a narrow tube resulting from the attractive forces between particles and glass
- i. \_\_\_\_\_ - process by which a liquid or solid changes to a gas is vaporization; process by which particles escape from the surface of a non-boiling liquid and enter into gaseous phase is evaporation. Some particles are moving faster than other particles and can escape spontaneously into gaseous phase. This lowers the liquid's  $E_K$  and allows it to absorb energy from the surroundings

thus giving more particles enough energy to escape. This is a continuous process until all liquid is evaporated. No noticeable temperature change. (Good short answer question)

-This also causes vapor pressure in a closed system. Equilibrium is reached when as many particles are going to vapor as are returning to liquid. This is continuous thus dynamic. Rate are equal not particles. When equilibrium is reached the vapor pressure is called the \_\_\_\_\_ . Vapor pressure is directly related to intermolecular attractive forces. Substances with strong IM forces have \_\_\_\_\_ vapor pressures; weak IM forces have \_\_\_\_\_ vapor pressures. Why? (another good short answer)

j. \_\_\_\_\_ - When bubbles appear throughout liquid ; this occurs when the equilibrium vapor pressure equals the atmospheric pressure. The temp at which this occurs is called the boiling point. The temperature at which the vapor pressure of a solid equals the vapor pressure of the corresponding liquid is melting point. Depends on IM forces - weak IM forces (non polar, low molar mass) - low boiling point and melting point ; strong IM forces (polar) - high boiling and melting points. If a solid has a high enough vapor pressure the solid will sublime - turn to gas with out passing through the liquid phase (iodine). How does boiling differ from evaporation? \_\_\_\_\_ . A liquid with a very low boiling point and evaporates readily at room temp is said to be volatile, high vapor pressures.

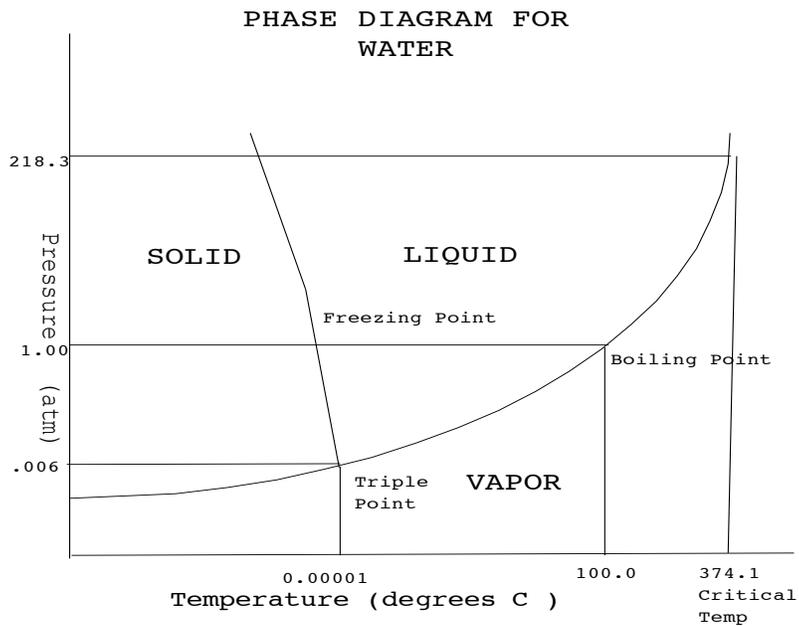
k. \_\_\_\_\_ & \_\_\_\_\_ - Condensation is vapor to liquid while liquefaction is gas to liquid. Cooling gas / vapor slows the particles until the IM forces ca take hold. Compression helps since van der Waals forces are effective for short distances. Associated terms - \_\_\_\_\_ ( $T_c$ ) - temp above which no amount of pressure can liquefy gas. Indication of strength of IM forces. ( Table 11.7 ) critical pressure ( $P_c$ ) - minimum pressure needed to liquefy gas at  $T_c$  .

l. \_\_\_\_\_ - As liquid is cooled  $E_K$  is small enough for IM forces to hold particle about a fixed point with an orderly arrangement (solid). All liquids freeze. Wax freezes at room temp

m. \_\_\_\_\_ - equilibrium - In a closed system we have liquid + energy  $\rightleftharpoons$  vapor and then vapor  $\rightleftharpoons$  liquid + energy once equilibrium is reached ( rates are equal ) we have liquid + energy  $\rightleftharpoons$  vapor

If this equilibrium is disturbed by a stress, the equilibrium will shift to relieve that stress and form a new equilibrium at a new rate – called \_\_\_\_\_.

- n. \_\_\_\_\_ - A diagram used to show the relationships between solids, liquids and gases. It is a graph of temperature vs. pressure that indicates the conditions under which gaseous, liquid or solid phase of a substance exists. The negative slope of solid - liquid line means if pressure drops ice melts at a higher temperature. As temp drops below triple point, the temp of sublimation decreases with decreasing pressure. This is how food is



freeze - dried.. vaporize ice and lower press after freezing.

- o. Most common liquid - \_\_\_\_\_.  
 Why does ice float? \_\_\_\_\_. Why? Water's bond angle is  $105^\circ$ , this creates openings in crystal. Water has strong hydrogen bonds - strong IM forces - high boiling point - low vapor pressure and high critical temp. Water is most dense at  $4^\circ\text{C}$ ; as ice melts particles are closest at  $4^\circ\text{C}$  then move apart and break hydrogen bonds

C. Solids

1. General definition - \_\_\_\_\_ shape, \_\_\_\_\_ volume
2. KMT description
  - a. Consists of particles ( ions, atoms, molecules )
  - b. Avg. kinetic energy is \_\_\_\_\_ than gas and liquid particles thus \_\_\_\_\_ motion; particles move back and forth about fixed position.

- c. More \_\_\_\_\_ packed than gases or liquids thus IM forces are more important ; dipole-dipole, dispersion interaction, hydrogen bonds; hold solid particles in place.
  - d. Much more orderly since held in place and therefore very low \_\_\_\_\_.
3. Particle model of a solid
- a. description in terms of the arrangement and motion of the solid particles and the attractive forces between them.
  - b. two types of solids
    - 1) \_\_\_\_\_ solid - most common - consists of crystal, particles are arranged in an orderly geometric repeating pattern.
    - 2) \_\_\_\_\_ solids - super cooled liquids; from Greek a(\_\_\_\_\_) + Morphe(\_\_\_\_); particles arranged randomly
4. Properties
- a. \_\_\_\_\_ - crystalline solids are geometrically regular; amorphous solids do maintain a definite shape (temporary), but no geometry and can have any shape.
  - b. \_\_\_\_\_ - since particles are very closely packed, strong IM forces; all surfaces are free
  - c. \_\_\_\_\_ - particles are held in fixed position; some amorph do flow very slowly - very viscous
  - d. \_\_\_\_\_ - temp where a solid becomes a liquid;  $E_k$  of particles overcome IM forces; amorphous solids have no definite m.p.( this is why they are called super cooled liquids - retain certain liquid properties at temperatures which appear to be solid
  - e. \_\_\_\_\_ - particles are tightly packed
  - f. \_\_\_\_\_ - particles are as close as they are going to get at normal conditions ( FYI- Discover article - Element Smith pounds elements closer together than normal. Older article)
  - g. \_\_\_\_\_ - particles are still in motion - at surface some can migrate to other solids; rate is a million times slower than in liquids.
5. Closer look at crystalline solids
- a. classified by \_\_\_\_\_ and shape -Done by X-ray crystallography , as X-ray passes through crystal it gives a distinct pattern similar to light passing through a prism as in spectroscope lab.
  - b. \_\_\_\_\_ structure is the total 3-D arrangement of particles
  - c. \_\_\_\_\_ is the smallest portion of a crystal lattice that shows the pattern ( simplest repeating pattern ).
  - e. \_\_\_\_\_ basic shapes

NAME	Axes	Angles between axes
	all equal axes	all $90^\circ$
	2 equal	all $90^\circ$
	all equal	all not $90^\circ$
	all unequal	all $90^\circ$
	all unequal	2 = $90^\circ$ , 1 not
	all unequal	no $90^\circ$
	3 equal	1 = $90^\circ$ , 3 = $60^\circ$

- f. central shapes may contain different type of unit cells for example : simple cubic - each particle has \_\_\_\_\_ immediate neighbors and contains \_\_\_\_\_ atom; face-centered - each particle has \_\_\_\_\_ immediate neighbors and contains \_\_\_\_\_ atoms; body centered - each particle has \_\_\_\_\_ immediate neighbors and contains \_\_\_\_\_ atoms.
- g. Forces that keep crystals together (4 basic types of crystals)
- 1) \_\_\_\_\_ Crystal -contains ions, Group1 or 2 w/Group16 or 17, strong binding (+) (-) attraction results in hard & brittle, high melting points and good insulators.
  - 2) \_\_\_\_\_ (Network) Crystals - contains single atoms covalently bonded to neighbor, very large # of atoms e.g. diamonds ( $C_x$ ), quartz ( $SiO_2$ )<sub>x</sub> , carbide ( $SiC$ )<sub>x</sub> where x = indefinite chains; giant or macromolecules; hard & brittle, high melting points, nonconductors or semiconductors.
  - 3) \_\_\_\_\_ Crystals - positive ions of metal surrounded by a cloud of valence electrons donated by metal atoms and belonging to the crystal as a whole, binding force is between (+) ions and electrons; electrons are free to migrate through crystal lattice thus high electrical conductivity; m.p. vary
  - 4) \_\_\_\_\_ Crystals - covalently bonded molecules held together by IM forces between molecules; non polar molec.= dispersion interaction; polar molec. = dipole-dipole and dispersion; low melting points, easily vaporized, soft and good insulators
6. More about Morphs
- a. amorphous forms of \_\_\_\_\_ alloys prepared by rapid cooling of thin films of metals- atoms do not have time to arrange themselves - resulting in “\_\_\_\_\_”, strong flexible more resistant to corrosion than crystals of same composition.

- b. \_\_\_\_\_ - crystals of different solids with same structural shape
- c. \_\_\_\_\_ - crystals of same substance with 2 or more shapes

7. Nothing's Perfect

- a. \_\_\_\_\_ or \_\_\_\_\_ either within unit cell structure (ions missing others in place) or extra layer.
- b. Some defects are on purpose and helpful
  - 1) man made defects - crystals are "\_\_\_\_\_"; used to make \_\_\_\_\_. Si & Ge can be made to conduct electricity, if As is added - extra electron available to move across crystal. If Ga is added - short an electron and electrons fill "hole."
- c. Other impurities
  - 1) If water is in crystal that is if ions are chemically bonded to water (\_\_\_\_\_ ions) & then crystallize - \_\_\_\_\_ crystal or \_\_\_\_\_. To remove water - heat, lower pressure, then they become \_\_\_\_\_.  $\text{CuSO}_4 \cdot 5\text{H}_2\text{O}$
  - 2) some substances absorb water from air - \_\_\_\_\_
  - 3) some absorb enough to dissolve - \_\_\_\_\_

8. What the heck is a \_\_\_\_\_ crystal?

- a. crystalline solid very orderly, liquid very unorderedly - random in all three dimensions; solid \_\_\_\_\_ crystalline order in only one or two \_\_\_\_\_ at melting point form liquid crystals. (smectic - retain 2 orders; nematic - retains 1 order)

D. Energy involved in Phase Changes

1. Substances boil and melt at constant temps. This is due to energy needed for phase change to occur. IM forces are overcome.
2. Energy required to melt one gram of a substance at its melting point is called the \_\_\_\_\_ ( $\Delta H_f$ ) ( $\text{H}_2\text{O} = 334 \text{ J/g}$ )
3. Energy required to vaporize 1 g of a substance at its boiling point is called the \_\_\_\_\_ of \_\_\_\_\_ ( $\Delta H_v$ ) ( $\text{H}_2\text{O} = 2260 \text{ J/g}$ )
4. Also involved is the \_\_\_\_\_ heat (amount of heat needed to raise 1 g of a substance  $1 \text{ C}^0$ .) since substance needs to reach its m.p. or b.p. ( $C_p$ )

$$\text{H}_2\text{O}_{(l)} = \frac{\quad \text{J/g C}^0}{\quad \text{H}_2\text{O}_{(v)}} \quad \text{H}_2\text{O}_{(s)} = \frac{\quad \text{J/g C}^0}{\quad \text{H}_2\text{O}_{(v)}}$$

5. Formula  $q = mC_p(\Delta T)$  not needed just catch the fever!  
 $q = \text{Heat (J)}$      $m = \text{mass (g)}$      $\Delta T = \text{change in temp (C}^0\text{)}$   
 $C_p = \text{specific heat}$