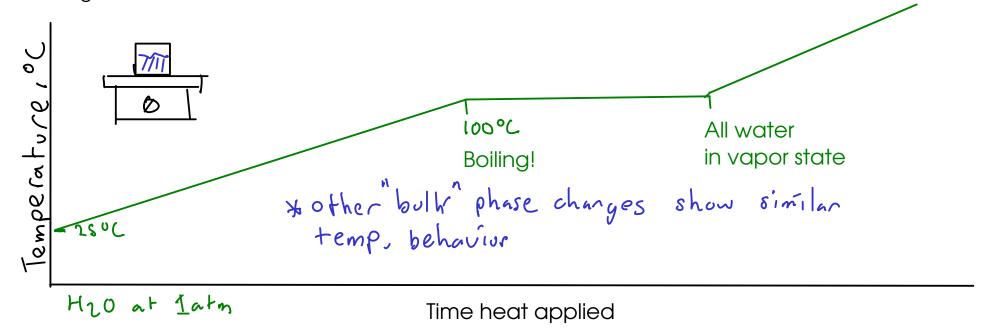


- For a molecule to move from the liquid phase to the gas phase, it must acquire enough KINETIC ENERGY (which depends on molecular SPEED) to break away from the INTERMOLECULAR FORCES holding the molecule in the liquid.
- The AVERAGE KINETIC ENERGY of molecules is proportional to the TEMPERATURE. On average, molecules in both the liquid and the solid state move faster at higher temperatures.
- Even at room temperature, some of the molecules have enough kinetic energy to escape into the gas phase. This accounts for the tendency of volatile liquids to evaporate from open containers even well away from the boiling point.
- As we increase temperature, the gaussian curve above shifts to the right, and a higher fraction of molecules have enough energy to vaporize.

- To increase the rate of vaporization, we can increase the temperature. At the BOILING POINT, the pressure of the vaporizing water is enough to push back against the liquid water and bubbles of vapor form in the liquid: BOILING
 - If we look at the temperature changes up to and through the boiling point, we see something interesting...



DURING THE BOILING PROCESS, as long as you have some liquid water remaining, the temperature will remain constant - EVEN AS YOU CONTINUE TO APPLY HEAT!

How do we explain this behavior?

- The VAPORIZATION itself requires an energy input. What's that energy doing? Breaking water molecules away from one another (breaking apart the water's intermolecular forces).

LIQUIDS

- FLUID, DENSE, INCOMPRESSIBLE
- Posess a few unique properties
 - U SURFACE TENSION
 - a measure of the tendency of a liquid to minimize its surface area, or the resistance to the breaking of a liquid surface.

Liquid droplets tend to be spehrical...

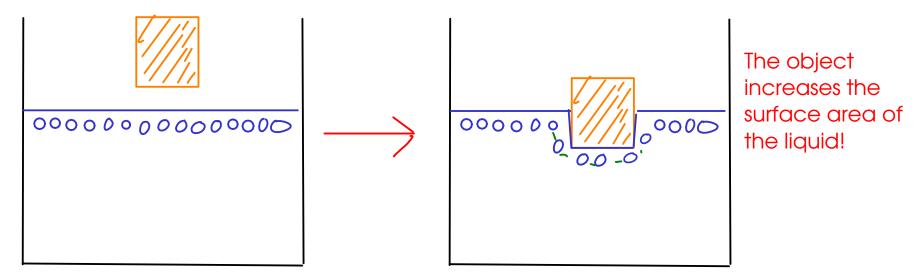
... but friction causes falling droplets to take on the traditional "drop" shape.

A sphere has the smallest surface area for a given volume.

- (2) VISCOSITY (also for gases)
 - a measure of a liquids resistance to flow, or "thickness"

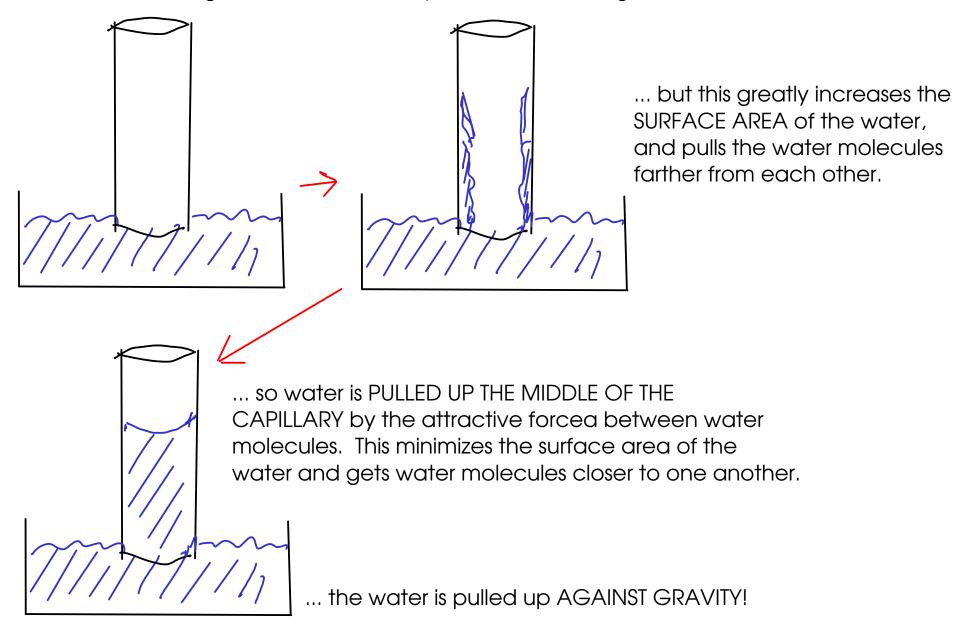
SURFACE TENSION

- Surface tension can be explained by looking at liquid molecules as being attracted to each other by INTERMOLECULAR FORCES.

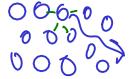


For the object to penetrate the liquid surface, it must push water molecules at the surface apart. Since these water molecules ARE ATTRACTED TO ONE ANOTHER, the liquid will resist!

³⁷Surface tension also explains CAPILLARY ACTION, the drawing up of WATER into a glass tube. Water is attracted to glass, and will climb up the surface of a glass tube.



- viscosity can also be explained (at least partially) by looking at INTERMOLECULAR FORCES!
- For a liquid to FLOW, its molecules must move past one another. This means that some of the molecules must move farther away from other molecules. Since the molecules in the liquid state are ATTRACTED TO ONE ANOTHER, that means the flowing will be slowed.
- Viscosity is also determined by STRUCTURE. Liquids with large chains (like oils) which can rotate and tangle in one another will also be viscous.



Molecules have to move past one another to flow, and stronger attractions between molecules make that more difficult!

INTERMOLECULAR FORCES IN LIQUIDS

- "Intermolecular forces" is a generic term. It refers to any number of forces that exist between molecules!
- In liquids, there are three main types of intermolecular force
 - 1 DIPOLE-DIPOLE INTERACTIONS

- only for polar molecules

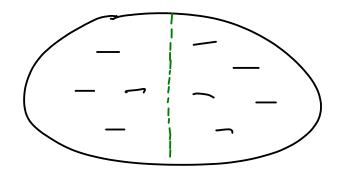
van der Waals forces...

- 2 LONDON DISPERSION FORCES
 - exist in all liquids
- 3 HYDROGEN BONDS
 - exist only when hydrogen is directly bonded to a highly electronegative atom.

- attractions between the positive pole of one polar molecule and the negative pole of another

- Dipole-dipole interactions occur only between POLAR molecules
- Dipole-dipole interactions are weak relative to the other two kinds of intermolecular forces in liquids
- The more polar a molecule is (the larger the dipole moment), the stronger its dipole-dipole interactions.

- often called "London forces" for short.
- occurs because electron density is at any given point in time likely to be uneven across a molecule due to the simple fact that electrons are MOVING!



Let's say that at one point in time, the electrons around a molecule are here. At this moment, there is a TEMPORARY (INSTANTANEOUS) DIPOLE.

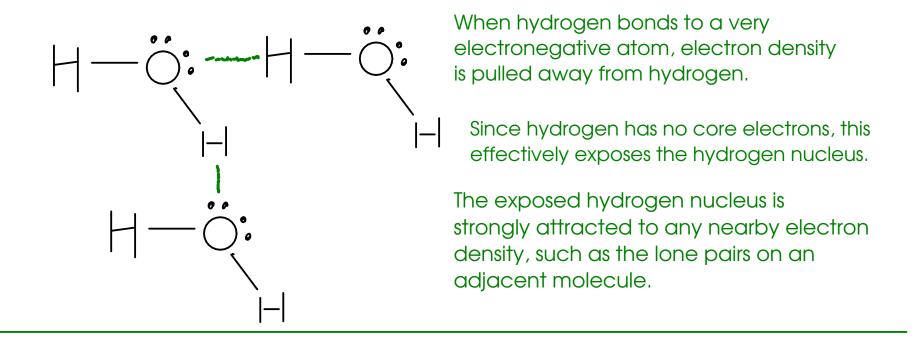
This dipole can interact (and even induce) dipoles on other, nearby molecules.

What happens when the electrons move around some more? The dipole may flip, and the surrounding INDUCED DIPOLES flip along with it. This results in a net ATTRACTIVE FORCE between molecules.

- London forces occur in all molecules, polar or nonpolar.
- London forces increase in strength as molecules get larger. The larger a molecule is, the more easily instantaneous dipoles form and the more easily dipoles can be induced.
- London forces are often the most important interaction between molecules in the liquid state, unless the molecules are capable of HYDROGEN BONDING.

HYDROGEN BONDING

- can happen when there is a HYDROGEN ATOM bonded DIRECTLY to either O, N, or F and that O, N, or F atom has at least one lone pair.



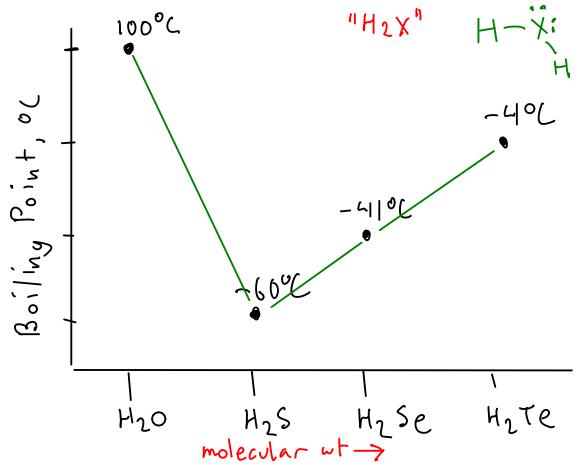
- Hydrogen bonds are the strongest type of intermolecular force in the liquid state. They are ALMOST as strong as a covalent bond.
- Hydrogen bonds form in several important molecules: water, ethanol, nucleic acids (they hold the DNA helix together)

FORCES AND OBSERVABLE PROPERTIES

- These intermolecular forces affect observable liquid properties!
 - (I) MELTING AND BOILING POINTS
 - Stronger forces mean higher melting and boiling points.
 - 2 SURFACE TENSION
 - Stronger forces mean greater surface tension.
 - (3) VISCOSITY
 - Stronger forces mean a more viscous (thick) substance.
 - ... but viscosity is also affected by molecular structure!
 - 4 VAPOR PRESSURE
 - ... is the PARTIAL PRESSURE of vapor over a liquid's surface. The liquid boils when its vapor pressure equals atmospheric pressure.
 - Stronger forces mean a LOWER vapor pressure.

COMPARING MOLECULES

- In general, heavier molecules have stronger intermolecular forces than lighter molecules. (London forces depend on size!)
- For molecules OF SIMILAR SIZE, a polar molecule will have stronger intermolecular forces than a nonpolar one.
- Molecules that hydrogen bond will have much stronger intermolecular forces than you would otherwise expect.



These molecules are similar in structure.

The boiling points of the sulfide, selenide, and telluride increase with molecular weight, as we expect.

Water is capable of HYDROGEN BONDS, so its boiling point is much greater than the others in the series, even though it has a much smaller molecular weight!

- RIGID, DENSE, and INCOMPRESSIBLE
 - Properties of interest:
 - (1) MELTING POINT
 - Temperature at which the bulk phase change from solid to liquid occurs
 - (2) HARDNESS and BRITTLENESS
 - hardness: resistance of a solid to deformation (shape change) caused by the application of a force
 - brittleness: tendency of a material to fracture or break rather than to deform.
 - (3) CONDUCTIVITY
 - ability of a material to conduct an electric current

... these properties will be influenced by the KINDS OF FORCES holding the solid together!

- Solids may be classified either by the type of forces holding the solid together or by structure. We'll discuss forces first.
- Some solids are held together by the same sorts of forces found in liquids. But there are more options for solids!
- There are four kinds of solids when classified by forces.
- MOLECULAR SOLIDS
 - held together by the same kinds of forces that hold liquids together:
 - van der Waals forces: London dispersion forces and dipole-dipole interactions
 - (B) hydrogen bonds

... generally, these forces are the weakest.

Examples: candle wax, water ice

Generally, molecular solids:

- have LOW MELTING POINTS
- are SOFT
- are NONCONDUCTORS

- held together by METALLIC BONDS, which involve electron sharing throughout the body of the metal..
- ... strength of these metallic bonds is variable.

Examples: iron, gold, copper, zinc, other metals

Generally, metallic solids:

- have a wide range of MELTING POINTS, though almost all melt above room temperature.
- range from SOFT to HARD. Many are MALLEABLE, meaning they deform before breaking.
- are good CONDUCTORS of both heat and electricity

3 IONIC SOLIDS

- held together by IONIC BONDS:
- ... generally, these forces are much stronger than the ones in molecular solids.

Examples: sodium chloride, any ionic compound

Generally, ionic solids:

- have HIGH MELTING POINTS, well over room temperature
- are HARD
- are NONCONDUCTORS of electricity in the solid phase, but CONDUCT when melted or dissolved into a liquid solution.

(4) COVALENT NETWORK SOLIDS

- held together by COVALENT BONDS.
- are, in essence, giant molecules where the entire solid (not simply individual molecules WITHIN the solid) are held together by covalent bonds.

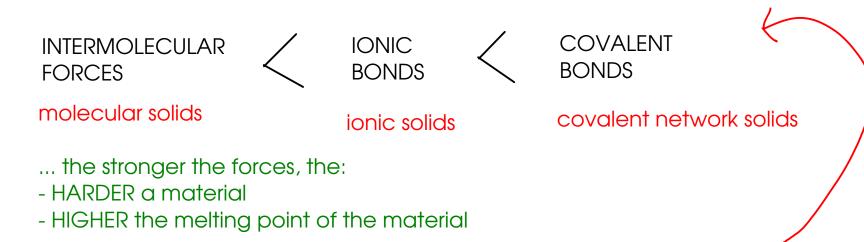
... these are the strongest kind of forces holding solids together.

Example: diamond

Generally, covalent network solids:

- have EXTREMELY HIGH MELTING POINTS. Most don't melt at all they decompose before melting.
- are EXTREMELY HARD. The hardest materials known are covalent network solids.
- are NONCONDUCTORS

Relative strengths of the forces holding solids together:



Metallic bonds vary considerably, so they have been left out of the comparison!

- Solids may also be classified by structure. A more in-depth look at solids is something you would find in a materials science class, but we'll discuss two broad categories of solid materials.

(1) AMORPHOUS SOLIDS

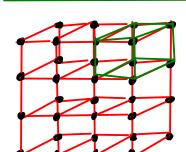
- have a disordered structure at the microscopic level.
- a very small amount of solids are completely amorphous, but quite a few plastics are at least partially amorphous.

2 CRYSTALLINE SOLIDS

- have a well-defined three dimensional structure at the microscopic level.
- structure is made up of a regular, repeating arrangement of points in space a CRYSTAL LATTICE



• • • The simplest repeating pattern that describes the entire crystal is called the UNIT CELL. It's outlined in GREEN here.



Here's a crystal lattice in three dimensions. This one is called a SIMPLE CUBIC lattice. This simple structure can be found in some solid metals like polonium. A polonium atom occupies each lattice point.

The unit cell, again, is highlighted in GREEN.

See pages 449-450 for more types of crystal systems and more unit cells.

- Natural crystals almost always have some DEFECTS in their structure.
 - Holes in the crystal lattice, where an atom should be but isn't
 - Misaligned planes in the crystal
 - Substitutions of one atom for another in the crystal lattice
- Often defects are undesirable, but not always:

Alumina: Al203

- clear / white in color
- usually used as the "grit"in cleaners like Comet andSoft Scrub!

ruby: A1203 with some A1
replaced with Cr

- red in color
- valuable gemstone!