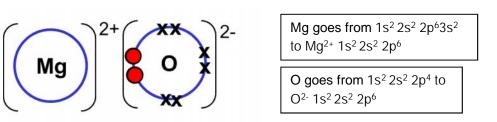
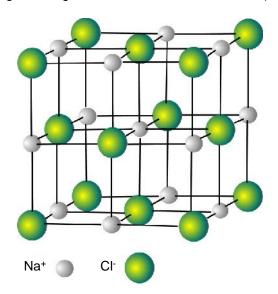
2.2.2 Bonding and Structure

Ionic Bonding

Definition: lonic bonding is the **electrostatic force** of attraction between **oppositely charged ions** formed by electron transfer.



lonic bonding is stronger and the melting points higher when **the ions are smaller** and/ or have **higher charges.** E.g. MgO has a higher melting point than NaCl as the ions involved ($Mg^{2+} \& O^{2-}$ are smaller and have higher charges than those in NaCl, Na⁺ & Cl⁻)



The ions in an ionic solid are arranged in a regular 3D pattern called a giant ionic lattice

Metal atoms lose electrons to form

Non-metal atoms gain electrons to

+ve ions.

form -ve ions.

The sticks in this diagram are there to help show the arrangements of the ions. They do not represent the ionic bonds.

Ionic bonding is between ions and all their surrounding oppositely charged ions. Each sodium ion in this structure is surrounded and equally attracted by six chloride ions. The ionic bond is the attraction between all these ions

Typical Physical properties of Ionic Compounds

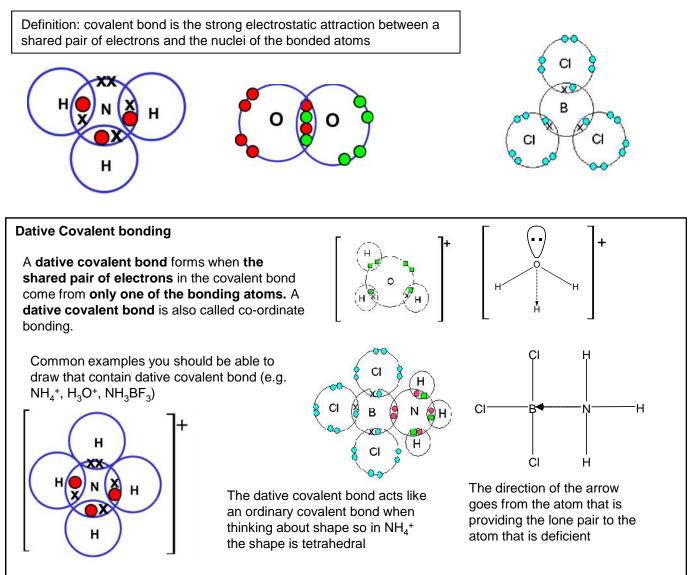
•High melting points - There are strong electrostatic attractive forces between the oppositely charged ions in the lattice

•Non conductor of electricity when solid- The ions are held together tightly in the lattice and can not move so no charge is conducted

•Good conductor of electricity when in solution or molten – The ions are free to move when in solution and molten. Charge can be carried

• They are **usually** soluble in aqueous solvents.

Covalent Bonding



The term *average bond enthalpy is used* as a measurement of covalent bond strength. The larger the value of the average bond enthalpy, the stronger the covalent bond.

Bonding and Structure

| Bonding | Structure | Examples | |
|---|--|--|--|
| Ionic : electrostatic force of attraction between oppositely charged ions | Giant ionic lattice | Sodium chloride Magnesium oxide | |
| Covalent : shared <u>pair</u> of electrons | Simple molecular: With intermolecular forces (Induced dipole–dipole, permanent dipole-dipole, hydrogen bonds) between molecules | Iodine Ice Carbon dioxide Water Methane | |

Only use the words molecules and intermolecular forces when talking about simple molecular substances

| Property | Giant Ionic | Molecular (simple) | |
|----------------------------|---|---|--|
| boiling and melting points | high- because of giant lattice of ions with strong electrostatic forces between oppositely charged ions. | low- because of weak intermolecular forces between molecules (specify type e.g Induced dipole–dipole/hydrogen bond) | |
| Solubility in water | generally good | generally poor | |
| conductivity when solid | poor: ions can't move/ fixed in lattice | poor: no ions to conduct and electrons are localised (fixed in place) | |
| conductivity when molten | good: ions can move | poor: no ions | |
| general description | crystalline solids | mostly gases and liquids | |

Shape of molecules

| Name | No bonding pairs | No lone pairs | Diagram | Bond angle | Examples |
|-----------------------|------------------------|------------------|--|------------|---|
| linear | 2 | 0 | CIBeCI | 180 | CO _{2,} CS ₂ , HCN, BeF ₂ |
| Trigonal planar | 3 | 0 | Cl B Cl | 120 | BF ₃ , AlCl ₃ , SO ₃ , NO ₃ ⁻ , CO ₃ ²⁻ |
| Tetrahedral | 4 | 0 | H (1)(1)(1)(1)(1)(1)(1)(1)(1)(1)(1)(1)(1)(| 109.5 | SiCl ₄ , SO ₄ ²⁻ , ClO ₄ ⁻ , NH ₄ ⁺ |
| Trigonal pyramidal | 3 | 1 | H ^{NWWWWWWW} H | 107 | NCI ₃ ,PF ₃ ,CIO ₃ ,H ₃ O ⁺ |
| Bent | 2 | 2 | Н | 104.5 | OCl ₂ , H ₂ S, OF ₂ , SCl ₂ |
| Octahedral | 6 | 0 | F F F F | 90 | SF ₆ |

How to explain shape

- 1. State number of bonding pairs and lone pairs of electrons.
- 2. State that electron pairs repel and try to get as far apart as possible (or to a position of minimum repulsion.)
- 3. If there are no lone pairs state that the electron pairs repel equally
- 4. If there are lone pairs of electrons, then state that lone pairs repel more than bonding pairs.
- 5. State actual shape and bond angle.

Remember lone pairs repel more than bonding pairs and so reduce bond angles (by about 2.5° per lone pair in above examples)

Definition

Electronegativity is the relative tendency of an atom in a covalent bond in a molecule to attract electrons in a covalent bond to itself.

Factors affecting electronegativity

Electronegativity increases across a period as the number of protons increases and the atomic radius decreases because the **electrons in the same shell** are pulled in more.

It decreases down a group because the distance between the nucleus and the outer electrons increases and the shielding of inner shell electrons increases

A compound containing elements of similar electronegativity and hence a small electronegativity difference will be purely covalent

Formation of a permanent dipole – (polar covalent) bond

A polar covalent bond forms when the elements in the bond have different electronegativities.

When a bond is a polar covalent bond it has an unequal distribution of electrons in the bond and produces a charge separation, (dipole) ends.

Symmetric molecules

A symmetric molecule (all bonds identical and no lone pairs) will not be polar even if individual bonds within the molecular are polar.

The individual dipoles on the bonds 'cancel out' due to the symmetrical shape of the molecule. There is no **net** dipole moment: the **molecule** is non-polar.

e.g. CCl₄ will be non-polar whereas CH₃Cl will be polar

electronegativity difference will be ionic

0 = C = 0

CO₂ is a symmetrical molecule and is a non-polar molecule

A compound containing elements of very different electronegativity and hence a very large

F, O, N and Cl are the most electronegative atoms

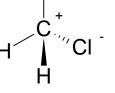
> The element with the larger electronegativity

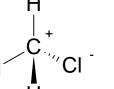
in a polar compound

will be the - end

+

H – Cl





Intermolecular bonding

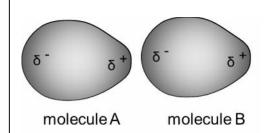
Induced dipole-dipole interactions

Induced dipole–dipole interactions occur between all molecular substances and noble gases. They do not occur in ionic substances.

Induced dipole–dipole interactions are also called **London forces**. They occur between all simple covalent molecules and the separate atoms in noble gases.

In any molecule the electrons are moving constantly and randomly. As this happens the electron density can fluctuate and parts of the molecule become more or less negative i.e. small temporary or transient dipoles form.

These temporary dipoles can cause dipoles to form in neighbouring molecules. These are called induced dipoles. The induced dipole is always the opposite sign to the original one.



Main factor affecting size of Induced dipole-dipole interactions

The **more electrons** there are in the molecule the higher the chance that temporary dipoles will form. This makes the **induced dipole-dipole interactions stronger between the molecules** and so boiling points will be greater.

The increasing boiling points of the halogens down the group 7 series can be explained by the increasing number of electrons in the bigger molecules causing an increase in the size of the induced dipole–dipole interactions between the molecules. This is why I_2 is a solid whereas CI_2 is a gas.

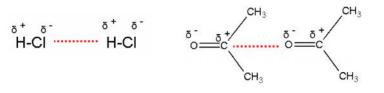
The increasing boiling points of the alkane homologous series can be explained by the increasing number of electrons in the bigger molecules causing an increase in the size of the induced dipole–dipole interactions between molecules.

The shape of the molecule can also have an effect on the size of the induced dipole–dipole interactions forces. Long chain alkanes have a larger surface area of contact between molecules for induced dipole–dipole interactions to form than compared to spherical shaped branched alkanes and so have induced dipole–dipole interactions.

Permanent dipole-dipole forces

•Permanent dipole-dipole forces occurs between polar molecules

It is stronger than induced dipole–dipole interactions and so the compounds have higher boiling points
Polar molecules have a permanent dipole. (commonly compounds with C-CI, C-F, C-Br H-CI, C=O bonds)
Polar molecules are asymmetrical and have a bond where there is a significant difference in electronegativity between the atoms.

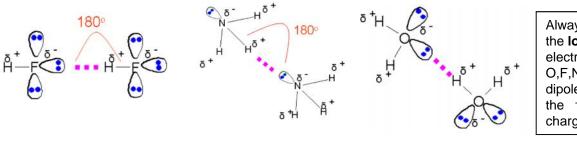


Permanent dipole forces occur in addition to induced dipole–dipole interactions

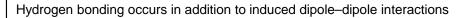
Permanent dipole–dipole and induced dipole–dipole interactions can **both** be referred to as van der Waals' forces.

Hydrogen bonding

It occurs in compounds that have a **hydrogen atom attached to** one of the three **most electronegative** atoms of **nitrogen**, **oxygen and fluorine**, which must have an available lone pair of electrons. e.g. a –O-H -N-H F- H bond. There is a **large electronegativity difference** between the H and the O,N,F



Always show the **lone pair** of electrons on the O,F,N and the dipoles and all the ⁻ ⁺ charges

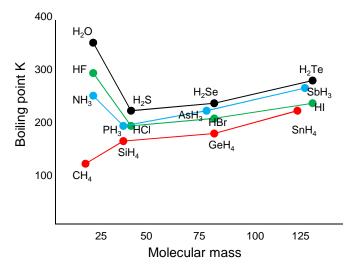


Hydrogen bonding is stronger than the other two types of intermolecular bonding.

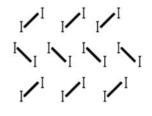
The **anomalously high** boiling points of H_2O , NH_3 and HF are caused by the hydrogen bonding between the molecules.

The general increase in boiling point from H_2S to H_2Te is caused by increasing induced dipoledipole interactions between molecules due to an increasing number of electrons.

Alcohols, carboxylic acids, proteins, amides all can form hydrogen bonds.



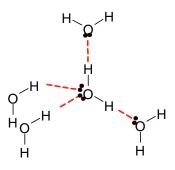
Molecular: lodine



There are covalent bonds between the iodine atoms in the I_2 molecule.

The crystals contain a regular arrangement of I_2 molecules held together by weak induced dipole– dipole interactions intermolecular forces.

Molecular: Ice



Water can form two hydrogen bonds per molecule, because oxygen is very electronegative, and it has two lone pairs of electrons.

The molecules are held further apart than in liquid water and this explains the lower density of ice.