AN INTRODUCTION TO BONDING

A guide for A level students





KNOCKHARDY PUBLISHING

BONDING

INTRODUCTION

This *Powerpoint* show is one of several produced to help students understand selected topics at AS and A2 level Chemistry. It is based on the requirements of the AQA and OCR specifications but is suitable for other examination boards.

Individual students may use the material at home for revision purposes or it may be used for classroom teaching if an interactive white board is available.

Accompanying notes on this, and the full range of AS and A2 topics, are available from the KNOCKHARDY SCIENCE WEBSITE at...

www.knockhardy.org.uk/sci.htm

Navigation is achieved by...

- either clicking on the grey arrows at the foot of each page
 - or using the left and right arrow keys on the keyboard

BONDING

CONTENTS

- Introduction
- Chemical and physical bonding
- Ionic bonding
- Covalent bonding
- Simple molecules
- Van der Waals' forces
- Electronegativity & dipole-dipole interaction
- Hydrogen bonding
- Co-ordinate (dative covalent) bonding
- Molecular solids
- Covalent networks
- Metallic bonding

STRUCTURE AND BONDING

The physical properties of a substance depend on its structure and type of bonding present. Bonding determines the type of structure.

Basic theory

- the noble gases (He, Ne, Ar, Kr, Xe and Rn) are in Group VIII
- they are all relatively, or totally, inert
- their electronic structure appears to confer stability
- they have just filled their 'outer shell' of electrons
- atoms without the electronic structure of a noble gas try to get one
- various ways are available
- the method depends on an element's position in the periodic table

STRUCTURE AND BONDING

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TYPES OF BOND

CHEMICAL ionic (or electrovalent) strong bonds covalent dative covalent (or co-ordinate) metallic

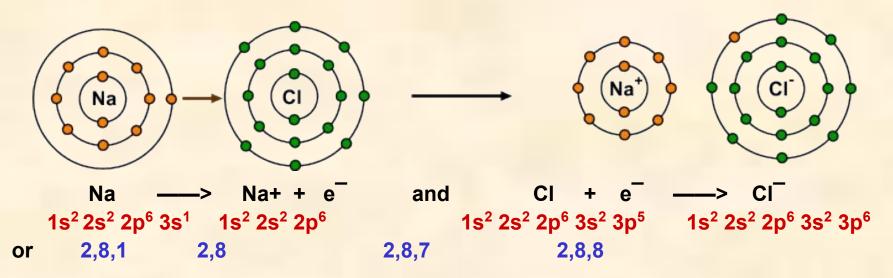
PHYSICAL van der Waals' forces - weakest weak bonds dipole-dipole interaction hydrogen bonds - strongest

IONIC BONDING

lonic bonds tend to be formed between elements whose atoms need to "lose" electrons to gain the nearest noble gas electronic configuration (n.g.e.c.) and those which need to gain electrons. The electrons are transferred from one atom to the other.

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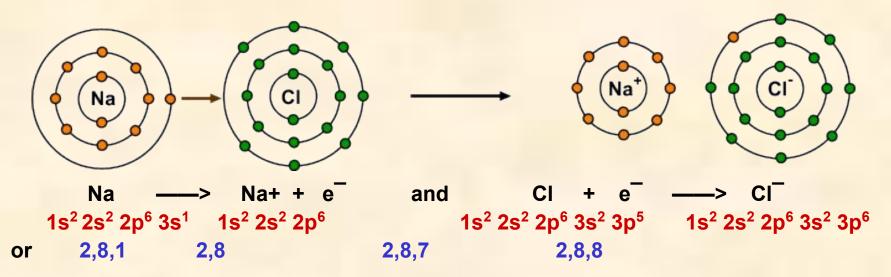
Sodium Chloride





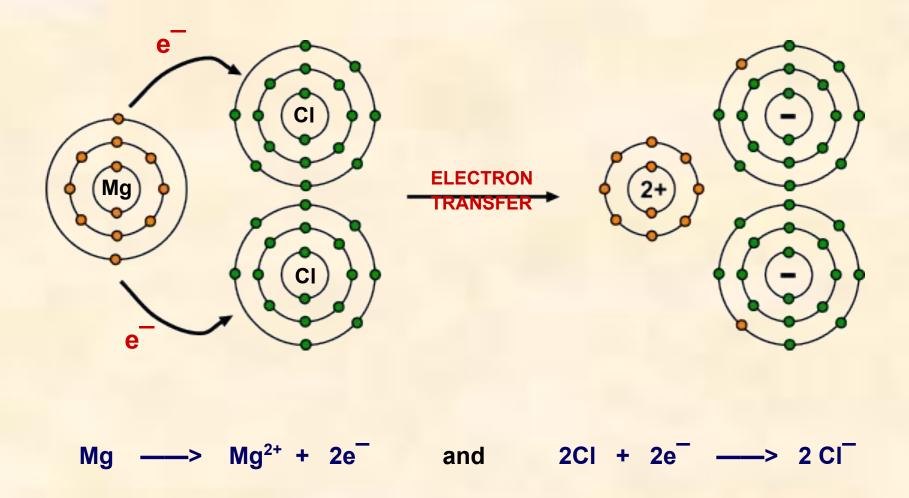
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Sodium Chloride



An electron is transferred from the 3s orbital of sodium to the 3p orbital of chlorine; both species end up with the electronic configuration of the nearest noble gas the resulting ions are held together in a crystal lattice by electrostatic attraction.

FORMATION OF MAGNESIUM CHLORIDE



THE FORMATION OF IONS

Positive ions

- also known as cations; they are smaller than the original atom.
- formed when electrons are removed from atoms.
- the energy associated with the process is known as the ionisation energy

1st IONISATION ENERGY (1st I.E.)

The energy required to remove one mole of electrons (to infinity) from the one mole of gaseous atoms to form one mole of gaseous positive ions.

Other points

Successive IE's get larger as the proton:electron ratio increases.

Large jumps in value occur when electrons are removed from shells nearer the nucleus because there is less shielding and more energy is required to overcome the attraction. If the I.E. values are very high, covalent bonding will be favoured (e.g. beryllium).



THE FORMATION OF IONS

Negative ions

- known as anions
- are larger than the original atom due to electron repulsion in outer shell
- formed when electrons are added to atoms
- energy is released as the nucleus pulls in an electron
- this energy is the electron affinity.

ELECTRON AFFINITY

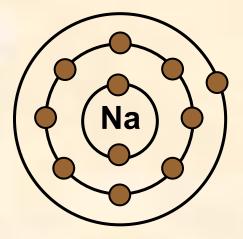
The energy change when one mole of gaseous atoms acquires one mole of electrons (from infinity) to form one mole of gaseous negative ion

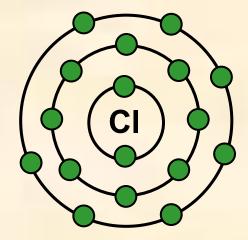
e.g.
$$CI(g) + e^{-} \longrightarrow CI(g)$$
 and $O(g) + e^{-} \longrightarrow O(g)$

The greater the effective nuclear charge (E.N.C.) the easier an electron is pulled in.

IONIC BONDING Animations

SODIUM CHLORIDE

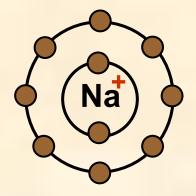


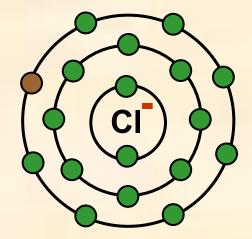


SODIUM ATOM 2,8,1

CHLORINE ATOM 2,8,7

SODIUM CHLORIDE



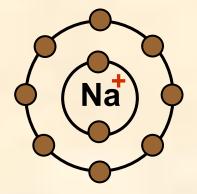


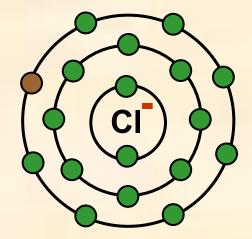
SODIUM ION 2,8

CHLORIDE ION 2,8,8

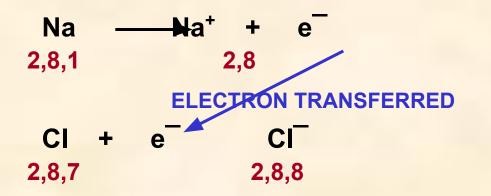
both species now have 'full' outer shells; ie they have the electronic configuration of a noble gas

SODIUM CHLORIDE

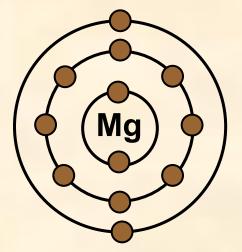




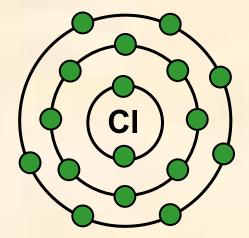
SODIUM ION 2,8 CHLORIDE ION 2,8,8



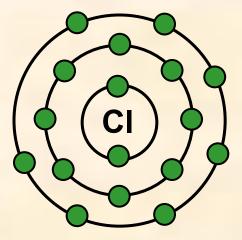
MAGNESIUM CHLORIDE



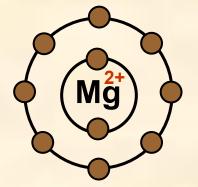
MAGNESIUM ATOM 2,8,2



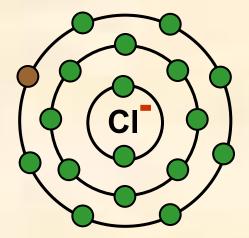
CHLORINE ATOMS 2,8,7



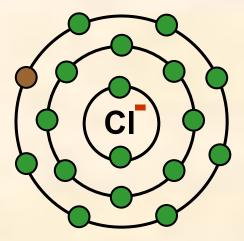
MAGNESIUM CHLORIDE



MAGNESIUM ION 2,8



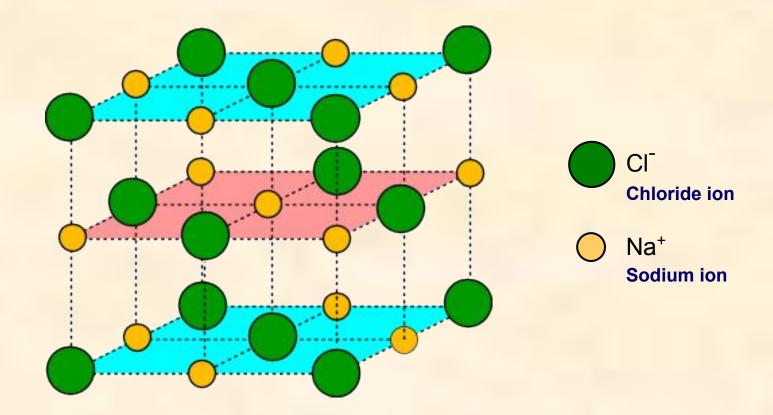
CHLORIDE IONS 2,8,8



GIANT IONIC CRYSTAL LATTICE

Oppositely charged ions held in a regular 3-dimensional lattice by electrostatic attraction

The arrangement of ions in a crystal lattice depends on the relative sizes of the ions

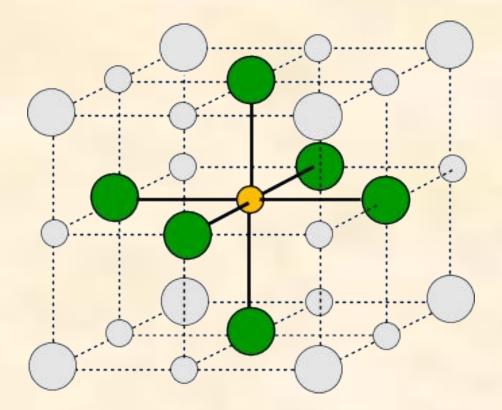


The Na⁺ ion is small enough relative to a Cl⁻ ion to fit in the spaces so that both ions occur in every plane.

GIANT IONIC CRYSTAL LATTICE

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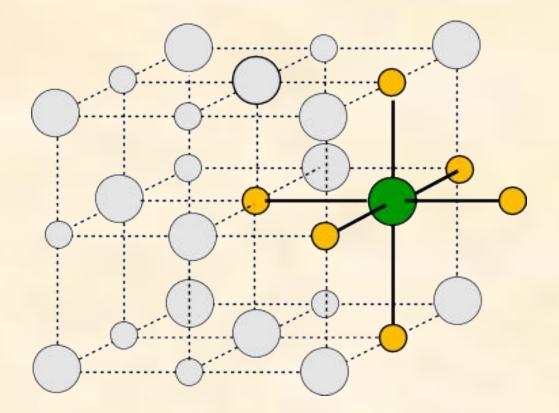


Each Na⁺ is surrounded by 6 Cl⁻ (co-ordination number = 6) and each Cl⁻ is surrounded by 6 Na⁺ (co-ordination number = 6).

GIANT IONIC CRYSTAL LATTICE

Oppositely charged ions held in a regular 3-dimensional lattice by electrostatic attraction

The arrangement of ions in a crystal lattice depends on the relative sizes of the ions



Each Na⁺ is surrounded by 6 Cl⁻ (co-ordination number = 6) and each Cl⁻ is surrounded by 6 Na⁺ (co-ordination number = 6).

Physical properties of ionic compounds

Melting point very high A large amount of energy must be put in to overcome the strong electrostatic attractions and separate the ions.

Strength

Very brittle Any dislocation leads to the layers moving and similar ions being adjacent. The repulsion splits the crystal.

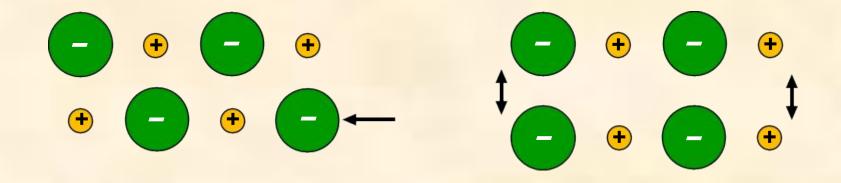
Electrical don't conduct when solid - ions held strongly in the lattice conduct when molten or in aqueous solution - the ions become mobile and conduction takes place.

Solubility Insoluble in non-polar solvents but soluble in water Water is a polar solvent and stabilises the separated ions.

Much energy is needed to overcome the electrostatic attraction and separate the ions stability attained by being surrounded by polar water molecules compensates for this

IONIC BONDING

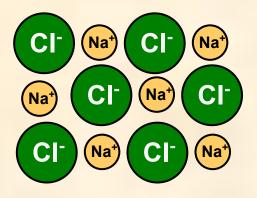
BRITTLE IONIC LATTICES



IF YOU MOVE A LAYER OF IONS, YOU GET IONS OF THE SAME CHARGE NEXT TO EACH OTHER. THE LAYERS REPEL EACH OTHER AND THE CRYSTAL BREAKS UP.

IONIC COMPOUNDS - ELECTRICAL PROPERTIES

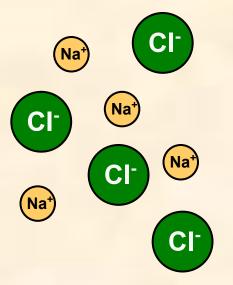
SOLID IONIC COMPOUNDS DO NOT CONDUCT ELECTRICITY



IONS ARE HELD STRONGLY TOGETHER + IONS CAN'T MOVE TO THE CATHODE - IONS CAN'T MOVE TO THE ANODE

MOLTEN IONIC COMPOUNDS DO CONDUCT ELECTRICITY

SOLUTIONS OF IONIC COMPOUNDS IN WATER DO CONDUCT ELECTRICITY



IONS HAVE MORE FREEDOM IN A LIQUID SO CAN MOVE TO THE ELECTRODES

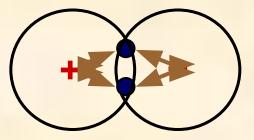
DISSOLVING AN IONIC COMPOUND IN WATER BREAKS UP THE STRUCTURE SO IONS ARE FREE TO MOVE TO THE ELECTRODES

COVALENT BONDING

COVALENT BONDING

Definition consists of a shared pair of electrons with one electron being supplied by each atom either side of the bond. compare this with dative covalent bonding

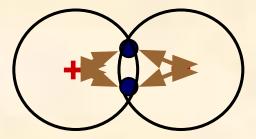
> atoms are held together because their nuclei which have an overall positive charge are attracted to the shared electrons



COVALENT BONDING

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> atoms are held together because their nuclei which have an overall positive charge are attracted to the shared electrons



Formation between atoms of the same element N₂, O₂, diamond, graphite

between atoms of different elements CO_2 , SO_2 on the RHS of the table;

when one of the elements is in the CCI_4 , $SiCI_4$ middle of the table;

with head-of-the-group elements with high ionisation energies; BeCl₂

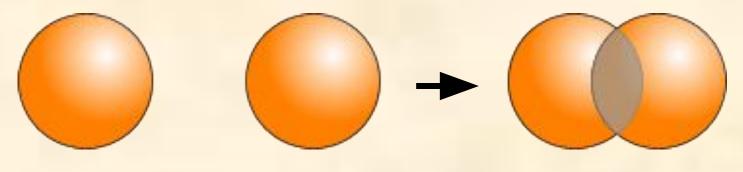
COVALENT BONDING

- atoms share electrons to get the nearest noble gas electronic configuration
- some don't achieve an "octet" as they haven't got enough electrons eg Al in AICI₃
- others share only some if they share all they will exceed their "octet" eg NH₃ and H₂O
- atoms of elements in the 3rd period onwards can exceed their "octet" if they wish as they are not restricted to eight electrons in their "outer shell" eg PCI₅ and SF₆

SIMPLE MOLECULES

Orbital theory

Covalent bonds are formed when orbitals, each containing one electron, overlap. This forms a region in space where an electron pair can be found; new molecular orbitals are formed.



orbital containing 1 electron orbital containing 1 electron overlap of orbitals provides a region in space which can contain a pair of electrons

The greater the overlap the stronger the bond.



HYDROGEN





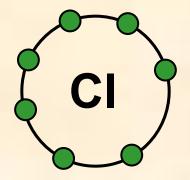
Hydrogen atom needs one electron to complete its outer shell atomAmshiaee ayotriogefre latomons to formation raceiols lancoelalatio bood A hydrogen MOLECULE is formed

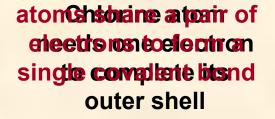
WAYS TO REPRESENT THE MOLECULE





HYDROGEN CHLORIDE







Hydrogen atom also needs one electron to complete its outer shell

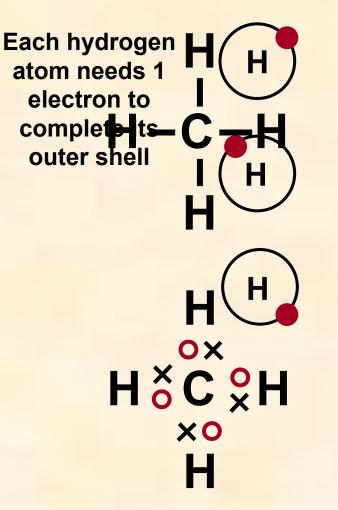
WAYS TO REPRESENT THE MOLECULE

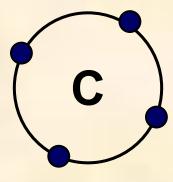
H × colo

H - CI

METHANE

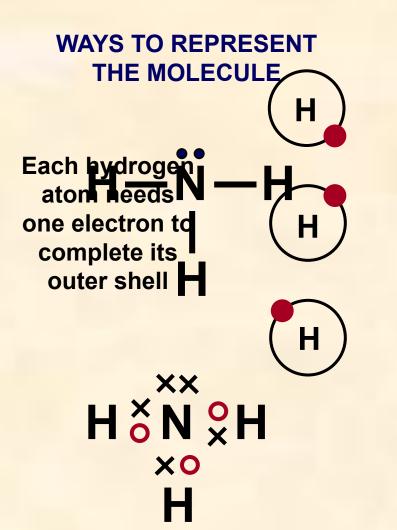


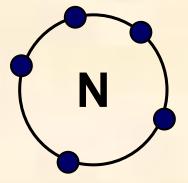




A advoors bares raded s f4 ideeteotre to fop het sing its coutale shelonds

AMMONIA

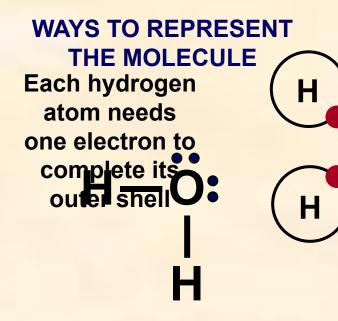


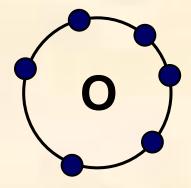


Nitrogen atom nevelsate feotrions 5 telectrople tel iteroviseris will exceed the maximum of 8

A LONE PAIR REMAINS

WATER



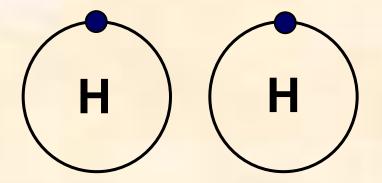


H $\stackrel{\times \times}{\stackrel{\circ}{_{\circ}}} O \stackrel{\times}{_{\times}} H$

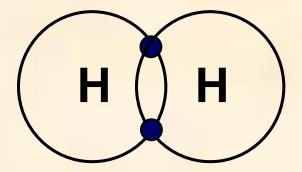
Oxygen atomonéeds 2 relêctfötss6 teleotropistetits ovitær is heill exceed the maximum of 8

2 LONE PAIRS REMAIN

HYDROGEN



both atoms need one electron to complete their outer shell



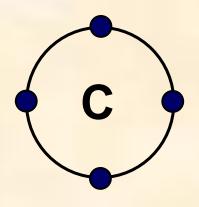
atoms share a pair of electrons to form a single covalent bond

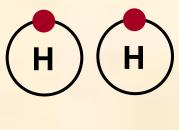
CROSS DIAGRAM

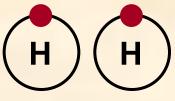
H - H

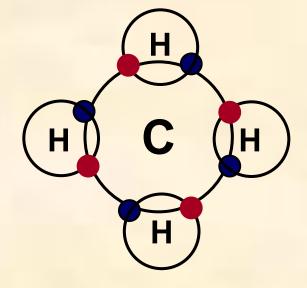


METHANE





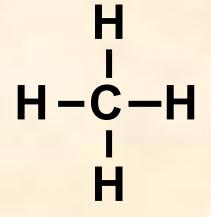




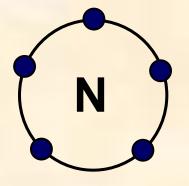
atom needs four electrons to complete its outer shell

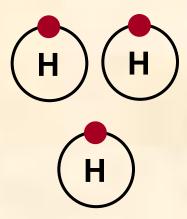
each atom needs one electron to complete its outer shell Carbon shares all 4 of its electrons to form 4 single covalent bonds

H CROSS DIAGRAM H COX H COX COX COX COX COX COX H COX COX H COX COX H COX COX H COX



AMMONIA





H Nitrogen can only share 3 of

Ν

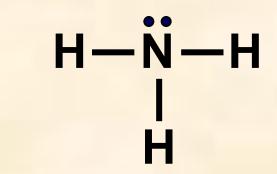
Η

atom needs three electrons to complete its outer shell each atom needs one electron to complete its outer shell

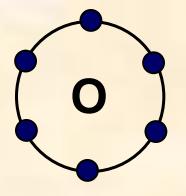
its 5 electrons otherwise it will exceed the maximum of 8

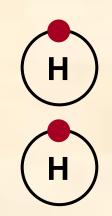
A LONE PAIR REMAINS

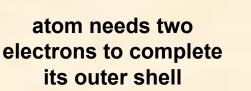
H ^{××} N [°] H



WATER

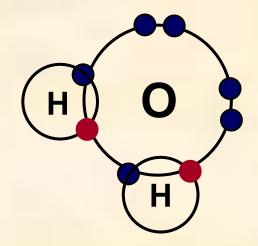






each atom needs one electron to complete its outer shell

H-Ö:



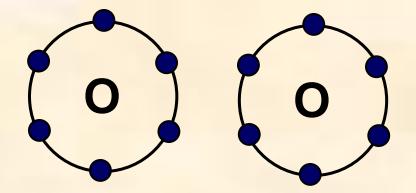
Oxygen can only share 2 of its 6 electrons otherwise it will exceed the maximum of 8

TWO LONE PAIRS REMAIN

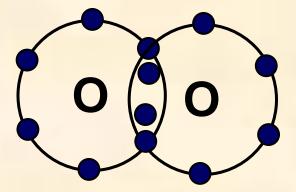
H ^{××} O × × O H

OXYGEN

 $\mathbf{O} = \mathbf{O}$



each atom needs two electrons to complete its outer shell



each oxygen shares 2 of its electrons to form a DOUBLE COVALENT BOND

SIMPLE COVALENT MOLECULES

- **Bonding** Atoms are joined together within the molecule by covalent bonds.
- **Electrical** Don't conduct electricity as they have no mobile ions or electrons
- Solubility Tend to be more soluble in organic solvents than in water; some are hydrolysed

Boiling point Low - intermolecular forces (van der Waals' forces) are weak; they increase as molecules get a larger surface area

e.g.
$$CH_4$$
 -161°C C_2H_6 - 88°C C_3H_8 -42°C

as the intermolecular forces are weak, little energy is required to to separate molecules from each other so boiling points are low

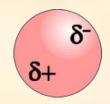
some boiling points are higher than expected for a given mass because you can get additional forces of attraction

VAN DER WAALS' FORCES INSTANTANEOUS DIPOLE-INDUCED DIPOLE FORCES

Although the bonding within molecules is strong, that between molecules is weak. Molecules and monatomic noble gases are subject to weak attractive forces.

Instantaneous dipole-induced dipole forces

Because electrons move quickly in orbitals, their position is constantly changing; at any given instant they could be anywhere in an atom. The possibility will exist that one side will have more electrons than the other. This will give rise to a dipole...



VAN DER WAALS' FORCES INSTANTANEOUS DIPOLE-INDUCED DIPOLE FORCES

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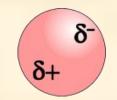
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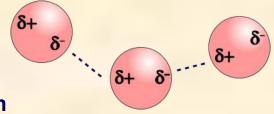
Because electrons move quickly in orbitals, their position is constantly changing; at any given instant they could be anywhere in an atom. The possibility will exist that one side will have more electrons than the other. This will give rise to a dipole...

The dipole on one atom induces dipoles on nearby atoms

Atoms are now attracted to each other by a weak forces

The greater the number of electrons, the stronger the attraction and the greater the energy needed to separate the particles.







VAN DER WAALS' FORCES INSTANTANEOUS DIPOLE-INDUCED DIPOLE FORCES

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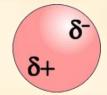
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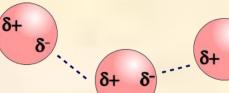
The dipole on one atom induces dipoles on others

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NOBLE GASES ALKANES Electrons B pt. B pt. Electrons 10 -161°C He 2 -269°C 10 -246°C 18 - 88°C Ne 18 -186°C 26 - 42°C Ar C,H, Kr 36 -152°C





ELECTRONEGATIVITY

'The ability of an atom to attract the electron pair in a covalent bond to itself'

Non-polar bond similar atoms have the same electronegativity they will both pull on the electrons to the same extent the electrons will be equally shared

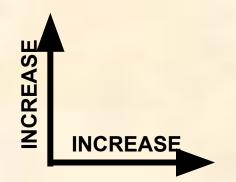
Polar bond different atoms have different electronegativities one will pull the electron pair closer to its end it will be slightly more negative than average, dthe other will be slightly less negative, or more positive, d+ a dipole is formed and the bond is said to be polar greater electronegativity difference = greater polarity

Pauling Scale a scale for measuring electronegativity

ELECTRONEGATIVITY

'The ability of an atom to attract the electron pair in a covalent bond to itself'

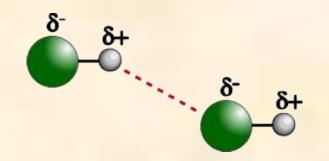
Pauling Scale a scale for measuring electronegativity values increase across periods values decrease down groups fluorine has the highest value



н							
2.1							
Li	Be	В	С	Ν	0	F	
1.0	1.5	2.0	2.5	3.0	3.5	4.0	
Na	Mg	ΑΙ	Si	Р	S	CI	
0.9	1.2	1.5	1.8	2.1	2.5	3.0	
Κ				Br			
0.8				2.8			

DIPOLE-DIPOLE INTERACTION

Occurrence occurs between molecules containing polar bonds acts in addition to the basic van der Waals' forces the extra attraction between dipoles means that more energy must be put in to separate molecules get higher boiling points than expected for a given mass

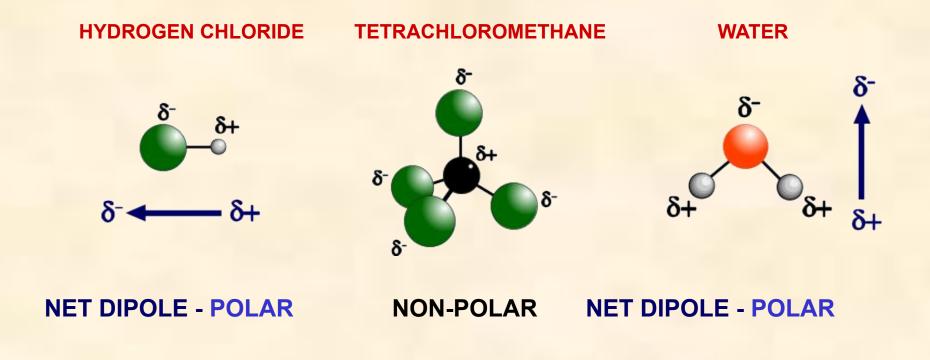


Boil	ing	poi	nts
of	hyd	rid	es

	Mr	°C		Mr	°C
CH₄	16	-161	H ₂ O	18	+100
SiH ₄		-117	H ₂ S	34	-61
GeH ₄	77	-90	H ₂ Se	81	-40
SnH ₄			H ₂ Te	130	-2
NH ₃	17	-33	HF	20	+20
PH ₃	34	-90	HCI	36.5	-85
AsH ₃	78	-55	HBr	81	-69
SbH,	125	-17	HI	128	-35

POLAR MOLECULES

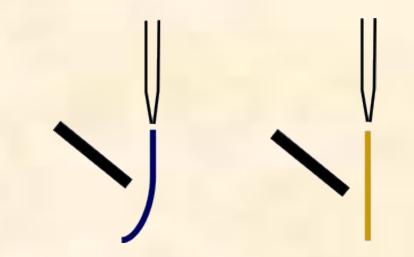
Occurrence not all molecules containing polar bonds are polar overall if bond dipoles 'cancel each other' the molecule isn't polar if there is a 'net dipole' the molecule will be polar



POLAR MOLECULES

Evidence place a liquid in a burette allow it to run out place a charged rod alongside the stream of liquid polar molecules are attracted by electrostatic attraction non-polar molecules will be unaffected



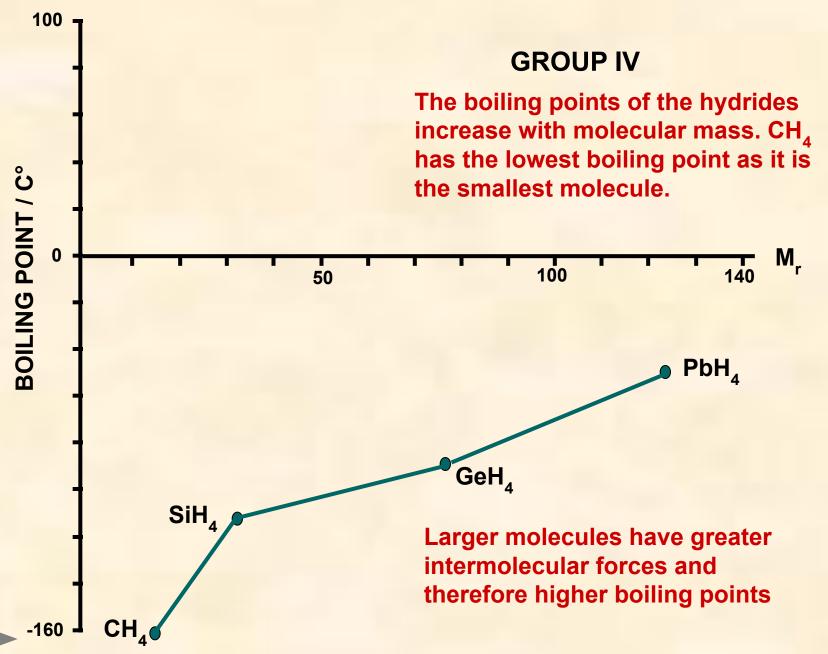


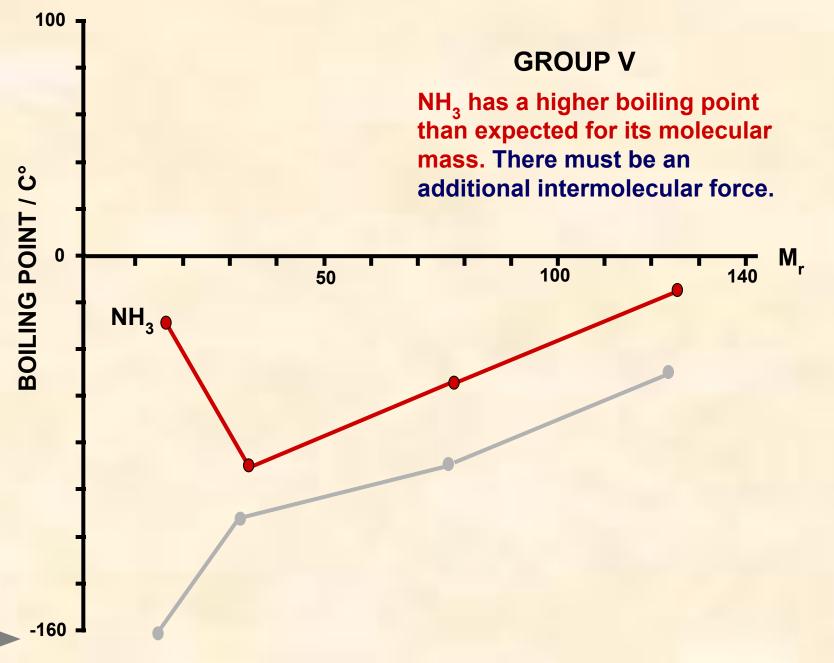
NET DIPOLE - POLAR

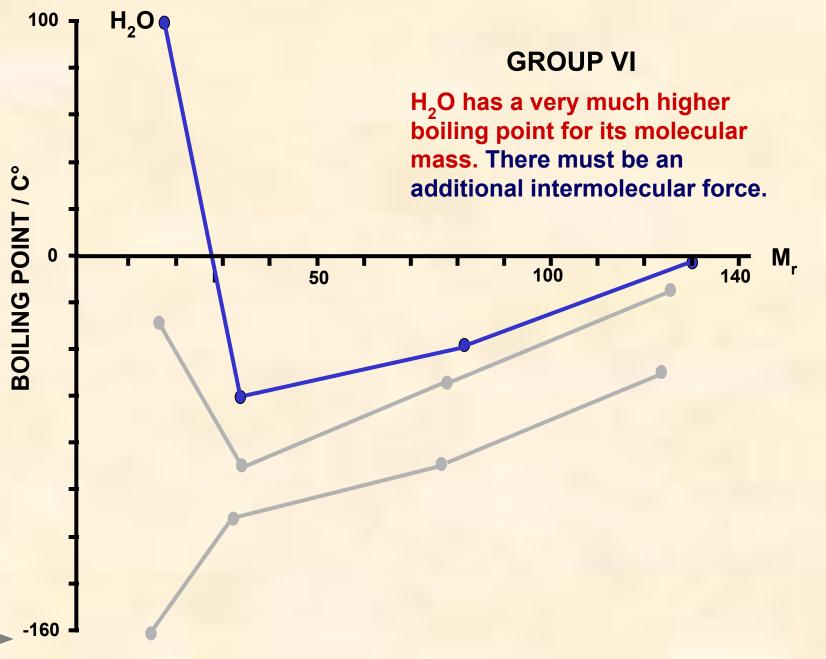
NON-POLAR

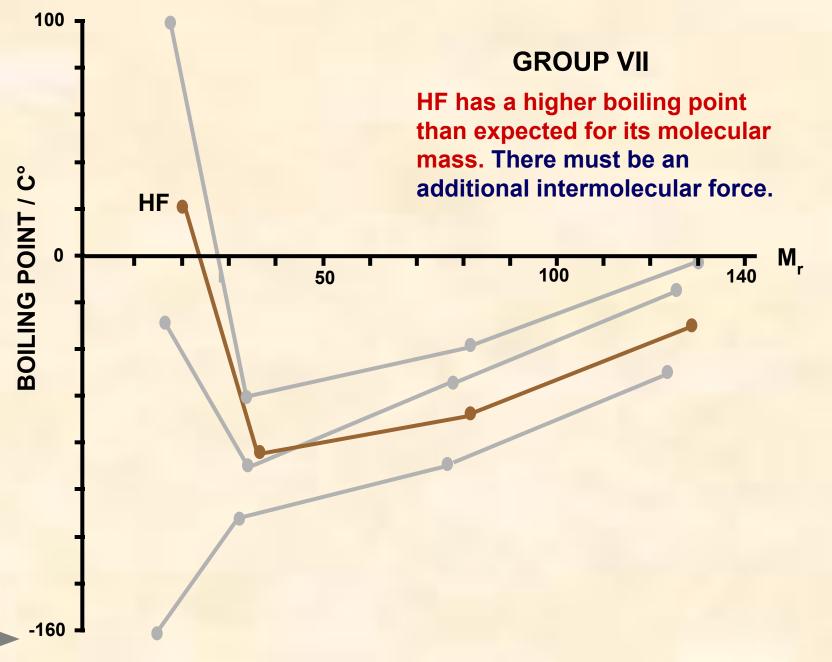
	Mr	°C		Mr	°C	
	CH₄ 16	-161		H ₂ O18	+100	
GROUP	SiH₄	32 -117	GROUP	$H_{2}S 34$	-61	
IV	GeH₄	77 -90	VI	H ₂ Se	81 -40	
	SnH ₄	123 -50		H ₂ Te	130 -2	
	NH ₃ 17			HF 20		
GROUP	PH ₃ 34		GROUP		HCI 36.5-85	
V	AsH ₃	78 -55	VII	HBr 81	-69	
	SbH ₃	125 -17		HI 128	-35	

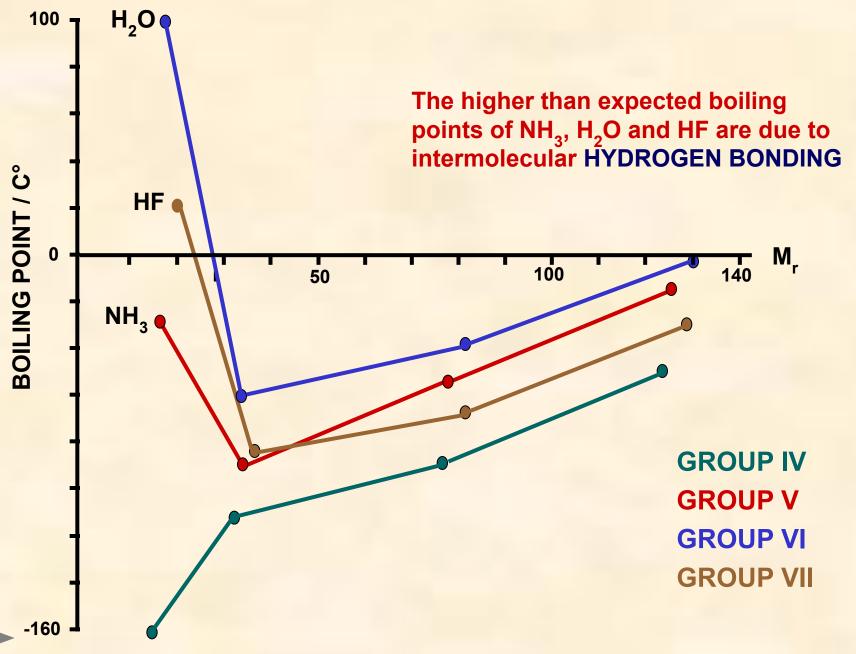
The values of certain hydrides are not typical of the trend you would expect

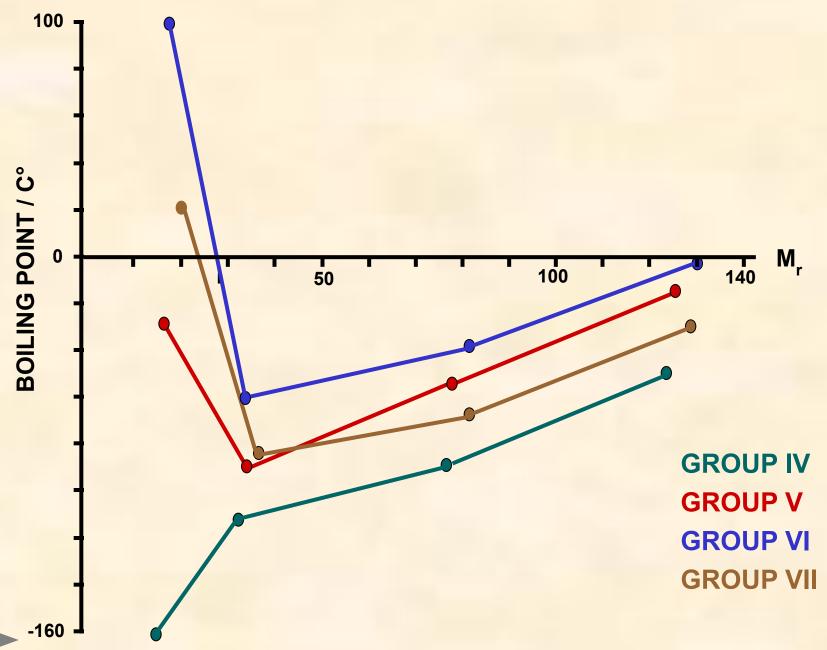












HYDROGEN BONDING

- an extension of dipole-dipole interaction
- gives rise to even higher boiling points
- bonds between H and the three most electronegative elements,
 F, O and N are extremely polar
- because of the small sizes of H, F, N and O the partial charges are concentrated in a small volume thus leading to a high charge density
- makes the intermolecular attractions greater and leads to even higher boiling points

HYDROGEN BONDING - ICE

each water molecule is hydrogen-bonded to 4 others in a tetrahedral formation

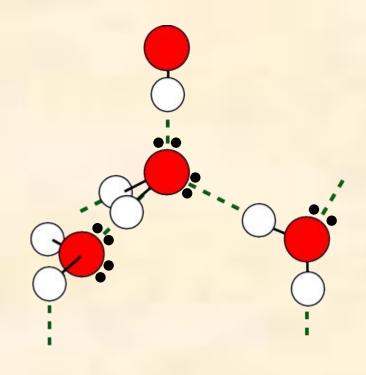
ice has a "diamond-like" structure

volume is larger than the liquid making it

when ice melts, the structure collapses slightly and the molecules come closer; they then move a little further apart as they get more energy as they warm up

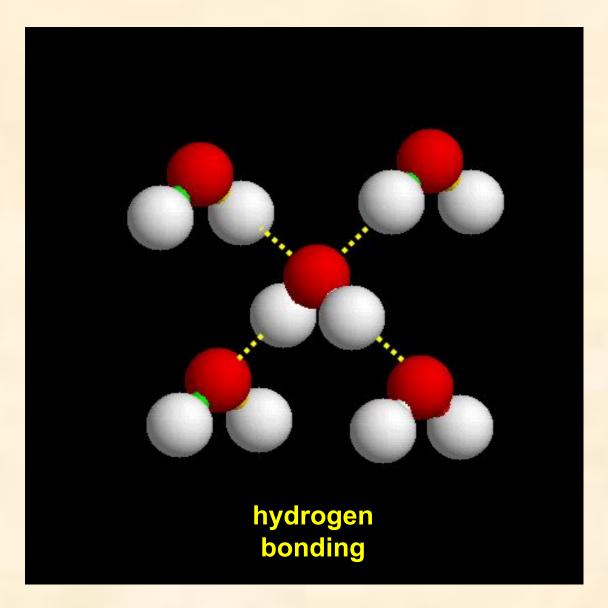
this is why...

- a) water has a maximum density at 4°C
- b) ice floats.



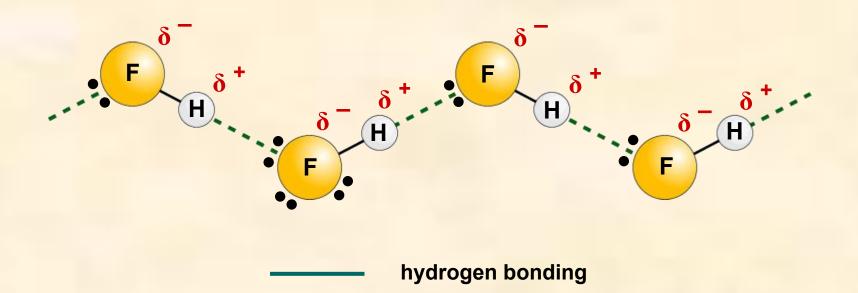
- hydrogen bondinglone pair
 - ione pa

HYDROGEN BONDING - ICE





HYDROGEN BONDING - HF



Hydrogen fluoride has a much higher boiling point than one would expect for a molecule with a relative molecular mass of 20

Fluorine has the highest electronegativity of all and is a small atom so the bonding with hydrogen is extremely polar

DATIVE COVALENT (CO-ORDINATE) BONDING

A dative covalent bond differs from covalent bond only in its formation Both electrons of the shared pair are provided by one species (donor) and it shares the electrons with the acceptor

Donor species will have lone pairs in their outer shells

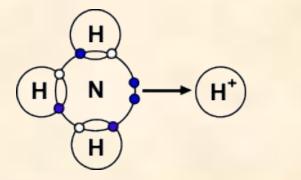
Acceptor species will be short of their "octet" or maximum.

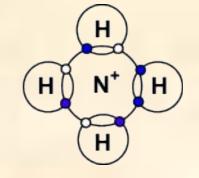
Lewis base a lone pair donor Lewis acid a lone pair acceptor

Ammonium ion, NH_{4}^{+}

The lone pair on N is used to share with the hydrogen ion which needs two electrons to fill its outer shell.

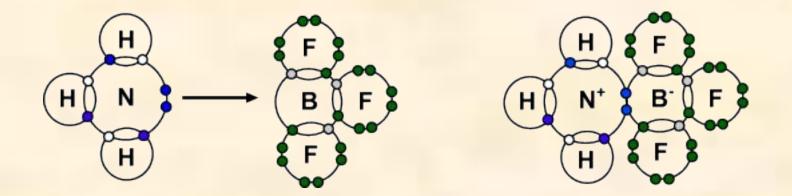
The N now has a +ive charge as - it is now sharing rather than owning two electrons.





Boron trifluoride-ammonia NH₃BF₃

Boron has an incomplete shell in BF_3 and can accept a share of a pair of electrons donated by ammonia. The B becomes -ive as it is now shares a pair of electrons (i.e. it is up one electron) it didn't have before.

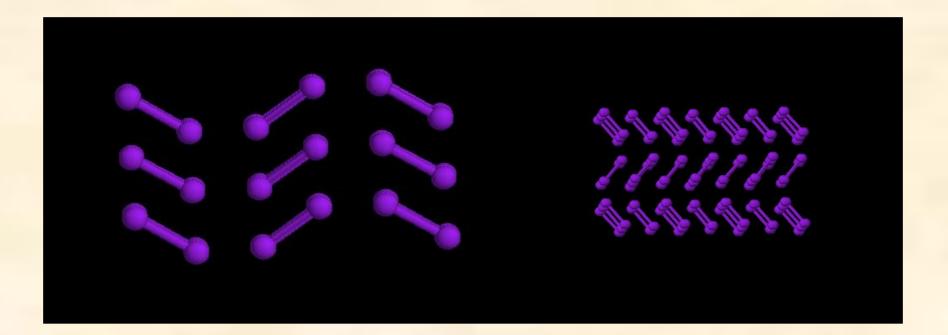


MOLECULAR SOLIDS

MOLECULAR SOLIDS

IODINE

At room temperature and pressure, iodine is a greyish solid. However it doesn't need to be warmed much in order to produce a purple vapour. This is because iodine is composed of diatomic molecules (I_2) which exist in an ordered molecular crystal in the solid state. Each molecule is independent of the others, only being attracted by van der Waals' forces. Therefore, little energy is required to separate the iodine molecules.



COVALENT NETWORKS GIANT MOLECULES MACROMOLECULES

They all mean the same!

DIAMOND, GRAPHITE and SILICA

Many atoms joined together in a regular array by a large number of covalent bonds

GENERAL PROPERTIES

MELTING POINT Very high structure is made up of a large number of covalent bonds, all of which need to be broken if atoms are to be separated

ELECTRICAL Don't conduct electricity - have no mobile ions or electrons but... Graphite conducts electricity

STRENGTH Hard - exists in a rigid tetrahedral structure Diamond and silica (SiO₂)... but Graphite is soft

DIAMOND

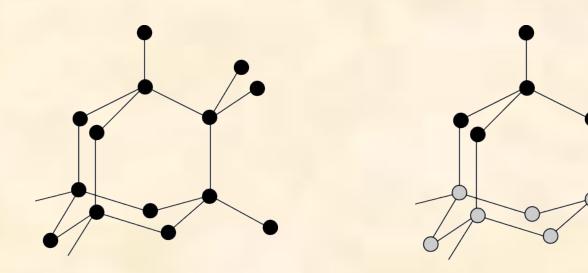
MELTING POINT VERY HIGH many covalent bonds must be broken to separate atoms

STRENGTH STRONG

each carbon is joined to four others in a rigid structure Coordination Number = 4

ELECTRICAL NON-CONDUCTOR

No free electrons - all 4 carbon electrons used for bonding



GRAPHITE

MELTING POINT VERY HIGH

many covalent bonds must be broken to separate atoms

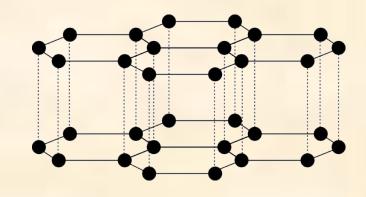
STRENGTH SOFT

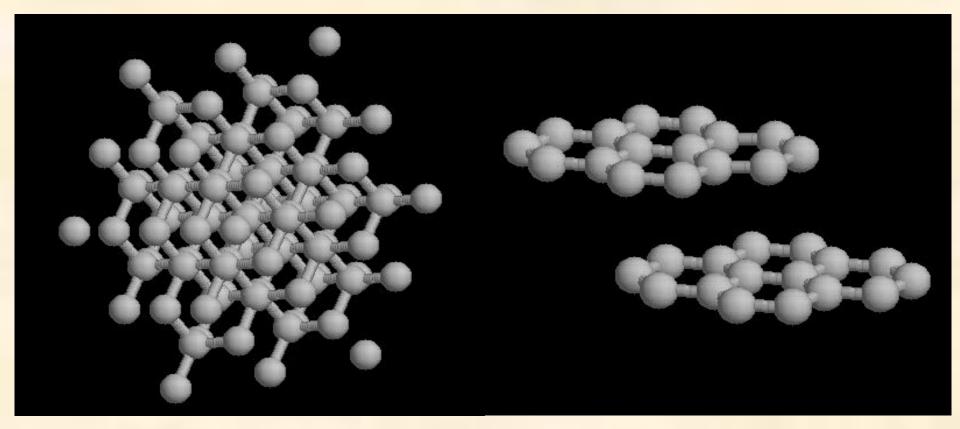
each carbon is joined to three others in a layered structure Coordination Number = 3 layers are held by weak van der Waals' forces can slide over each other

ELECTRICAL CONDUCTOR

Only three carbon electrons are used for bonding which leaves the fourth to move freely along layers

layers can slide over each other used as a lubricant and in pencils





DIAMOND



SILICA

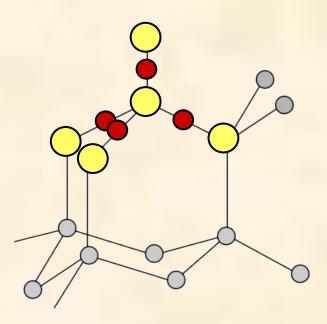
MELTING POINT VERY HIGH

many covalent bonds must be broken to separate atoms

STRENGTH STRONG

each silicon atom is joined to four oxygens - C No. = 4 each oxygen atom are joined to two silicons - C No = 2

ELECTRICAL NON-CONDUCTOR - no mobile electrons



METALLIC BONDING

METALLIC BONDING

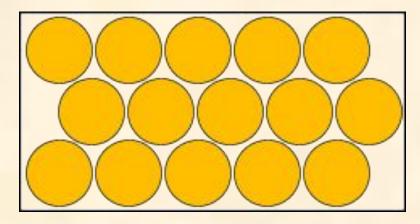
Involves a lattice of positive ions surrounded by delocalised electrons

Metal atoms achieve stability by "off-loading" electrons to attain the electronic structure of the nearest noble gas. These electrons join up to form a mobile cloud which prevents the newly-formed positive ions from flying apart due to repulsion between similar charges.

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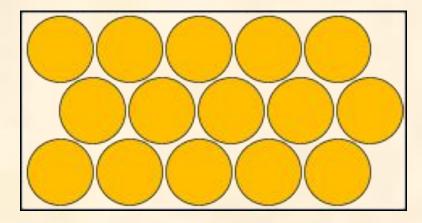


Atoms arrange in regular close packed 3-dimensional crystal lattices.

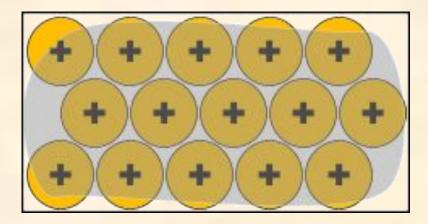
METALLIC BONDING

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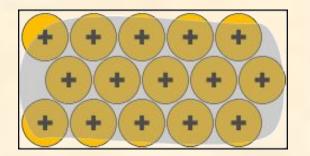
Atoms arrange in regular close packed 3-dimensional crystal lattices.



The outer shell electrons of each atom leave to join a mobile "cloud" or "sea" of electrons which can roam throughout the metal. The electron cloud binds the newly-formed positive ions together.

METALLIC BOND STRENGTH

Depends on the number of outer electrons donated to the cloud and the size of the metal atom/ion.

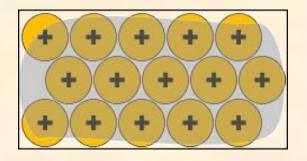


Na

The strength of the metallic bonding in sodium is relatively weak because each atom donates one electron to the cloud.

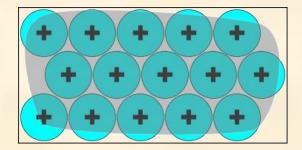
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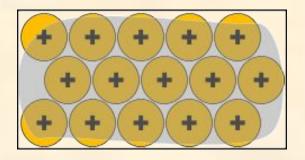


Κ

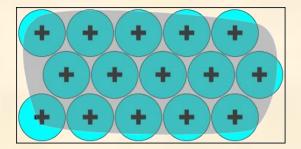
The metallic bonding in potassium is weaker than in sodium because the resulting ion is larger and the electron cloud has a bigger volume to cover so is less effective at holding the ions together.

METALLIC BOND STRENGTH

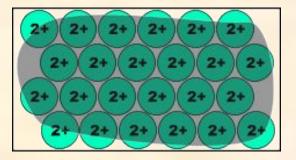
Depends on the number of outer electrons donated to the cloud and the size of the metal atom/ion.



Na



Κ



Mg

The strength of the metallic bonding in sodium is relatively weak because each atom donates one electron to the cloud.

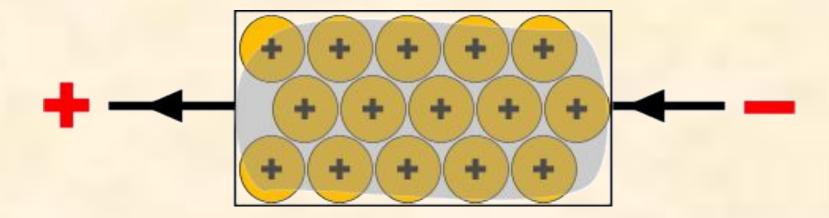
The metallic bonding in potassium is weaker than in sodium because the resulting ion is larger and the electron cloud has a bigger volume to cover so is less effective at holding the ions together.

The metallic bonding in magnesium is stronger than in sodium because each atom has donated two electrons to the cloud. The greater the electron density holds the ions together more strongly.

Metals are excellent conductors of electricity

For a substance to conduct electricity it must have mobile ions or electrons.

Because the ELECTRON CLOUD IS MOBILE, electrons are free to move throughout its structure. Electrons attracted to the positive end are replaced by those entering from the negative end.



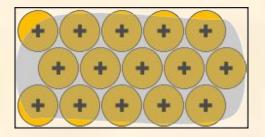
MOBILE ELECTRON CLOUD ALLOWS THE CONDUCTION OF ELECTRICITY

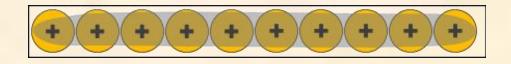
Metals can have their shapes changed relatively easily

MALLEABLE CAN BE HAMMERED INTO SHEETS

DUCTILE CAN BE DRAWN INTO RODS AND WIRES

As the metal is beaten into another shape the delocalised electron cloud continues to bind the "ions" together.





Some metals, such as gold, can be hammered into sheets thin enough to be translucent.

HIGH MELTING POINTS

Melting point is a measure of how easy it is to separate individual particles. In metals it is a measure of how strong the electron cloud holds the + ions.

The ease of separation of ions depends on the...

ELECTRON DENSITY OF THE CLOUD IONIC / ATOMIC SIZE

 PERIODS Na (2,8,1)
 <</td>
 Mg (2,8,2)
 <</td>
 AI (2,8,3)

 m.pt
 98°C
 650°C
 659°C

 b.pt
 890°C
 1110°C
 2470°C

 Na⁺

MELTING POINT INCREASES ACROSS THE PERIOD

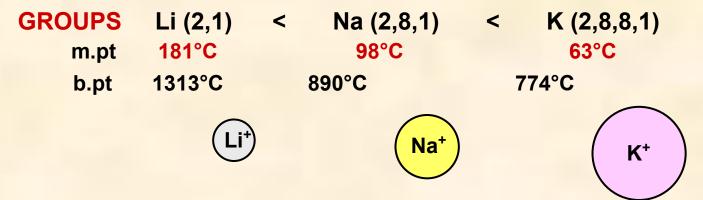
THE ELECTRON CLOUD DENSITY INCREASES DUE TO THE GREATER NUMBER OF ELECTRONS DONATED PER ATOM. AS A RESULT THE IONS ARE HELD MORE STRONGLY.

HIGH MELTING POINTS

Melting point is a measure of how easy it is to separate individual particles. In metals it is a measure of how strong the electron cloud holds the + ions.

The ease of separation of ions depends on the...

ELECTRON DENSITY OF THE CLOUD IONIC / ATOMIC SIZE



MELTING POINT INCREASES DOWN A GROUP

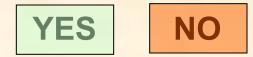
IONIC RADIUS INCREASES DOWN THE GROUP. AS THE IONS GET BIGGER THE ELECTRON CLOUD BECOMES LESS EFFECTIVE HOLDING THEM TOGETHER SO THEY ARE EASIER TO SEPARATE.

REVISION CHECK

What should you be able to do?

Recall the different types of physical and chemical bonding
Understand how ionic, covalent, dative covalent and metallic bonding arise
Recall the different forms of covalent structures
Understand how the physical properties depend on structure and bonding
Understand how different types of physical bond have different strengths
Recall and explain the variation in the boiling points of hydrides
Balance ionic equations
Construct diagrams to represent covalent bonding

CAN YOU DO ALL OF THESE?



You need to go over the relevant topic(s) again Click on the button to return to the menu



WELL DONE!

Try some past paper questions

AN INTRODUCTION TO BONDING THE END



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