

From Models to Materials

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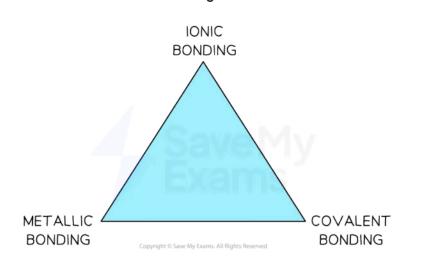
Bonding Models

Bonding Models

Bonding Models

- Models simplify complex systems and allow us to predict and test theories
- As we have seen, chemical bonding broadly falls into three types: ionic, covalent and metallic
- The bonding types can be used to explain chemical and physical properties of substances
- However, like all models, there are limitations and inaccuracies that arise from oversimplification
- For example, take a substance like aluminium chloride, AICI_{3.} The compound consists of a metal and non-metal, so the traditional bonding approach would be to predict it has ionic bonding and the associated properties of an ionic compound such as high melting point and boiling point
 - Aluminium chloride actually melts at 192°C, so it does not behave like an ionic compound
- The fact we know polar covalent bonds exist arise provides evidence that bonding type is not something that can be easily compartmentalised
- Bonding is best thought of as a continuum of the three different bonding types like the area of an
 equilateral triangle

A bonding model



Chemical bonding is a continuum of ionic, covalent and metallic bonding

- The location of an element or compound in the bonding triangle is determined by the electronegativity values of the elements present
- The bonding triangle is anchored by two axes, electronegativity difference on the y-axis and average electronegativity on the x-axis
- The symbol for electronegativity is the Greek letter (chi) χ, pronounced 'ky' as in 'sky'
 - The average electronegativity of two elements, A and B would be: $\Sigma \chi = \frac{\chi_A + \chi_B}{2}$

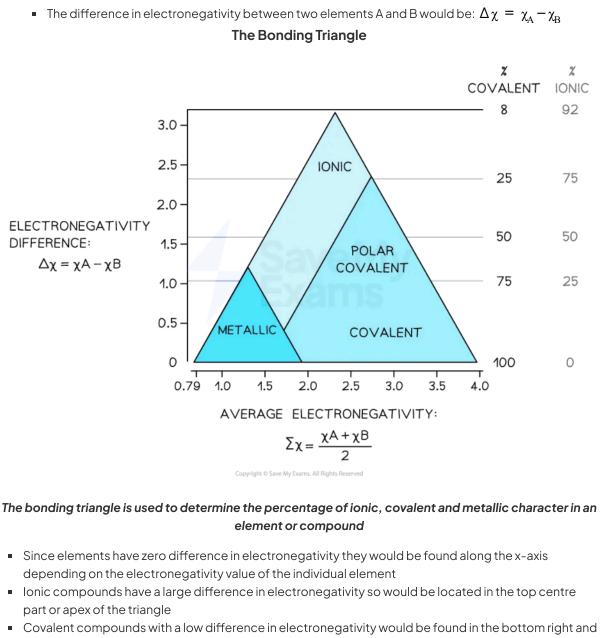
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Your notes



polar covalent compounds in the zone between ionic and covalent

😧 Examiner Tip

- You don't need to learn the bonding triangle as it is found in Section 17 of the Data Booklet
- Electronegativity values are found in the Periodic Table in Section 9.

Bonding & Properties

Bonding & Properties

Determining the position of a compound in the bonding triangle

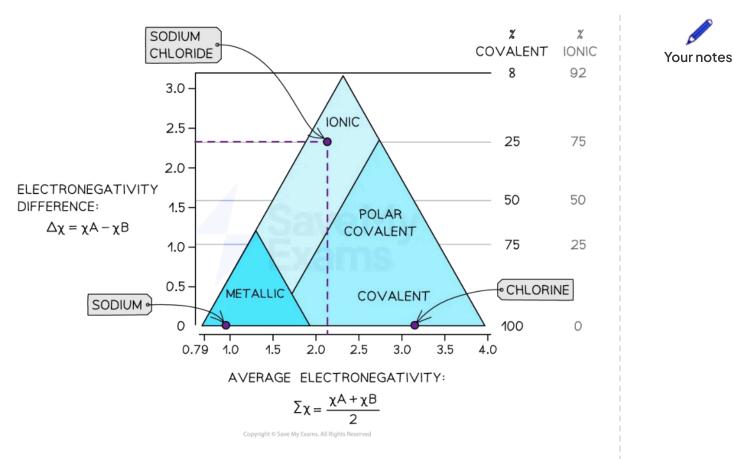
- We can use the electronegativity of elements and binary compounds to find their position in the bonding triangle
- For example, take sodium chloride:

Substance	Average electronegativity	Electronegativity difference	Where it is in the triangle
Na	0.9	0.9 - 0.9 = 0.0	Bottom left of the triangle. Na is 100%
			metallic
Cl ₂	3.2	3.2 - 3.2 = 0.0	Bottom right corner. Cl ₂ is 100% covalent
NaCl	$\Sigma \chi = \frac{\chi_A + \chi_B}{2} = \frac{3.2 + 0.9}{2} = 2.1$	$\Delta \chi = \chi_{\rm A} - \chi_{\rm B} = 3.2 - 0.9 = 2.3$	Centre top of the triangle. It is about 70% ionic on
			the triangle

Sodium chloride bonding triangle



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The location of sodium chloride on the bonding triangle using electronegativity values

Worked example

Use the bonding triangle and electronegativity values from Section 9 of the data booklet to mark the location for the following substances:

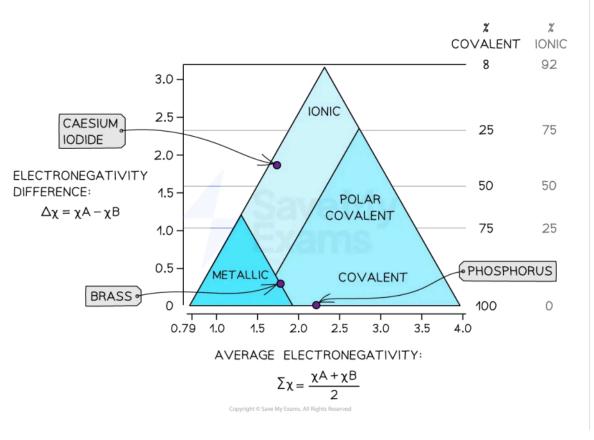
a) phosphorus

b) caesium iodide

c) brass (an alloy of copper and zinc)

Answer

Substance	Average electronegativity	Electronegativity difference
a) phosphorus	2.2	0
b) caesium chloride	$\Sigma \chi = \frac{\chi_{\rm A} + \chi_{\rm B}}{2} = \frac{0.8 + 2.7}{2} = 1.8$	$\Delta \chi = \chi_{\rm A} - \chi_{\rm B} = 2.7 - 0.8 = 1.9$
c) brass (an alloy of copper and zinc)	$\Sigma \chi = \frac{\chi_{\rm A} + \chi_{\rm B}}{2} = \frac{1.9 + 1.6}{2} = 1.8$	$\Delta \chi = \chi_{\rm A} - \chi_{\rm B} = 1.9 - 1.6 = 0.3$



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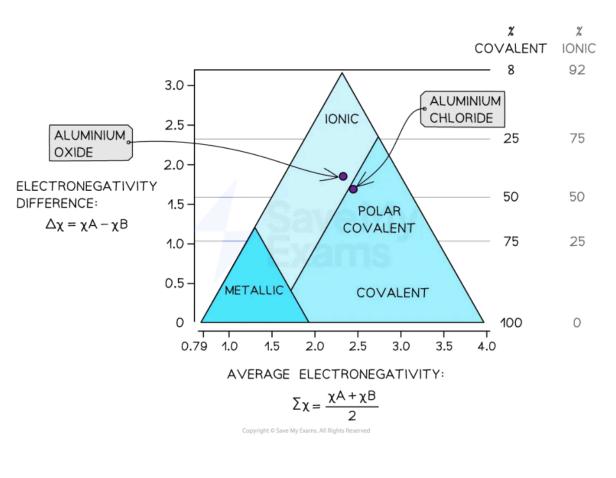


Percentages of Bonding Type

- We can also use the bonding triangle to assess the percentage of ionic or covalent character in a compound
- Take aluminium chloride and aluminium oxide as an example:

Substance	Average electronegativity	Electronegativity difference		
aluminium oxide	$\Sigma \chi = \frac{\chi_A + \chi_B}{2} = \frac{1.6 + 3.4}{2} = 2.5$	$\Delta \chi = \chi_{A} - \chi_{B} = 3.4 - 1.6 =$		
aluminium chloride	$\Sigma \chi = \frac{\chi_A + \chi_B}{2} = \frac{1.6 + 3.2}{2} = 2.4$	$\Delta \chi = \chi_{A} - \chi_{B} = 3.2 - 1.6 =$		





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The percentage of bonding type character in aluminium oxide and aluminium chloride can be assessed using the bonding triangle

- From this analysis, we can see why there is a significant difference in properties between aluminium oxide and aluminium chloride
 - The melting point of the oxide is 2072 °C, whereas the chloride is 192°C
- The bonding triangle continuum allows for a more accurate assessment of bonding type and the prediction of associated properties

Examiner Tip

Calculations of percentage ionic character are not required.

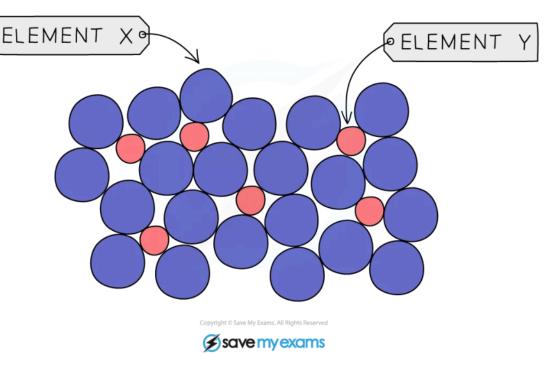


Properties of Alloys

Alloys

What is an alloy?

- Alloys are mixtures of metals, where the metals are mixed together physically but are not chemically combined
- They can also be made from metals mixed with nonmetals such as carbon
- lons of the different metals are spread throughout the lattice and are bound together by the delocalized electrons
- It is possible to form alloys because of the **non-directional** nature of the metallic bonds



Particle Diagram of an Alloy

In a metallic lattice the regular structure of metal cations (shown by Element Y) is disrupted by the presence of another element (Element X)

Why do alloys have different properties to pure metals?

- Alloys have distinct properties due to the different packing of the cations in the lattice
- Alloys often have properties that can be very **different** to the metals they contain, for example they can have greater **strength**, **hardness** or **resistance** to **corrosion** or extreme **temperatures**

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- Alloys contain atoms of **different** sizes, which **distorts** the regular arrangements of cations
- This makes it more difficult for the layers to slide over each other, so they are usually much harder than the pure metal
- Below is a table of some common alloys and their uses:

Common Alloys and their Uses Table

Alloy	Elements present	Properties	Uses
Brass	copper and zinc	strong and resistant to corrosion	door handles, hinges, musical instruments
Steel	iron, carbon and other elements like chromium, vanadium and molybdenum	very strong	construction, bridges, cars
Stainless Steel	iron, chromium, nickel, carbon	corrosion resistant	cutlery, surgical instruments, cookware
Solder	lead and tin	low melting point	joining metals in electrical circuits and jewellery
Bronze	copper and tin	hard and strong resistance to corrosion	medals, sculptures, ship fittings

Examiner Tip

You don't need to learn the specific alloys, but you should be able to use examples you know to explain why alloys have the properties they do compared to pure metals



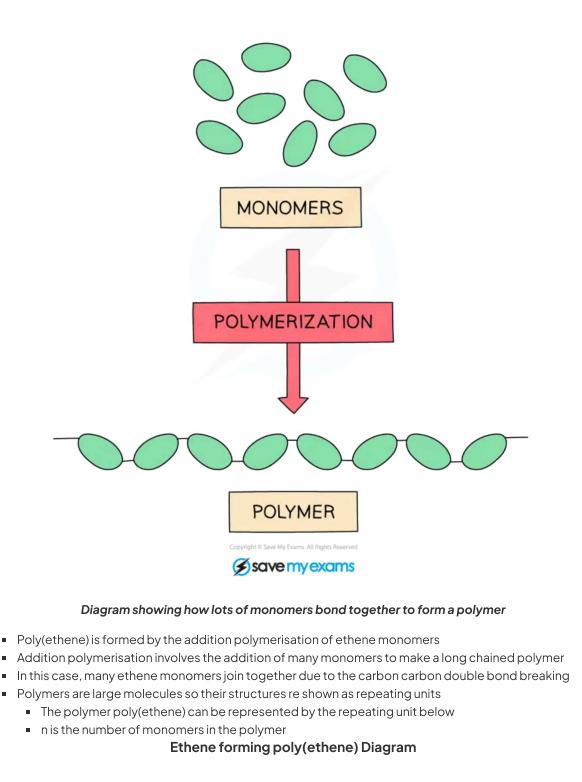
Polymers

Polymers

- Polymers are large molecules built by linking 50 or more smaller molecules called **monomers**
- Polymers are known as macromolecules as they are relatively large compared with other molecules
- Each repeat unit is connected to the adjacent units via covalent bonds
- Some polymers contain just one type of unit
 - Examples include poly(ethene) and poly(chloroethene), commonly known as PVC
- Others contain two or more different types of monomer units and which are called **copolymers**
 - Examples include nylon and biological proteins
- Different linkages also exist, depending on the monomers and the type of polymerisation
 - Examples of linkages are covalent bonds, amide links and ester links
 - Monomers Forming Polymers Diagram

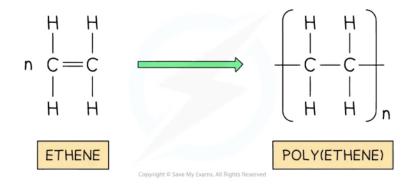


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Poly(ethene) is formed by addition polymerisation using ethene monomers

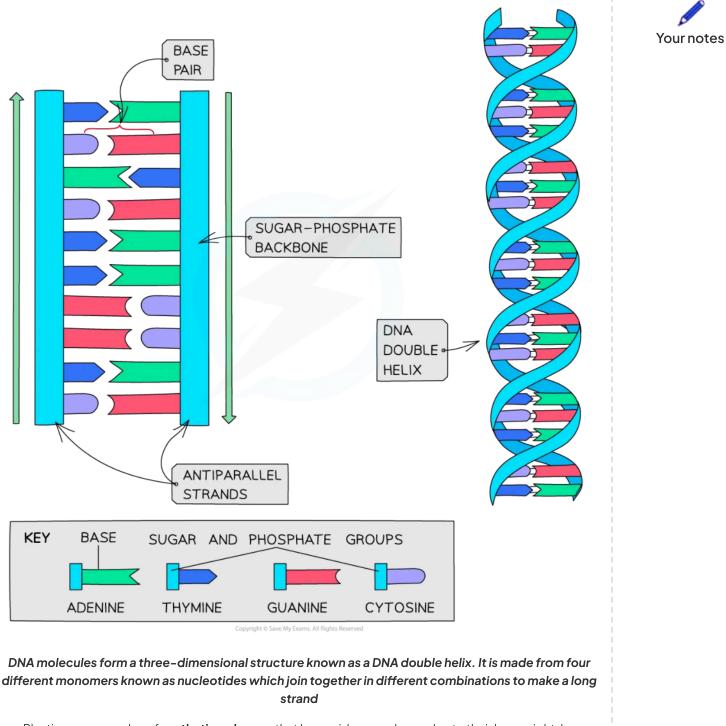
Properties of plastics

- Low weight
 - Polymers are loosely packed so will be less dense and lighter than other materials such as metals
- Unreactive
 - The addition polymers made from alkenes are saturated compounds because they do not contain double bonds and the main carbon chain is non-polar and will therefore are unreactive
- Water resistant
 - Polymers are hydrophobic so they repel water instead of absorbing it
- Strong
 - Polymers so are made up of many strong covalent bonds between the monomers
- These properties make plastics useful for packaging, construction, clothing and transportation

Natural and synthetic polymers

- Proteins, DNA and starch are examples of **natural polymers**
 - Understanding the structure and function of natural polymers has been a major focus of biochemical research for the last 100 years and have contributed to the advancement of vaccines and medicines

DNA is an example of a natural polymer



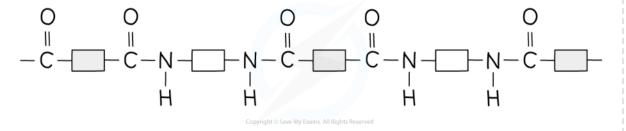
- Plastics are examples of synthetic polymers that have widespread uses due to their low weight, low reactivity, water resistance and strength
 - They have become widely distributed across the world
 - Their low reactivity means that they are **non-biodegradable** so will not break down naturally.

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• The extensive accumulation of plastics in natural environments remains one of the most pressing environmental concerns

Nylon-6,6 is an example of a synthetic polymer





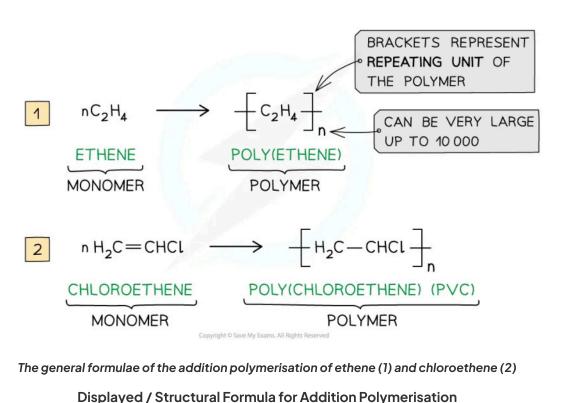
Nylon-6,6 is formed by condensation polymerisation

Addition Polymers

Addition Polymers

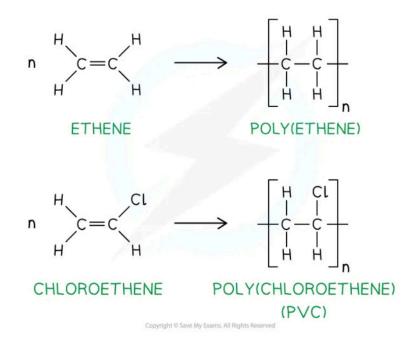
What is addition polymerisation?

- Addition polymerisation is one of the most important addition reactions of alkenes which form the basis of the plastic industry
- Addition polymerisation is the reaction in which many monomers containing at least one C=C double bond form long chains of polymers as the only product
 - Just like in other addition reactions of alkenes, the π-bond in each C=C bond breaks and then the monomers link together to form new C-C single bonds
- A **polymer** is a long-chain molecule that is made up of many repeating units
- The small, reactive molecules that react together to form the polymer are called **monomers**
- A polymerisation reaction can be represented by a general formula or by using structural / displayed formulae
 - E.g. poly(ethene) and poly(chloroethene) (also known as PVC) are polymers made up of the ethene and chloroethene monomers respectively and are commonly used in making plastics



General Formula Addition Polymerisation

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The addition polymerisation of ethene (1) and chloroethene (2)

Just like any other addition reaction of alkenes, addition polymerisation gives only one product

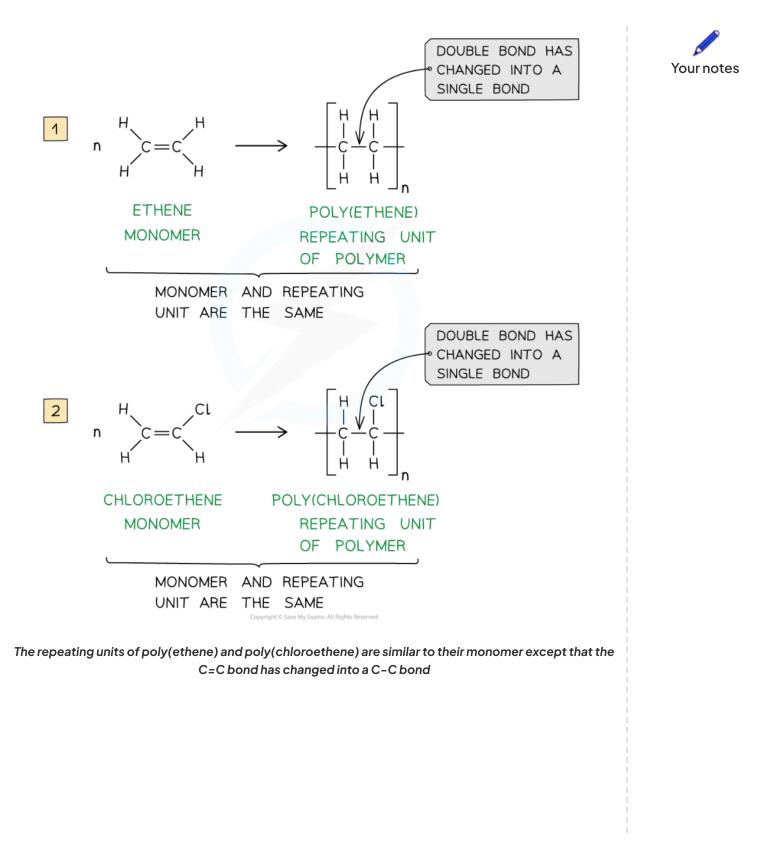
Deducing repeat units

- A **repeat unit** is the smallest group of atoms that when connected one after the other make up the polymer chain
 - It is represented by square brackets in the displayed and general formula
- In poly(alkenes) (such as poly(ethene)) and substituted poly(alkenes) (such as PVC) made of one type of monomer the repeating unit is the same as the monomer except that the C=C double bond is changed to a C-C single bond

Repeating Units for Addition Polymerisation

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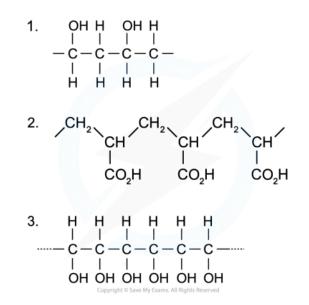
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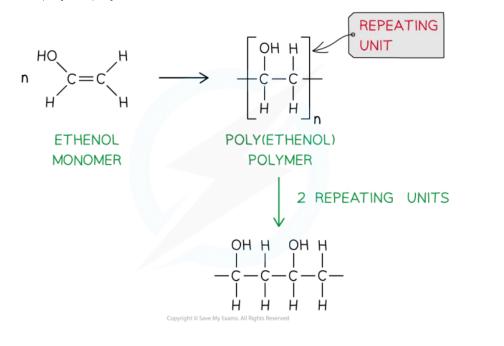
Worked example

Identify the monomers present in the given sections of addition polymer molecules:



Answer 1:

- When ethenol (CH(OH)=CH₂) is polymerised, the C-C double bond opens to produce a repeating unit of CH(OH)-CH₂
- This gives the polymer poly(ethenol)

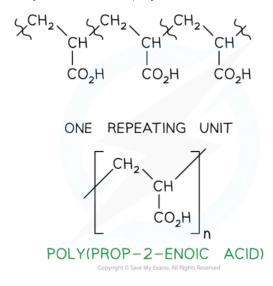


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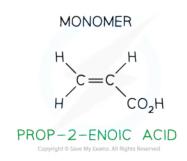


Answer 2:

- To find the monomer, first the repeating unit should be deduced
- Repeating units have only 2 carbons in the polymer main chain



Since the repeating unit is now found, it can be concluded that the monomer is prop-2-enoic acid



Answer 3:

- Again, the repeating unit only has 2 carbons in the polymer chain which in this case are two carbon atoms that each contain one OH group
- Thus, when ethene-1,2-diol (CH(OH)=CH(OH)) is polymerised, the C=C double bond opens to produce a repeating unit of CH(OH)-CH(OH) which gives the polymer poly(ethene-1,2-diol)

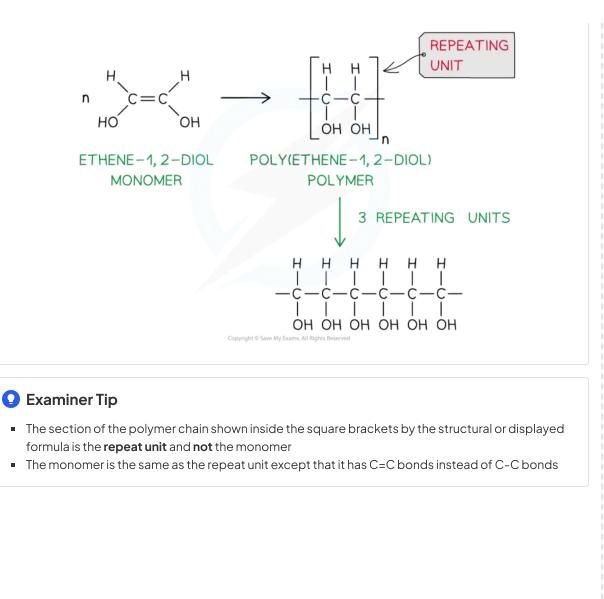


Your notes

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Condensation Polymers (HL)

Condensation Polymers

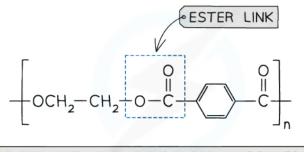
What is condensation polymerisation?

- Condensation polymerisation is another type of reaction whereby a polymer is produced by repeated condensation reactions between monomers
- Natural condensation polymers are all formed by elimination of water
 - Although the process of condensation polymerisation involves the elimination of a small molecule
- Condensation polymers can be identified because the monomers are linked by ester or amide links
- Condensation polymers can be formed by:
 - Dicarboxylic acids and diols
 - Hydroxycarboxylic acids
 - Dicarboxylic acids and diamines
 - Amino acids

What is polyester?

- Polyester is a polymer, most commonly formed by the condensation polymerisation of dicarboxylic acid monomers and diol monomers
- A polyester is produced by linking these monomers with ester bonds / links

Ester Link in a Polyester



THE POLYMER STRUCTURE FOR A POLYESTER

This polymer structure shows an ester functional group linking monomers together

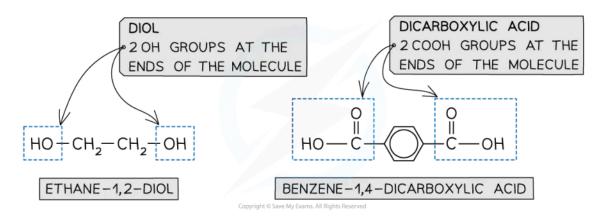
Formation of polyesters using diols and dicarboxylic acids

- A diol and a dicarboxylic acid are required to form a polyester
 - A diol contains two alcohol -OH groups
 - A dicarboxylic acid contains two carboxylic acid -COOH groups

Diol and dicarboxylic acid monomer structures

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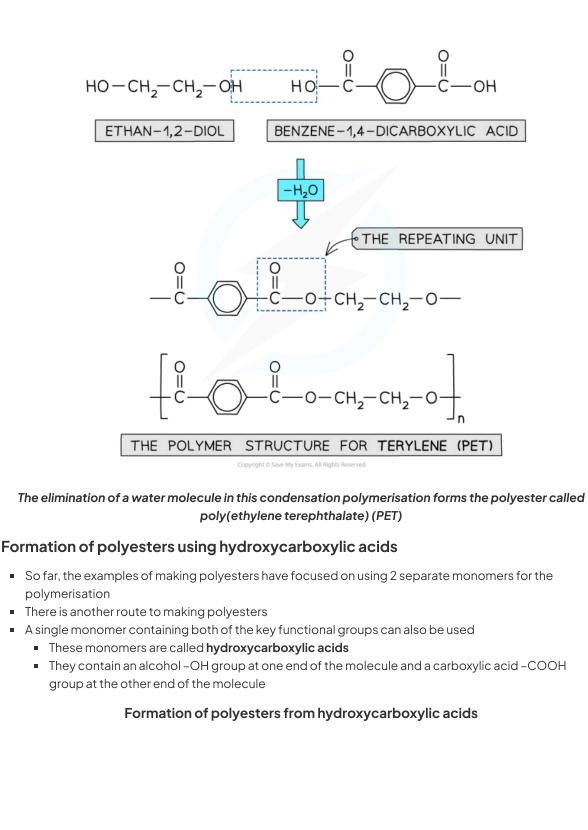
The position of the functional groups on both of these monomer molecules allows condensation polymerisation to take place effectively

- When the polyester is formed, one of the diol –OH groups and a hydrogen atom from one of the carboxylic acid –COOH groups are eliminated as a water molecule (H₂O)
- The resulting polymer is a polyester
 - For example, the polyester **poly(ethylene terephthalate)** or PET, which is sometimes known by its brand names of Terylene or Dacron

Formation of Poly(ethylene terephthalate)

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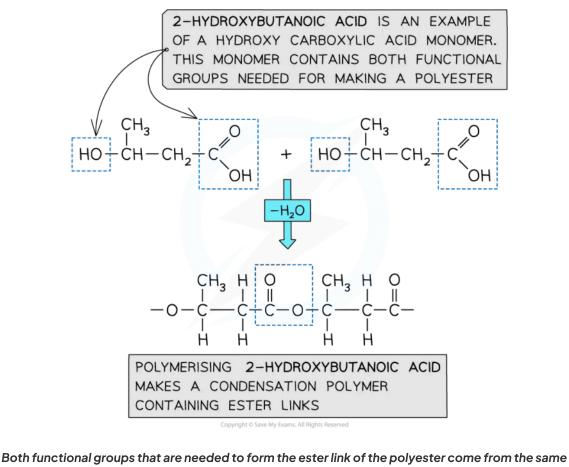
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Your notes

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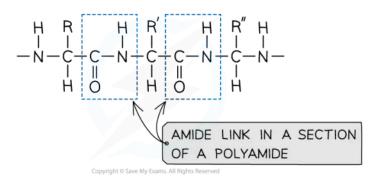


monomer

What are polyamides?

- Polyamides are polymers where the repeating units are bonded together by amide links
- The formula of an amide group is -CONH

Amide links in Polyamides





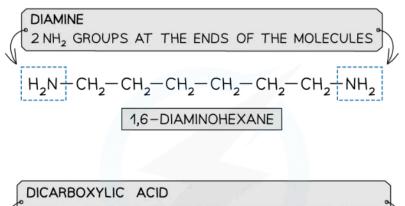
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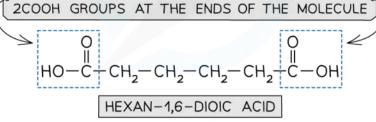
An amide link - also known as a peptide link - is the key functional group in a polyamide

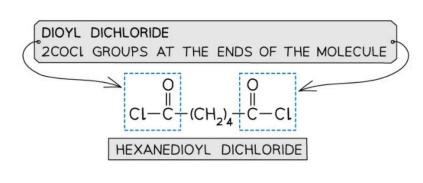
Polyamide monomers

- A diamine and a dicarboxylic acid are required to form a polyamide
 - A diamine contains 2 NH₂ groups
 - A dicarboxylic acid contains 2 COOH groups
- Dioyl dichlorides can also used to react with the diamine instead of the acid
 - A dioyl chloride contains 2 COCl groups
 - This is a more reactive monomer but more expensive than dicarboxylic acid

Polyamide monomers





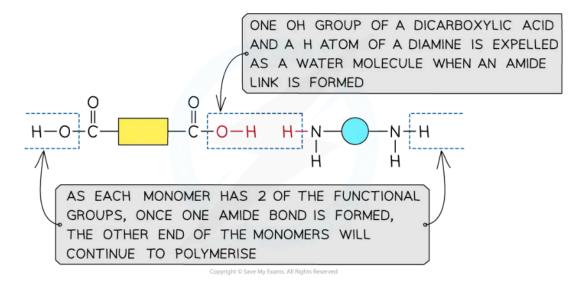


The monomers for making polyamides

Formation of polyamides

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Formation of Polyamides from Monomers



This shows the elimination of a small molecule as the amide link forms

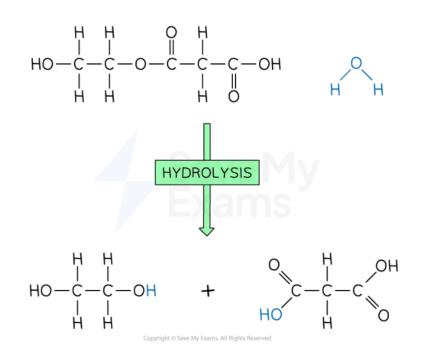
Biodegradable polymers

- Both polyesters and polyamides can be broken down using hydrolysis reactions
- This is a major advantage over the addition polymers produced using alkene monomers (polyalkenes)
- When polyesters and polyamides are taken to landfill sites, they can be broken down easily and their products used for other applications
- Hydrolysis occurs during
 - Digestion
 - Decomposition
- Both condensation and hydrolysis reactions are controlled by **enzymes**

Diagram to show hydrolysis of condensation polymers



Your notes



Water is added which causes the polymer to break down into the original monomers