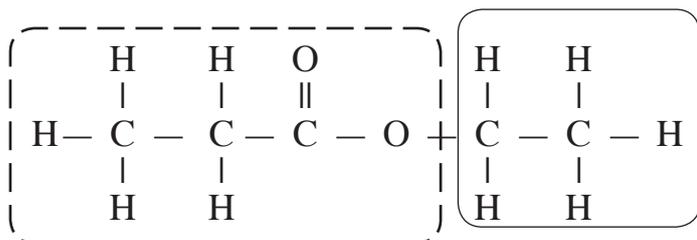
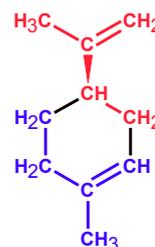


Higher Chemistry



ethyl propanoate



Limonene

(skin of citrus fruits)

Unit 2:

Nature's Chemistry

Student:

<p>A-PINENE</p> <p>ANTI-INFLAMMATORY BRONCHODILATOR AIDS MEMORY ANTI-BACTERIAL</p> <p>also found in pine needles</p> 	<p>LINALOOL</p> <p>ANESTHETIC ANTI-CONVULSANT ANALGESIC ANTI-ANXIETY</p> <p>also found in lavender</p> 	<p>BETA CARYOPHYLLENE</p> <p>ANTI-INFLAMMATORY ANALGESIC PROTECTS CELLS LINING THE DIGESTIVE TRACT</p> <p>also found in black pepper</p> 	<p>MYRCENE</p> <p>CONTRIBUTES TO SEDATIVE EFFECT OF STRONG INDICAS SLEEP AID MUSCLE RELAXANT</p> <p>also found in hops</p> 	<p>LIMONENE</p> <p>TREATS ACID REFLUX ANTI-ANXIETY ANTIDEPRESSANT</p> <p>also found in citrus</p> 
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Topic 6

Acids & Esters

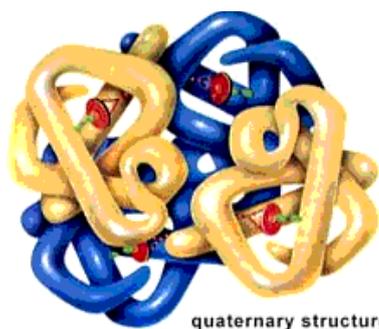
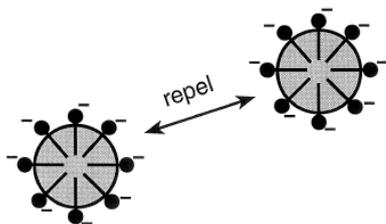
Fats & Oils

Soaps, Detergents &

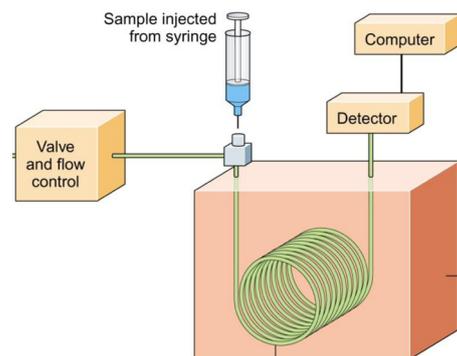
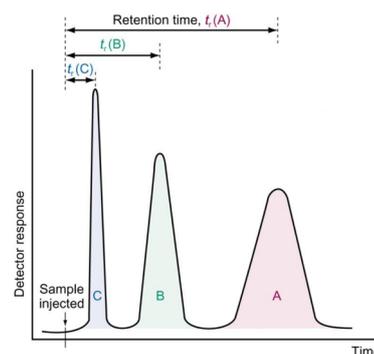
Emulsifiers

Terpenes

Proteins



quaternary structure
(aggregation of two or more peptides)



CFE New Higher

6.1 Alkanoic Acids

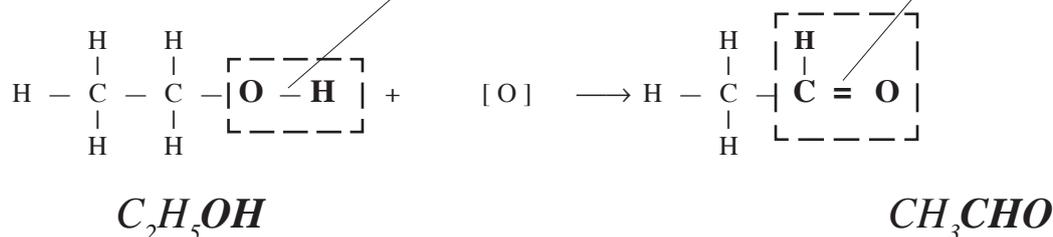
This first topic revises the names, formulae and structures of the family of acids called the alkanoic acids

Ethanoic Acid

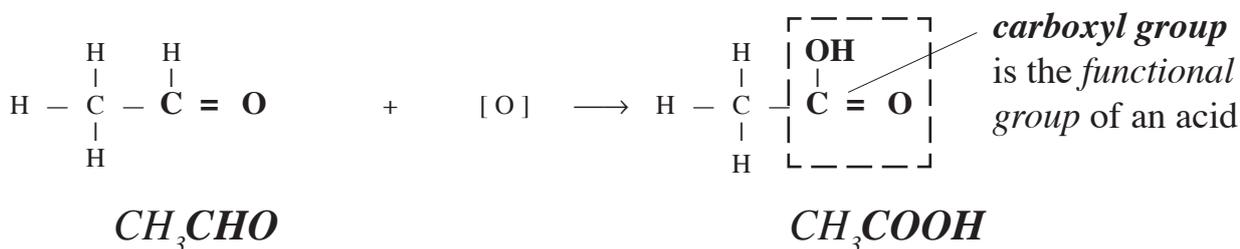
This activity considers the formation and structure of the alkanoic acid called ethanoic acid.

Ethanoic acid is normally manufactured from *ethanol*. The *oxygen* required for the *oxidation* reaction can come from the *air* or from an *oxidising agent* such as *copper(II) oxide*.

During the first *oxidation* step, the *hydroxyl group* is converted into a *carbonyl group*: *ethanol* is converted into *ethanal*



During the second *oxidation* step, an *oxygen* atom is inserted to convert the *carbonyl group* into a *carboxyl group*: *ethanal* is converted into *ethanoic acid*.

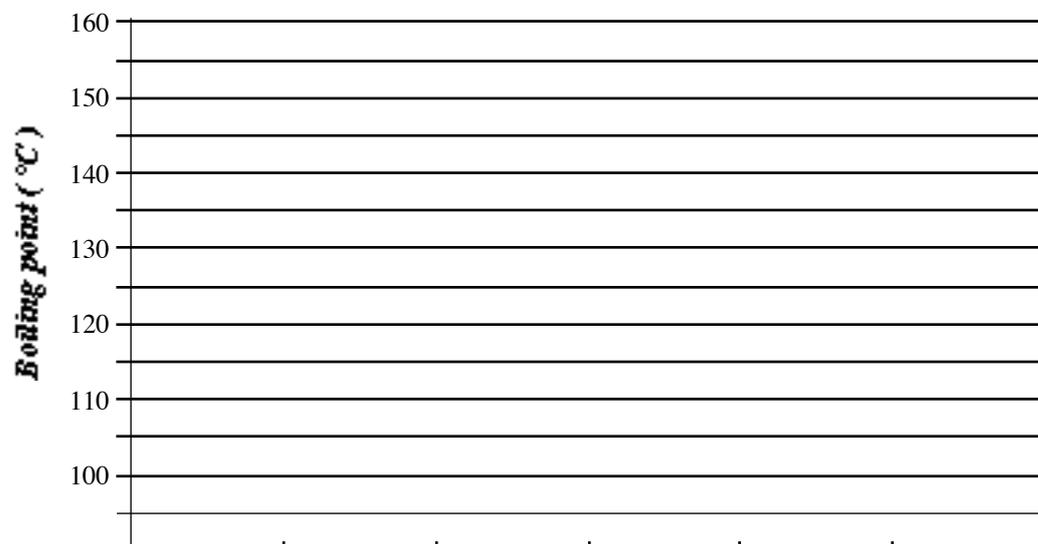


Alkanoic Acids

This activity considers the alkanoic acids as a homologous series.

a)	<i>Chemical properties</i>	<i>Ethanoic acid</i>	<i>Propanoic acid</i>
	<i>Smell & Appearance</i>	'vinegary' colourless liquid	'cheesy' colourless liquid
	<i>Universal indicator</i>	soluble pink, pH < 7	soluble pink, pH < 7
	<i>Magnesium</i>	'fizzes' (hydrogen gas)	'fizzes' (hydrogen gas)
	<i>Calcium carbonate</i>	'fizzes' (CO ₂ gas)	'fizzes' (CO ₂ gas)

b) **Boiling point trend**



c) **General formula**

The general formula for the alkanolic acids is:-



Notice that one of the **carbon** atoms is not included in the C_n 'chain'. This is to enable the **carboxyl functional group** to be emphasised. **WARNING!** - this means that for each acid n is one less than you'd expect; *methanoic* $n = 0$, *ethanoic* $n = 1$, etc.

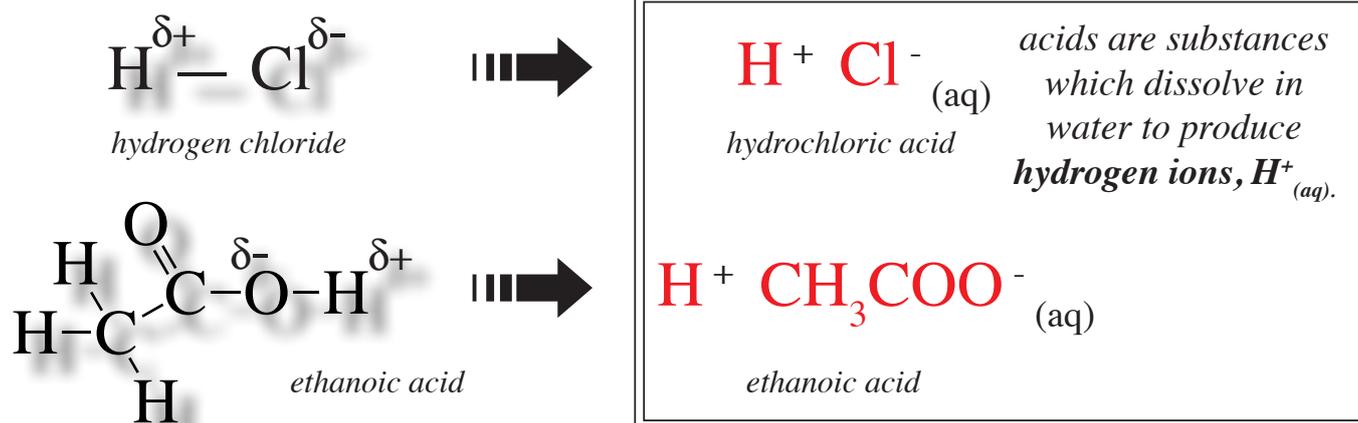
From all this it can be seen that the **alkanoic acids** have:

- ① **similar chemical** properties
- ② **physical** properties that show a **steady trend**

and ③ a **common general** formula, so they belong to a **homologous series**

d) **'Normal' Acid Reactions**

Alkanolic acids do all the 'normal' acid reactions

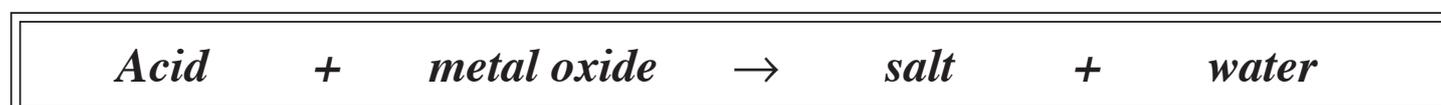
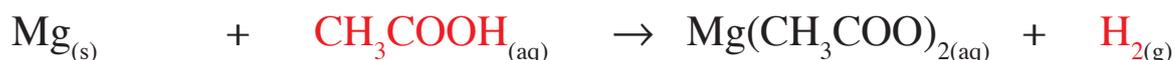




e.g. *magnesium* + *sulfuric acid* → *magnesium sulfate* + *hydrogen*



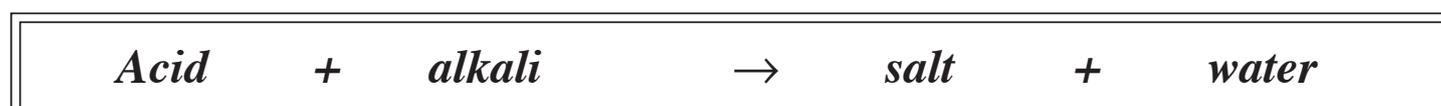
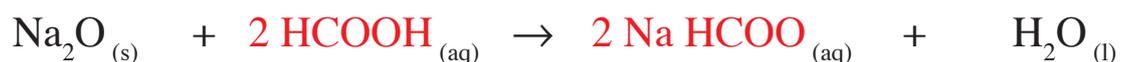
magnesium + *ethanoic acid* → *magnesium ethanoate* + *hydrogen*



e.g. *iron (III) oxide* + *nitric acid* → *iron(III) nitrate* + *water*



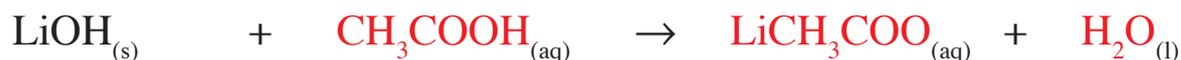
sodium oxide + *methanoic acid* → *sodium methanoate* + *water*



e.g. *potassium hydroxide* + *hydrochloric acid* → *potassium chloride* + *water*



lithium hydroxide + *ethanoic acid* → *potassium ethanoate* + *water*





e.g. calcium carbonate + hydrochloric acid \rightarrow calcium chloride + water + carbon dioxide



sodium carbonate + methanoic acid \rightarrow sodium methanoate + water + carbon dioxide



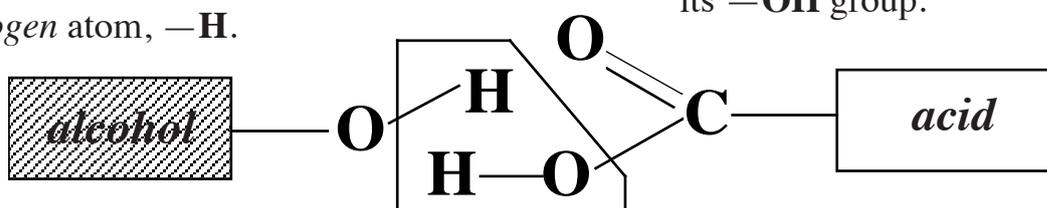
e) 'Organic' Acid Reactions *Alkanoic acids* also have reactions that are characteristic of *organic carboxylic acid* reactions

Condensation: reaction in which two molecules join together, usually in the presence of a catalyst, with elimination of water or some other simple molecule.

To *join together*, each molecule must *lose* some of the existing atoms attached to the *carbon* atom with the *functional group*.

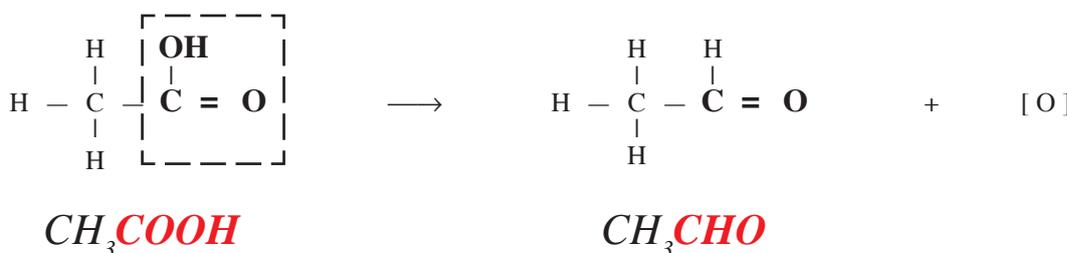
The *hydroxyl group* on the *alcohol* will have to lose its *hydrogen atom*, $-\text{H}$.

The *carboxyl group* on the *acid* loses its $-\text{OH}$ group.



The whole reaction is helped by the fact that an $-\text{OH}$ group and an $-\text{H}$ atom will then be able to form a *stable molecule*, water (H_2O).

Reduction: reaction in which the oxygen:hydrogen ratio decreases



In other words, it is possible to *reverse* the *oxidation* reactions met earlier using a *reducing agent* such as LiAlH_4 :



Structures, Names & Formulae

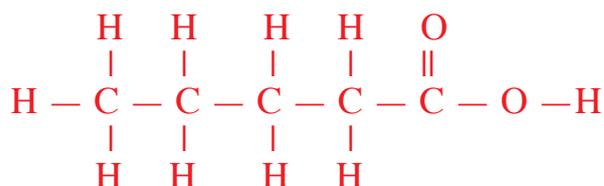
This activity considers the structures, formulae and systematic names of some of the members of the alkanolic acid family.

Any molecule that contains the *carboxyl* group, $-\text{COOH}$, can be considered as a *carboxylic acid*. The molecule could have a *chain* structure, a *ring* structure or even be *aromatic*, it could be *saturated* ($\text{C}-\text{C}$) or *unsaturated* ($\text{C}=\text{C}$ or $\text{C}\equiv\text{C}$).

The *alkanoic acids* are compounds which contain the *carboxyl* group joined to a hydrocarbon *chain* in which all the carbon atoms are joined by *single* bonds.

As usual, there are three ways to represent the formula of, for example, *pentanoic acid*

full structural
formula



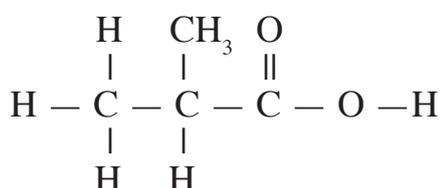
shortened structural
formula



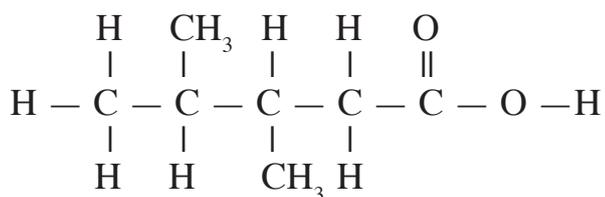
functional molecular
formula



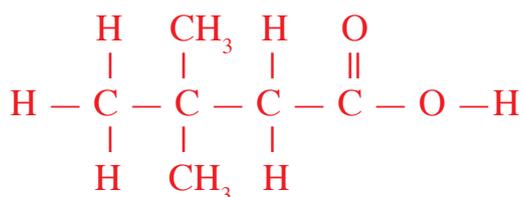
For naming purposes, the *carbon* of the *carboxyl functional* group is always taken as number 1, and the 'longest' chain always starts with the *functional* group. *Branches* must then be *numbered* accordingly. For example:



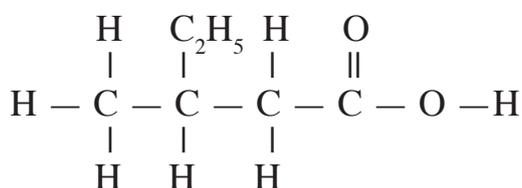
2-methylpropanoic acid



3,4-dimethylpentanoic acid



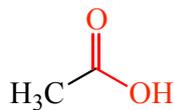
3,3-dimethylbutanoic acid



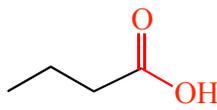
3-methylpentanoic acid

Uses of Acids

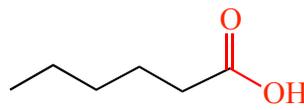
This activity looks at the variety of acids and some of their uses



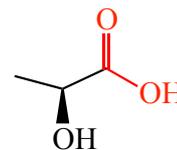
ethanoic acid
responsible for
the pungent smell
of vinegar



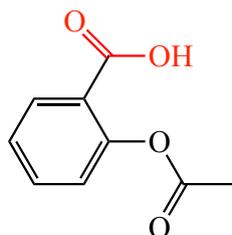
butanoic acid
responsible for
the rancid odour
of sour butter



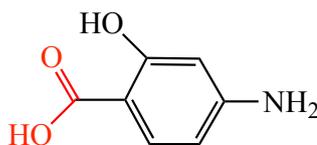
hexanoic acid
responsible for
the odour
of smelly feet



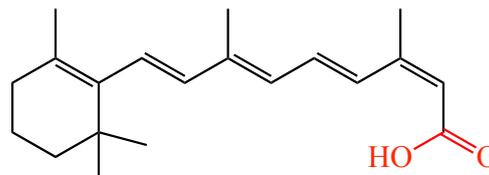
lactic acid
responsible for
the taste of
sour milk



acetylsalicylic acid
Aspirin: a widely
used analgesic



p-aminosalicylic acid
used in the treatment of
tuberculosis



isotretinoin
used in the treatment of
acne

Carbon atoms	Common name	IUPAC name	Chemical formula	Common location or use
1	Formic acid	Methanoic acid	HCOOH	Insect stings
2	Acetic acid	Ethanoic acid	CH ₃ COOH	Vinegar
3	Propionic acid	Propanoic acid	CH ₃ CH ₂ COOH	Preservative for stored grains
4	Butyric acid	Butanoic acid	CH ₃ (CH ₂) ₂ COOH	Butter
5	Valeric acid	Pentanoic acid	CH ₃ (CH ₂) ₃ COOH	Valerian
6	Caproic acid	Hexanoic acid	CH ₃ (CH ₂) ₄ COOH	Goat fat
7	Enanthic acid	Heptanoic acid	CH ₃ (CH ₂) ₅ COOH	
8	Caprylic acid	Octanoic acid	CH ₃ (CH ₂) ₆ COOH	Coconuts and breast milk
9	Pelargonic acid	Nonanoic acid	CH ₃ (CH ₂) ₇ COOH	Pelargonium
10	Capric acid	Decanoic acid	CH ₃ (CH ₂) ₈ COOH	Coconut and Palm kernel oil
11	Undecylic acid	Undecanoic acid	CH ₃ (CH ₂) ₉ COOH	
12	Lauric acid	Dodecanoic acid	CH ₃ (CH ₂) ₁₀ COOH	Coconut oil and hand wash soaps.
13	Tridecylic acid	Tridecanoic acid	CH ₃ (CH ₂) ₁₁ COOH	
14	Myristic acid	Tetradecanoic acid	CH ₃ (CH ₂) ₁₂ COOH	Nutmeg
15		Pentadecanoic acid	CH ₃ (CH ₂) ₁₃ COOH	
16	Palmitic acid	Hexadecanoic acid	CH ₃ (CH ₂) ₁₄ COOH	Palm oil
17	Margaric acid	Heptadecanoic acid	CH ₃ (CH ₂) ₁₅ COOH	
18	Stearic acid	Octadecanoic acid	CH ₃ (CH ₂) ₁₆ COOH	Chocolate, waxes, soaps, and oils
19		Nonadecylic acid	CH ₃ (CH ₂) ₁₇ COOH	Fats, vegetable oils, pheromone
20	Arachidic acid	Icosanoic acid	CH ₃ (CH ₂) ₁₈ COOH	Peanut oil

SELF CHECK

6.1

Q1. Which of the following is a word equation for the manufacture of ethanoic acid?

- A copper + ethanol \longrightarrow oxygen + ethanoic acid
- B oxygen + ethanol \longrightarrow copper + ethanoic acid
- C oxygen + ethanol \longrightarrow water + ethanoic acid
- D water + ethanol \longrightarrow oxygen + ethanoic acid.

Q2. The functional group in an ethanoic acid molecule is called the

- A methyl group
- B carboxyl group
- C carbonyl group
- D hydroxyl group.

Q3. Which result is obtained when magnesium is put in 0.1 M ethanoic and propanoic acids?

<i>ethanoic acid</i>	<i>propanoic acid</i>
----------------------	-----------------------

- | | |
|--------------|------------|
| A bubbles | bubbles |
| B no bubbles | bubbles |
| C bubbles | no bubbles |
| D no bubbles | no bubbles |

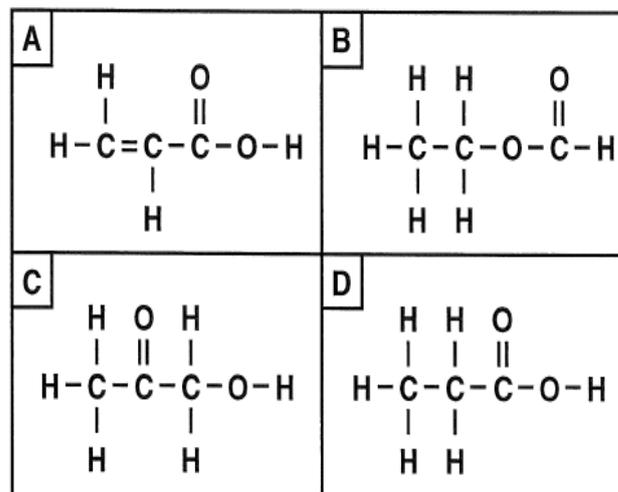
Q4. What is the general formula for an alkanolic acid?

- A $C_nH_{2n}COOH$
- B $C_nH_{2n+1}COOH$
- C $C_nH_{2n-1}COOH$
- D $C_nH_{2n+2}COOH$

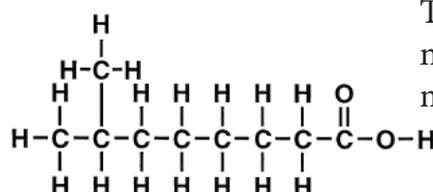
Q5. Which of the following formulae could represent hexanoic acid?

- A $C_6H_{13}COOH$
- B C_5H_9COOH
- C $C_5H_{11}COOH$
- D $C_6H_{11}COOH$

Q6. Which of the following structures represents an alkanolic acid?

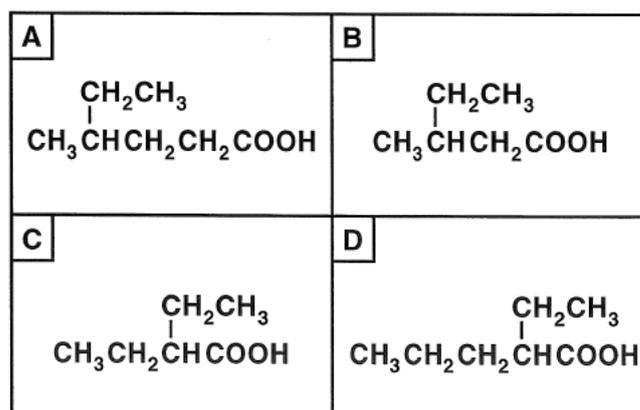


Q7. The systematic name for this molecule is



- A 2-methylheptanoic acid.
- B 2-methyloctanoic acid.
- C 6-methylheptanoic acid.
- D 7-methyloctanoic acid.

Q8. What is the shortened structural formula for 2-ethylbutanoic acid?



Q9. The conversion of butanoic acid to butanal is known as

- A condensation
- B oxidation
- C reduction
- D substitution

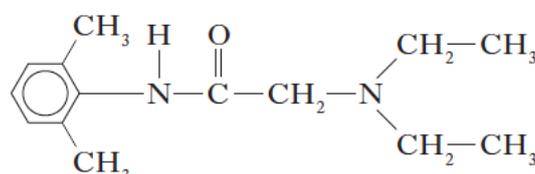
HOME PRACTICE

6.1

- Q1.** Name each of the following molecules.
- a) C_2H_5COOH 1
- b) $CH_3CH_2CH_2CH_2COOH$ 1
- Q2.** Draw the full structural formula for each of the following substances.
- a) methanoic acid 1
- b) 2-methylbutanoic acid 1
- Q3.** Draw the shortened structural formula for each of the following substances.
- a) hexanoic acid 1
- b) 4-methylpentanoic acid 1
- Q4.** All of the substances above are examples of one type of organic compound.
- a) Name the homologous series to which they belong. 1
- b) Name the wider class of organic compound to which they also belong. 1
- c) Name the functional group which characterises these compounds. 1
- d) **Explain** of the three acids - butanoic, octanoic, or hexanoic acid - would you expect to have the highest boiling point? 1

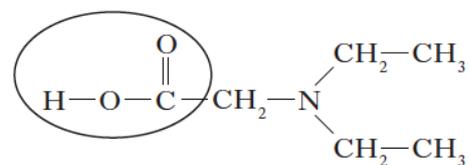
- Q5.** Dental anaesthetics are substances used to reduce discomfort during treatment.

Lidocaine is a dental anaesthetic.



Lidocaine causes numbness when applied to the gums. This effect wears off as the lidocaine is hydrolysed.

One of the products of the hydrolysis of lidocaine is compound C.



compound C

- a) Name the functional group circled above. 1
- b) Draw a structural formula for the organic compound formed when compound C reacts with $NaOH_{(aq)}$ 1
- c) Draw a structural formula for **both** organic compounds that could be formed when compound C is reduced by reacting with $NaAlH_4$. 2

Total (14)

6.2 ESTERS - Flavour Molecules

This second topic revises the group of substances known as esters and looks at their properties, their uses, and how they are related to alcohols and carboxylic acids.

Properties & Uses

This activity is about the properties and uses of esters, in particular an ester called pentyl ethanoate (amyl acetate)

Property	Result
Appearance	<i>clear, colourless liquid</i>
Smell	<i>sweet , pleasant - 'pear drops'</i>
Solubility	<i>immiscible in water - separate layers</i>
pH	<i>neutral , pH = 7 , no H⁺ or OH⁻</i>
Solvent action	<i>dissolves many non-polar & polar substances</i>

The 3 main uses of esters are as:

- ① *flavourings* - in foodstuffs
- ② *solvents* - e.g used in *nail varnish*
- and ③ *perfumes* - are *volatile*, so quickly releases *vapour*

Being *volatile* often makes them very *flammable*.

Ester Names

This activity considers the names of esters and how they relate to the alcohol and carboxylic acid from which the ester can be made

An *ester* is a substance which is formed by the *condensation* reaction of an alcohol with a carboxylic acid.

Each *ester* can thought of as having a '*parent*' *alcohol* from which it is formed. The '*parent alcohol*' provides the '*christian name*' of the *ester*. The *alcohol* name has the '-ol' ending replaced with an '-yl' ending.

'parent alcohol'	ester 'christian name'
<i>methanol</i>	<i>methyl</i>
<i>ethanol</i>	<i>ethyl</i>
<i>propanol</i>	<i>propyl</i>
<i>butanol</i>	<i>butyl</i>

Each **ester** also has a '**parent**' **carboxylic acid** from which it is formed. The '**parent acid**' provides the '**surname**' of the **ester**. The **acid** name has the '**—oic**' ending replaced with an '**—oate**' ending.

'parent acid'	ester 'surname'
methanoic	methanoate
ethanoic	ethanoate
propanoic	propanoate
butanoic	butanoate

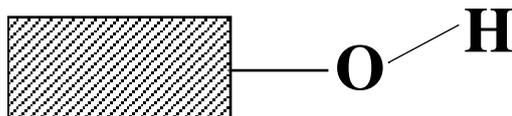
For example,



parent alcohol	parent carboxylic acid	ester name	ester flavour
ethanol	methanoic acid	ethyl methanoate	rum
ethanol	ethanoic acid	ethyl ethanoate	sweet wine
pentan-1-ol	ethanoic acid	pentyl ethanoate	pear drop
ethanol	butanoic acid	ethyl butanoate	pineapple

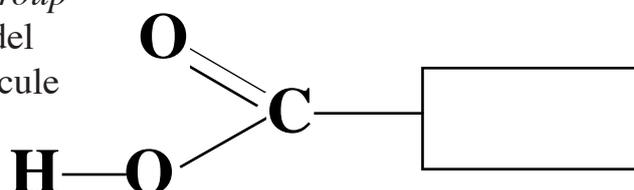
Functional Group

This activity considers the functional group in an ester molecule by looking at how it is formed from the parent alcohol and the parent carboxylic acid



Since all **alcohols** have the **hydroxyl** functional group they can be all be represented by the simple model shown. The block stands for the rest of the molecule which, in **alkanols**, would be a **hydrocarbon chain**.

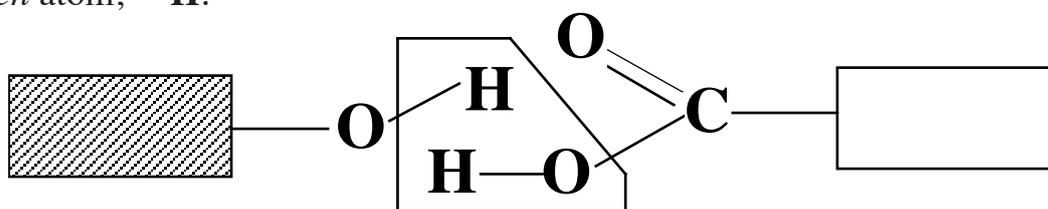
Since all **acids** have the **carboxoxyl** functional group they can be all be represented by the simple model shown. The block stands for the rest of the molecule which, in **alkanoic acids**, would be a **hydrocarbon chain**.



To **join together**, each molecule must **lose** some of the existing atoms attached to the **carbon** atom with the **functional group**.

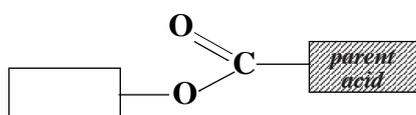
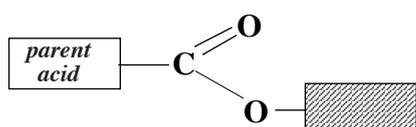
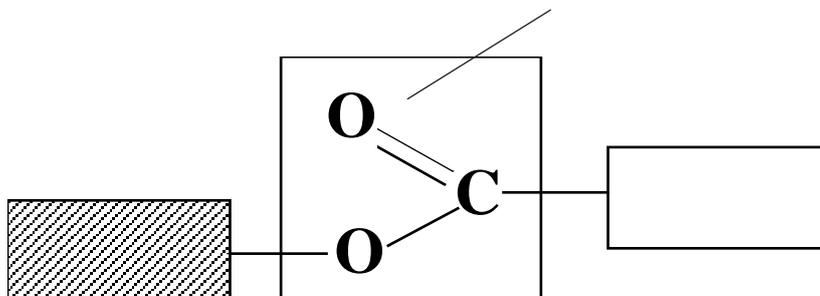
The **hydroxyl group** on the **alcohol** will have to lose its **hydrogen atom**, $-\text{H}$.

The **carboxyl group** on the **acid** loses its $-\text{OH}$ group.



The whole reaction is helped by the fact that an $-\text{OH}$ group and an $-\text{H}$ atom will then be able to form a **stable molecule**, *water* (H_2O).

The resulting **ester** molecule, formed by *joining* an **alcohol** to an **acid**, owes its *properties* to the group of atoms that now *link* the two molecules together. This can be called the **carboxylate group**, but is more often referred to as the '**ester link**'.



It is important to be able to recognise the **ester link** no matter how it is drawn.

You will also be expected to be able to redraw the **parent acid** and **parent alcohol** molecules, so whichever **carbon chain** is directly attached to the **carbonyl group**, $\text{C}=\text{O}$, must have been the **parent acid** and the other chain belonged to the **alcohol**.

SELF CHECK

6.2

- Q1.** Esters are substances which
- A often have poisonous vapours
 - B are very soluble in water
 - C often have pleasant tastes and smells
 - D are liquids with acidic properties.

- Q2.** Which of the following is **not** a use for pentyl ethanoate ?

- A colouring in food dyes
- B scent in perfume
- C flavouring in sweets
- D solvent in nail varnish.

- Q3.** An ester can be made from

- A a carboxylic acid and an alkane
- B an alcohol and a carboxylic acid
- C an alkane and an alcohol
- D a carboxylic acid and an alkene.

- Q4.** Which of the following substances is an example of an ester?

- A 2-methylbutane
- B methylpropanoic acid
- C methylpropanoate
- D 3-methylbutan-1-ol

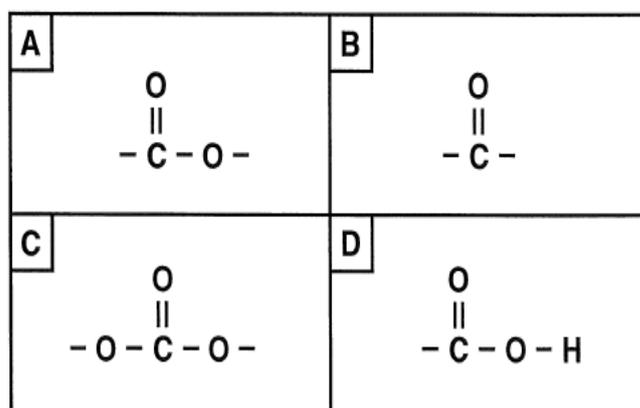
- Q5.** What is the parent alkanolic acid for pentyl ethanoate?

- A heptanoic acid
- B pentanoic acid
- C methanoic acid
- D ethanoic acid

- Q6.** What is the parent alkanol for pentyl ethanoate?

- A heptanol
- B pentanol
- C methanol
- D ethanol

- Q7.** What is the structure of the functional group in an ester molecule?



- Q8.** The functional group in an ester is called the

- A carbonate group
- B carboxyl group
- C carbonyl group
- D carboxylate group.

- Q9.** An ester has the following structural formula



The name of this ester is

- A propyl propanoate
- B ethyl butanoate
- C butyl ethanoate
- D ethyl propanoate.

HOME PRACTICE

6.2

Q1. Name the ester formed from each of the following pairs of compounds.

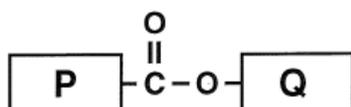
- a)** ethanol and butanoic acid 1
b) propanoic acid and butanol 1

Q2. Identify three typical uses for esters from the selection given below.

fertilisers	flavourings
pigments	perfumes
explosives	solvents
vitamins	neutralisers

3

Q3. The diagram shows a simplified model of an ester molecule.

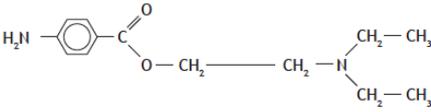
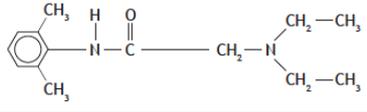
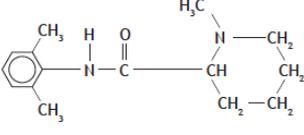
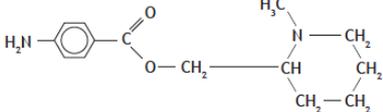


- a)** Name the functional group in this molecule. 1
b) Which part of the molecule, P or Q, came from the parent acid? 1

Q4. Name **i)** the parent acid and **ii)** the parent alcohol used to make each of the following esters.

- a)** propyl hexanoate 2
b) methyl octanoate 2

Q5. The table below shows the duration of numbness for common anaesthetics.

Name of anaesthetic	Structure	Duration of numbness (minutes)
procaine		7
lidocaine		96
mepivacaine		114
anaesthetic X		

- a)** Which of the anaesthetics are esters? 1
b) Which of the anaesthetics have amide links? 1
c) Estimate the duration of numbness, in minutes, for anaesthetic X. 1

Total (14)

6.3 ESTERS - Condensation & Hydrolysis Reactions

This topic looks in more detail at how esters can be made from their parent compounds and how esters can then be broken down to produce the original alcohol and carboxylic acid molecules.

Formation Reaction

This activity considers the structural formulae of the molecules involved in the formation of an ester by the condensation reaction between methanol and ethanoic acid

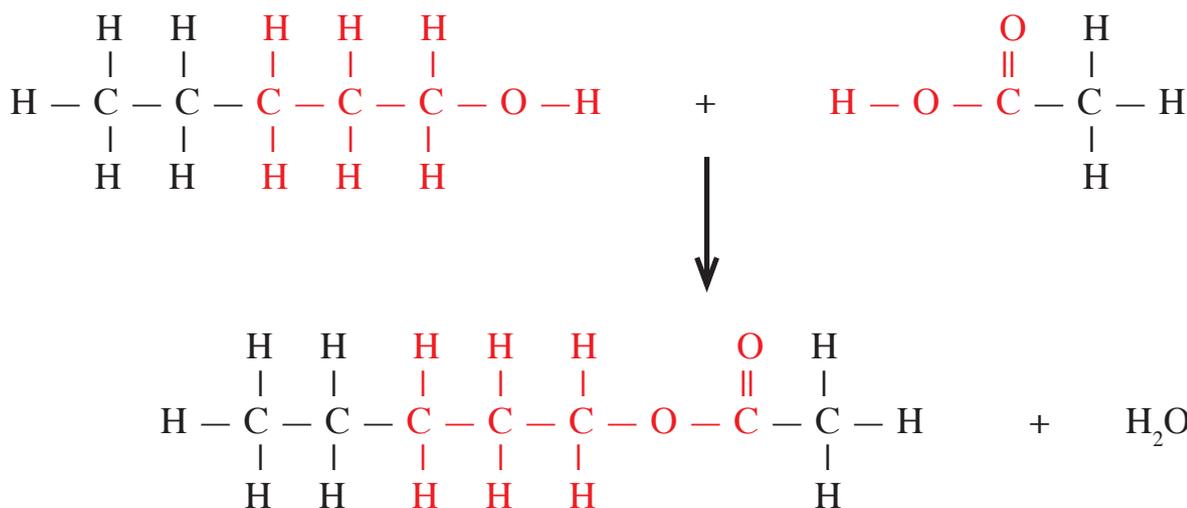
A **condensation reaction** occurs when **two molecules** each lose one or more atoms in order to **join together**. Another small molecule is also formed by the 'lost' atoms.

The other small molecule formed is often **water**, hence the use of the name **condensation**. However, other reactions that 'form water', such as **neutralisation** or the **dehydration of alkanols** to form alkenes, are **not** condensations because they **do not** result in the **joining together** of two molecules.

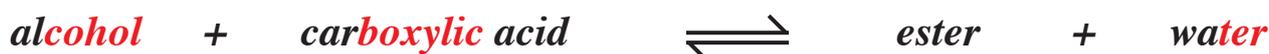
Word equation:



Equation using full structural formulae:



A **general word equation** can be written for the **formation** of an **ester**.



Condensation

The aim of this activity is to prepare an ester using the condensation reaction and to confirm the formation of an ester by its smell

conc.



The reagent needed for the **condensation** reaction is **concentrated** H_2SO_4 . It has two functions:

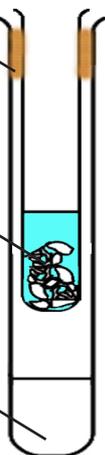
Catalyst: it provides H^+ (aq) which help **form** the **ester link** but are then **reformed**.

Dehydrating agent: it **absorbs** water molecules which **slows** down the **reverse** reaction and helps push the **equilibrium mixture** to the **right**.

wet **paper** towel

small test-tube with
ice and **water**

equal mixture of
alcohol and
carboxylic acid



Water Bath: A **beaker** of water is heated until **boiling** and then the **bunsen burner** is put out - both **alcohols** and **esters** are very **volatile / flammable** - **Safety**

Cold Finger: To **prevent** loss of **volatile** chemicals during **heating**.

conc H₂SO₄ : 5 drops of **conc.** H₂SO₄ were added by the **teacher** - **Safety**

By this stage, **two layers** may have formed. The **very polar alcohol** and **carboxylic acid** in one **layer** and the **less polar ester** floating above.

Adding **water** (with **hydrogen bonding**) will **dissolve** the **alcohol** and the **acid** as they also have **hydrogen bonding** groups. The **ester** only has **polar:polar (dipole:dipole) intermolecular** bonding.

The smell of the **ethanoic acid** (**vinegar**) is overpowering, so Na₂CO₃ solution was used instead - the acid is converted to **ethanoate salt** and **fizzing** can be used to tell when enough has been added.

By this stage, **two definite layers** will exist and the characteristic **smell** of the **ester** (top layer) will be obvious.

Alternatively, a **separating funnel** can be used - the **more dense water layer** (with **unreacted alcohol** and **acid**) can be let out of the **bottom** and then the **less dense ester** (top layer) is poured out of the **top**.

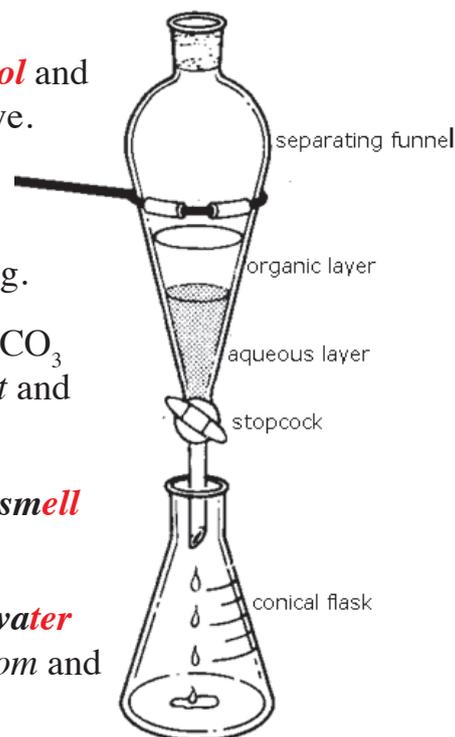
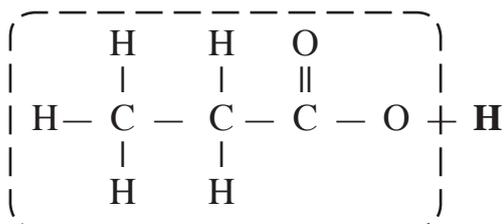


Table of esters and their smells

	methyl 1 carbon	ethyl 2 carbons	propyl 3 carbons	2-methyl propyl-	butyl 4 carbons	pentyl 5 carbons	hexyl 6 carbons	benzyl benzene ring	heptyl 7 carbons	octyl 8 carbons	nonyl 9 carbons
methanoate 1 carbon	ETHEREAL			ETHEREAL			"GREEN" 				?
ethanoate 2 carbons								JASMINE 			
propanoate 3 carbons											?
2-methyl propanoate 4 carbons, branched		ETHEREAL									?
butanoate 4 carbons											?

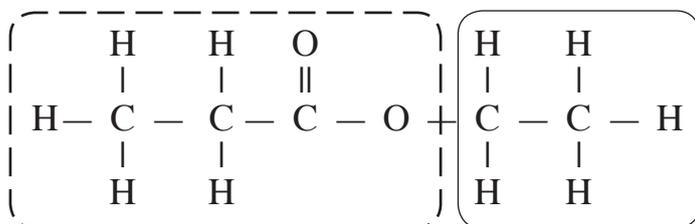
Formulae & Names

This activity deals with naming and drawing full and structural formulae for esters.



The 'best' way to think about an *ester* is to consider it as an *acid molecule* which has had its *hydrogen atom* replaced by a *carbon chain* (an *alkyl group*).

Learn to draw acids and you should find esters easy!



ethyl propanoate

As is often the case, we 'start' at the end of the name. *Identify the acid* (look for the *carbonyl* C = O) and give the *ester* its *surname* by changing the *-oic* ending to *-oate*.

The *carbon chain* (derived from the parent *alcohol*) is the '*christian*' name, *-ol* changed to *-yl*.

Ester Structures & Names

Name:	methylheptanoate
Full Structural Formula:	$\begin{array}{ccccccccccc} & \text{H} & \text{H} & \text{H} & \text{H} & \text{H} & \text{H} & \text{O} & & \text{H} & \\ & & & & & & & & & & \\ \text{H} & - \text{C} & - \text{O} & - & \text{C} & - \text{H} \\ & & & & & & & & & & \\ & \text{H} & \text{H} & \text{H} & \text{H} & \text{H} & \text{H} & & & \text{H} & \end{array}$
Shortened Structural Formula:	CH₃ CH₂ CH₂ CH₂ CH₂ CH₂ COO CH₃
Name:	
Full Structural Formula:	$\begin{array}{ccccccc} & \text{H} & \text{H} & \text{H} & \text{O} & & \text{H} \\ & & & & & & \\ \text{H} & - \text{C} & - \text{C} & - \text{C} & - \text{C} & - \text{O} & - \text{C} & - \text{H} \\ & & & & & & \\ & \text{H} & \text{H} & \text{H} & & & \text{H} \end{array}$
Shortened Structural Formula:	CH₃ CH₂ CH₂ COO CH₃
Name:	pentyl propanoate
Full Structural Formula:	$\begin{array}{ccccccccccc} & \text{H} & \text{H} & \text{H} & \text{H} & \text{H} & & \text{O} & \text{H} & \text{H} & \\ & & & & & & & & & & \\ \text{H} & - \text{C} & - \text{O} & - \text{C} & - \text{C} & - \text{C} & - \text{H} \\ & & & & & & & & & & \\ & \text{H} & \text{H} & \text{H} & \text{H} & \text{H} & & & \text{H} & \text{H} & \end{array}$
Shortened Structural Formula:	CH₃ CH₂ CH₂ CH₂ CH₂ OOC CH₂ CH₃

Hydrolysis

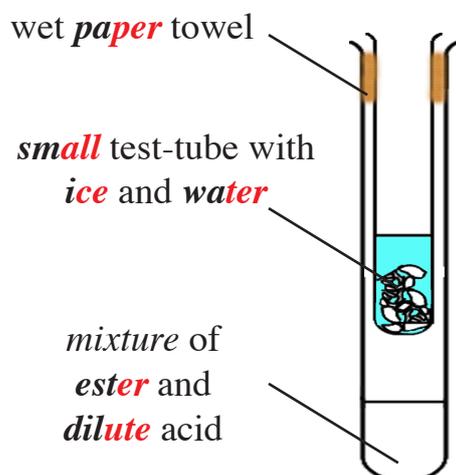
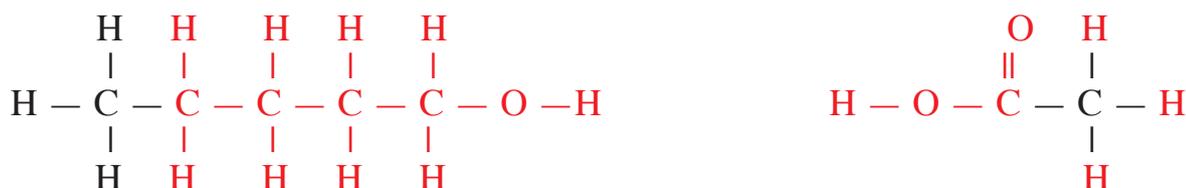
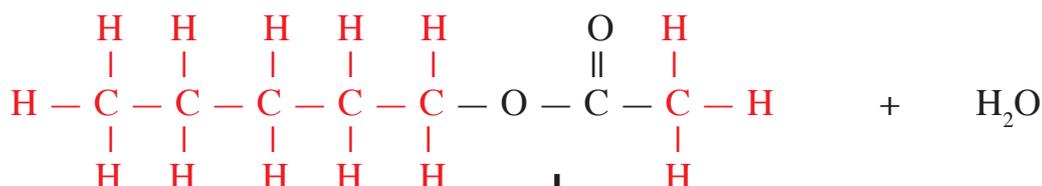
The aim of this activity is to consider the hydrolysis of an ester to produce an alcohol and carboxylic acid.



The reagent needed for the *hydrolysis* reaction is *dilute* H_2SO_4 . Again, it has two functions:

Catalyst: it provides $H^+_{(aq)}$ which help *break* the *ester link* but are then *reformed*.

Reactant: it *provides* the *water* molecules needed which *speeds* up the *forward* reaction and helps push the *equilibrium* mixture to the *right*.



Water Bath: A *beaker* of water is heated until *boiling* and then the *bunsen burner* is put out - both *alcohols* and *esters* are very *volatile* / *flammable* - *Safety*

Cold Finger: To *prevent* loss of *volatile* chemicals during *heating*.

dil. H₂SO₄ : to *provide water* needed for *hydrolysis* and $H^+_{(aq)}$ to *catalyse* the reaction.

At the beginning, *two layers* will have formed. The very *polar water* in one *layer* with the *less polar ester* floating above.

Gradually, the *ester layer* will get *smaller* and the *smell* of the *ethanoic acid* (*vinegar*) may become strong enough to be *detected* over the *smell* of the *ester* - *pear drops*.

A more reliable way of telling that the *hydrolysis* has been successful would be to add a few drops of *universal indicator* - it will turn *red*.

SELF CHECK

6.3

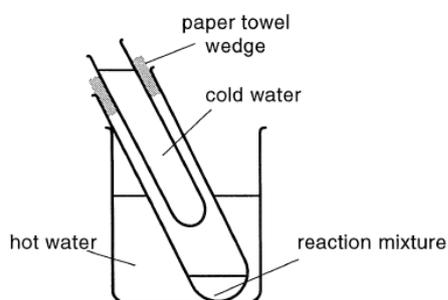
Q1. What is the general word equation for the formation of an ester?

- A ALCOHOL + CARBOXYLIC ACID
→ ESTER
- B ALCOHOL + CARBOXYLIC ACID
→ ESTER + WATER
- C SULPHURIC ACID + CARBOXYLIC ACID
→ ESTER + WATER
- D ALCOHOL + CARBOXYLIC ACID
→ ESTER + HYDROGEN

Q2. The formation of an ester is an example of

- A an addition reaction
- B a precipitation reaction
- C an oxidation reaction
- D a condensation reaction

Questions 3 and 4 refer to this diagram showing the preparation of an ester



Q3. The test-tube with cold water is there to act as a

- A catalyst
- B condenser
- C evaporator
- D purifier

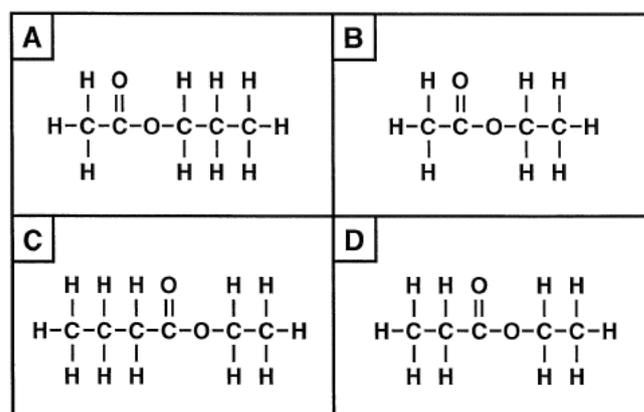
Q4. What substance is added to the reaction mixture to speed up the formation of ester?

- A concentrated nitric acid
- B deionised water
- C dilute hydrochloric acid
- D concentrated sulphuric acid

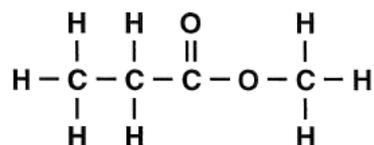
Q5. Which of the following is not true of the two molecules in a condensation reaction?

- A The two reactant molecules form a larger product molecule.
- B A small molecule is formed from atoms taken from two functional groups.
- C The reactant molecules add together to form a single product.
- D Each reactant molecule must have a reactive functional group

Q6. Which ester is formed by the reaction of CH_3COOH with $\text{CH}_3\text{CH}_2\text{CH}_2\text{OH}$?



Q7. What is the name of the following ester?



- A propyl methanoate
- B methyl ethanoate
- C methyl propanoate
- D ethyl methanoate

Q8. What is the formula of pentyl butanoate?

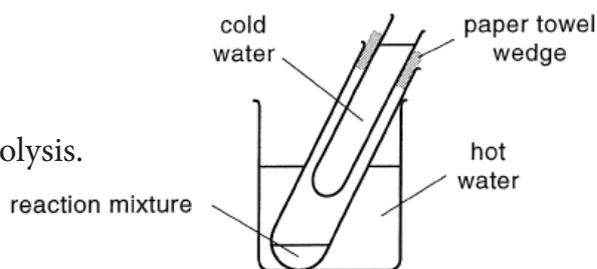
- A $\text{C}_3\text{H}_7\text{COOC}_5\text{H}_{11}$
- B $\text{C}_4\text{H}_9\text{COOC}_4\text{H}_9$
- C $\text{C}_3\text{H}_7\text{COOC}_5\text{H}_9$
- D $\text{C}_5\text{H}_{11}\text{COOC}_3\text{H}_7$

HOME PRACTICE

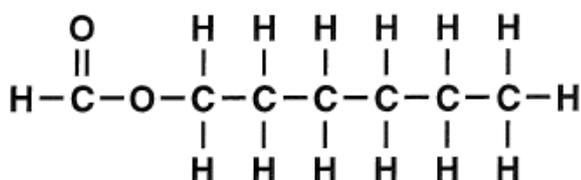
6.3

Q1. Butyl ethanoate can be hydrolysed using the apparatus shown

- What is the purpose of the cold tube? 1
- Write the word equation for this hydrolysis. 1
- Write the formula equation using shortened structural formulae. 2
- What does the term 'hydrolysis' mean? 1
- How does the hydrolysis equation compare with the equation for the formation of butyl ethanoate? 1



Q2. The diagram shows the structural formula of an ester.



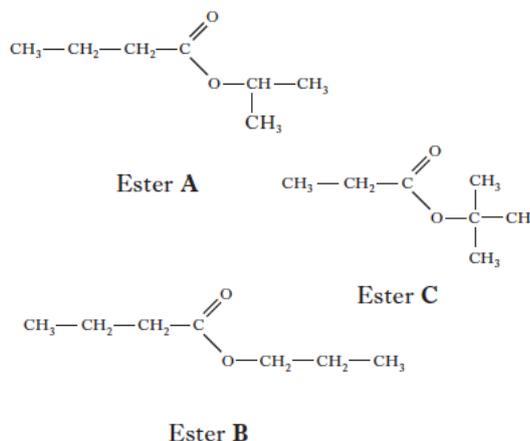
- Why can the hydrolysis of this ester be described as a *reversible reaction*? 1
- What *sign* is used in an equation to show that it is reversible? 1
- Name* the acid produced by the hydrolysis of this ester. 1
- Draw the *shortened* structural formula for the alcohol produced. 1

Q3. A team of chemists are developing a fragrance for use in a shower gel for men.

To give the gel a fruity smell the chemists are considering adding an ester. They synthesise six isomeric esters. Volunteers smell each ester and give it a rating out of one hundred depending on how fruity the smell is.

Structure	Fruit-smell rating	Structure	Fruit-smell rating
	100		92
	34		44
	0		32

a) Name the ester with the fruit-smell rating of 92. 1



b) Arrange the three esters (A , B and C) in order of *decreasing* fruit-smell rating. 1

Total (12)

6.4 Edible Fats & Oils

This lesson topic introduces edible fats and oils, substances which are both examples of esters and are closely related to each other

'Oil' is a much used word in Chemistry and can refer to quite different molecules from different parts of Chemistry.

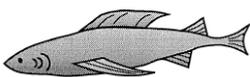
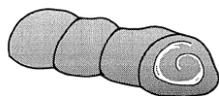
mineral oils:	mainly hydrocarbon molecules from crude oil
edible oils:	esters based on glycerol made by plants & animals
essential oils:	volatile aroma molecules that are characteristic of a particular plant

Sources of Fats & Oils

This activity considers the three main sources of edible fats and oils - animal, vegetable and marine.

Both edible **fats** and edible **oils** are esters - formed by **joining carboxylic acids** to **alcohols**. Both **fats** and **oils** are **greasy** to touch, but **fats are solid** while **oils are liquid** at **room temperature**.

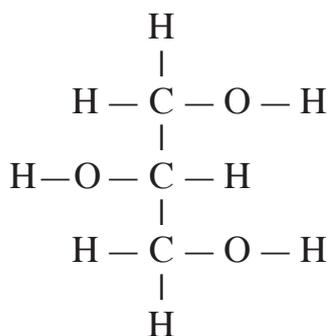
They are found in a variety of **living** things and form an important part of the **human diet**. They are usually **classified** according to whether their **source** is **animal**, **vegetable** or **marine**.



Types of Edible Fats & Oils (Source)		
Animal	Vegetable	Marine
dripping (beef)	fruits (olives)	whale blubber
lard (pork)	seeds (corn)	seal blubber
tallow (lamb)	seeds (peanuts)	sardine oil
fat (goose)	seeds (soya)	cod liver oil

Ester Molecules

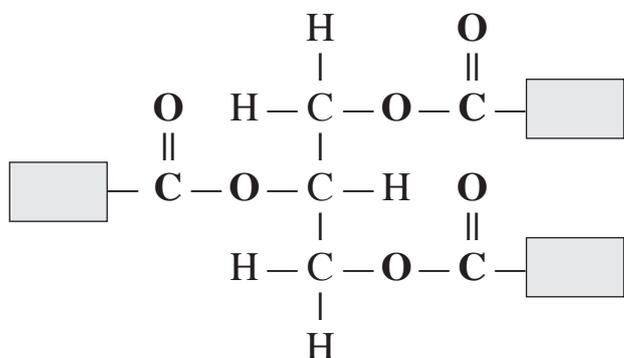
This activity looks at the structures of fats and oils as esters of the same alcohol.



Both **fats** and **oils** are esters - formed by **joining carboxylic acids** to **alcohols**.

Surprisingly, all **fat** and **oil** molecules are formed from the **same alcohol**. It has 3 carbon atoms each of which has a **hydroxyl group** (—OH) attached - it is the **triol** called **propan-1,2,3-triol**

It is a clear, colourless but very **viscous** liquid (very strong **hydrogen bonding** between molecules) and is better known as **glycerol**.



Having 3 **hydroxyl groups** allows 3 **acid** molecules to join onto **glycerol** - forming 3 **ester links** and fats & oils can be described as **triple esters** or as **triglycerides**.

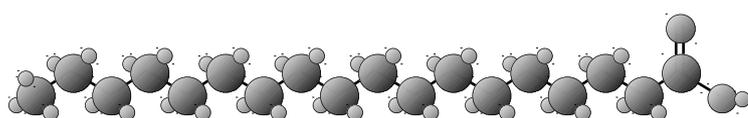
When **hydrolysed**, fats & oils always produce **3 moles** of acid molecules to **1 mole** of **glycerol**.

Different fats & oils produce different **acids** when **hydrolysed**. These acids are called **fatty acids** and differences in the **structures** of these **acid** molecules account for the different **properties** of fats and oils.

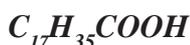
Fatty Acids

This activity looks at the structures of the carboxylic acids, i.e. the fatty acids, obtained by hydrolysing fats and oils

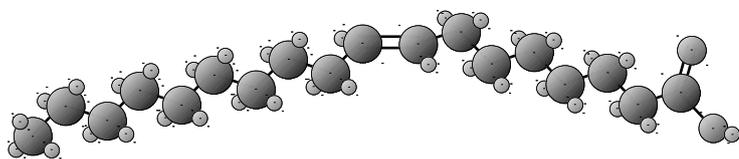
Fatty acids are saturated or unsaturated carboxylic acids, usually with long carbon chains, which are obtained from the hydrolysis of fats and oils



stearic acid



Stearic acid is a typical **saturated fatty acid** and is found in **animal fat**. All along its **carbon chain** are found C—C **single bonds**.



oleic acid

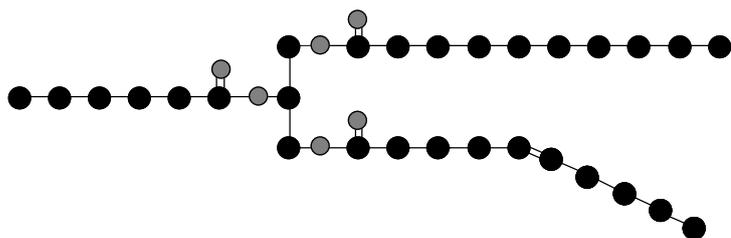


Oleic acid is a typical **unsaturated fatty acid** and is found in **olive oil**. Somewhere along its **carbon chain** is found a C = C **double bond**.

Fatty acids are quite **long chained carboxylic acids**, typically **18** carbons per molecule. **Animal fats** tend to have mainly **saturated fatty acids** while **marine oils** and **vegetable oils** usually have some **unsaturated fatty acids**.

Glycerides & Their Fatty Acid Content

This activity considers glycerides and the effect of their parent fatty acids on their properties.



Fatty acids with odd numbers of carbon atoms are rare in nature - they are usually in the range C_2 to C_{24} .

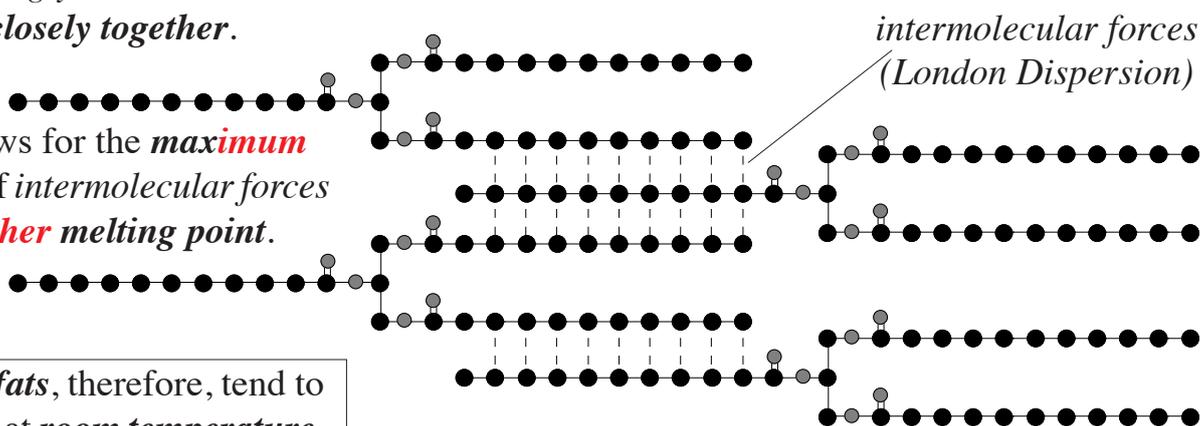
Fats and **oils** are **triple esters** based on the **alcohol glycerol**. They are often called **glycerides** or even **triglycerides**.

The 3 **acids** linked to the central **glycerol** molecule can be **identical** or **different**, **saturated** or **unsaturated**. This explains the variety of fats & oils that exist and their **different properties**.

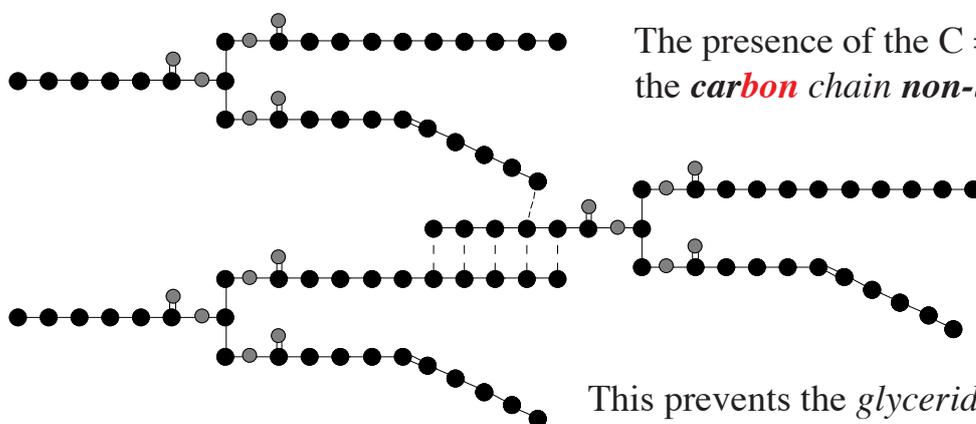
In **animal fats** most of the **fatty acids** are **saturated**. This makes their **carbon chains linear**. As a result, the **glyceride** molecules are able to **pack closely together**.

This allows for the **maximum** amount of **intermolecular forces** and a **higher melting point**.

Animal fats, therefore, tend to be **solids** at **room temperature**.



Vegetable oils and **marine oils** contain more **unsaturated fatty acids** in their **glyceride** molecules.



The presence of the **C = C double bond** makes the **carbon chain non-linear** or 'bent'.

This prevents the **glyceride** molecules **packing** so close together, so **fewer intermolecular forces** can be established, and the **melting point** is **lower**.

Vegetable oils and **marine oils** tend to be **liquids** at **room temperature**.

<i>Sample</i>	<i>Melting Point</i>	<i>Drops of bromine solution decolourised</i>
<i>lard</i>	<i>'high'</i>	<i>10</i>
<i>margarine</i>	<i>medium</i>	<i>15</i>
<i>corn oil</i>	<i>low</i>	<i>20</i>

The more drops of **bromine** that can be **decolourised**, the more **C = C double bonds** the molecules must contain, the **more unsaturated** they must be.

Conclusions:

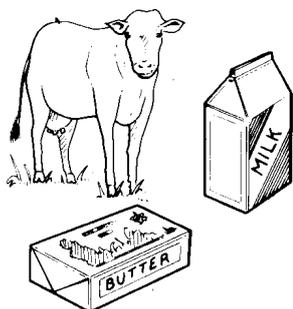
*fats are more **saturated** than oils and have **higher** melting points*

*oils are more **unsaturated** than fats and have **lower** melting points*

Hardening Oils

This activity deals with the way in which oils can be converted into solids by hardening

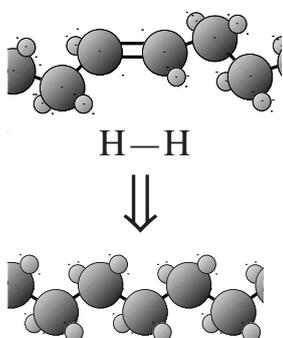
Hardening a fat or oil means hydrogenating it to increase its melting point.



Foods that are high in **animal fats**, such as **milk** and **butter**, pose **health risks** because of the **saturated fatty acids** they contain.

Vegetable oils are considered **healthier** because of the **higher level of unsaturated acids** they contain. In many cases, e.g. **frying food**, oils can do the same job that fats such as **lard** and **butter** used to do.

Vegetable oils are **runny liquids** and unsuitable for **spreading** on bread. However, if some of the **unsaturated acids** are converted into **saturated acids** by reacting with **hydrogen**, then the **melting point** of the oil will be **increased** and it will be **more solid** at **room temperature**.



This is the same **addition reaction** met earlier in the course that can be used to **convert an alkane** into an **alkene** and requires the same **catalyst, nickel**

As more and more of the **unsaturated acids** are **converted**, the **margarine** becomes **more and more solid**. However, it must not be allowed to become **too saturated** or it will be **too solid** and will lose its ability to 'spread straight from the fridge'.

Even more importantly, if allowed to become **too saturated** the **health advantages** that **unsaturated margarine** enjoys over **saturated butter** will be lost. For both these reasons only **partial hydrogenation** takes place

Health and Diet

This activity looks at some of the benefits and some of the problems associated with fats and oils



We all need some fat in our diet. But too much of a particular kind of fat – **saturated fat** – can raise our **cholesterol**, which **increases** the risk of **heart disease**. It's important to cut down on fat and choose foods that contain **unsaturated fat**.

Fats & Oils are **high energy** foods - **twice** as much as **carbohydrate** - and food high in fats & oils would be appropriate for people doing **very active** jobs, but for most people will be **stored** and can lead to **obesity** and the **health problems** associated with **obesity**.

The four **fat-soluble vitamins** namely **vitamin A, D, E and K** are, in fact, require fats and oils in the food to be **absorbed** through the gut. Inadequate fats may result in the **deficiency** of these **vitamins** leading to serious **health problems**. By law, **vitamins** are added to **margarine**.

Eating **unsaturated** fats instead of **saturated** can help lower blood cholesterol. **Unsaturated** fat, such as **omega-3 essential fatty acids**, is found in: **oily fish** such as **salmon** and **sardines**.

SELF CHECK

6.4

- Q1.** Which of the following is not a source of fats and oils for direct consumption?
- A animal
 - B mineral
 - C vegetable
 - D marine
- Q2.** Which of the following will provide a fat with a high melting point?
- A olives
 - B lamb
 - C sunflower seed
 - D sardines
- Q3.** Fats and oils are both examples of
- A carboxylic acids
 - B hydrocarbons
 - C alcohols
 - D esters
- Q4.** The most unsaturated fats or oils are those which decolourise
- A bromine solution the most quickly
 - B the least bromine solution
 - C bromine solution the most slowly
 - D the most bromine solution
- Q5.** When comparing a fat and an oil of equal molecular size
- A the more saturated one will have the higher melting point
 - B the more unsaturated one will have the higher melting point
 - C the less saturated one will have the higher melting point
 - D the two will have exactly the same melting point
- Q6.** In comparison to oils, fats are generally
- A more unsaturated
 - B equally unsaturated
 - C more saturated
 - D equally saturated
- Q7.** Compared with carbohydrates fats provide
- A slightly less energy per gram
 - B slightly more energy per gram
 - C more than double the energy per gram
 - D less than half the energy per gram
- Q8.** Compared with vegetable oils, butter in the human diet can
- A provide important vitamins
 - B cause less heart disease
 - C keep the arteries clear
 - D cause less obesity
- Q9.** Fats have higher melting points than oils because comparing fats and oils
- A fats have more hydrogen bonds
 - B fat molecules are more saturated
 - C fat molecules are more loosely packed
 - D fats have more cross-links between molecules.
- Q10.** Oils are generally
- A solid at room temperature and contain a high proportion of unsaturated molecules
 - B solid at room temperature and contain a high proportion of saturated molecules
 - C liquid at room temperature and contain a high proportion of unsaturated molecules
 - D liquid at room temperature and contain a high proportion of saturated molecules

SELF CHECK

6.4

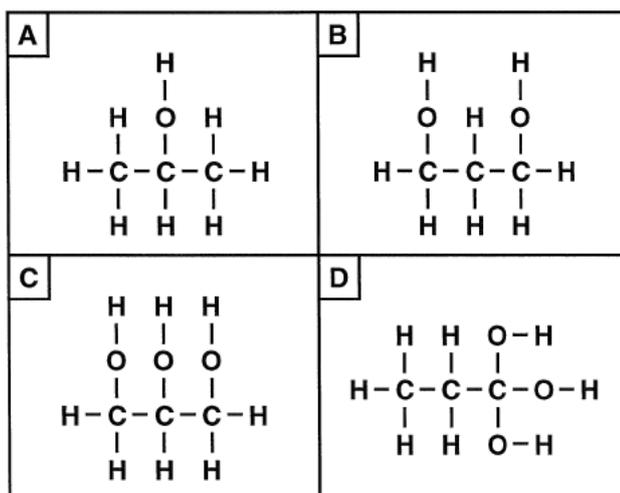
Q11. Which statement is *not* true of fats?

- A the parent acid molecule has three carboxyl groups.
- B there are three fatty acid chains in the fat molecule.
- C the parent alcohol molecule has three hydroxyl groups.
- D there are three ester links in the fat molecule

Q12. Hydrolysis of an oil is likely to produce

- A methanol
- B glycerol
- C ethanoic acid
- D methanoic acid

Q13. What is the structural formula of glycerol?



Q14. What is the ratio of alcohol to acid in the products from a hydrolysed fat?

- A 3 moles acid to 2 moles alcohol
- B 3 moles acid to 1 moles alcohol
- C 1 moles acid to 2 moles alcohol
- D 1 moles acid to 3 moles alcohol

Q15. Which of the following is a saturated fatty acid?

- A $C_{18}H_{33}COOH$
- B $C_{18}H_{35}COOH$
- C $C_{17}H_{33}COOH$
- D $C_{17}H_{35}COOH$

Q16. The conversion of an oil into a fat involves the removal of

- A single bonds
- B double bonds
- C hydrogen atoms
- D hydroxyl groups

Q17. Hardening of a fat or oil is done by

- A hydrolysis
- B dehydration
- C catalytic hydrogenation
- D fractional distillation

Q18. In the manufacture of margarine, why is only partial hydrogenation carried out?

- A to retain as much unsaturation as possible
- B to avoid destroying natural vitamins
- C to avoid losing the natural flavours
- D to keep the molecular size as large as possible

Q19. Which line in the table shows correct functional groups for aldehydes and ketones and fats and oils?

	Aldehydes and ketones	Fats and oils
A	carbonyl	hydroxyl
B	carboxyl	hydroxyl
C	carboxyl	ester link
D	carbonyl	ester link

HOME PRACTICE

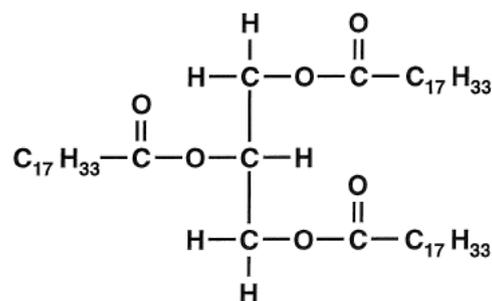
6.4

- Q1.** Olive oil and bacon fat are chemically related.
- To which class of organic compound do both belong? 1
 - Which of these two substances has the lower melting point? 1
 - Which of them is likely to be more unsaturated? 1
 - What test could you do to compare the degree of unsaturation in these two substances and what result would you expect? 2

- Q2.** Carbohydrates, fats and oils are all regarded as energy foods.
- How would you expect equal masses of olive oil and sugar to compare in energy content? 1
 - In what way can some fatty foods claim to be healthier than oils in the human diet? 1
 - In what way can unsaturated fats claim to be healthier than saturated fats in the human diet? 1
 - Explain what obesity is and how eating fats and oils can cause this. 1

- Q3.** Animal fats and vegetable oils are examples of esters.
- How many ester links are present in a single fat molecule? 1
 - Name the alcohol which is produced by the hydrolysis of any animal fat. 1
 - Draw the full structural formula for this alcohol. 1

- Q4.** The diagram shows a structural formula of an ester oil.



- Write the functional molecular formula of the parent fatty acid. 1
- Is this particular fatty acid saturated or unsaturated? Explain your answer. 2
- What will be the ratio of acid molecules to alcohol molecules in the products of hydrolysis of this oil? 1
- What type of reaction can be used to 'harden' this oil? 1
- What effect does hardening have on
 - the degree of saturation in the molecule,
 - and the melting point? 2

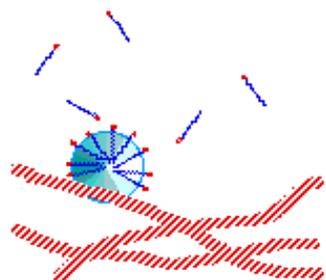
Total (19)

6.5 Soaps, Detergents & Emulsions

This lesson topic introduces other aspects of fats & oils including derivatives and special mixtures.

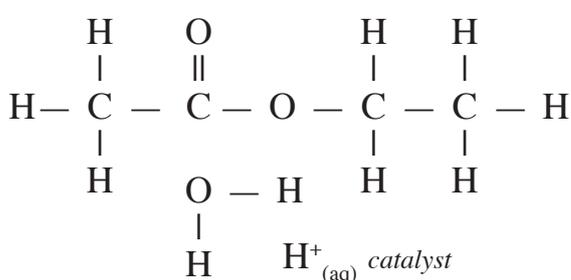
Soaps & Cleansing Action

This activity considers how soaps are made and how the structure of the soap molecule allows it to act as a cleansing agent



Soaps are molecules that help **water** cope with awkward 'greasy' molecules, such as **fats & oils**, that would be difficult for **water** alone to wash away.

The names of some **soaps**, e.g. *Palmolive*, betray the fact that **soaps** themselves are made from **fats** and **oils** - *palm oil* and *olive oil*.

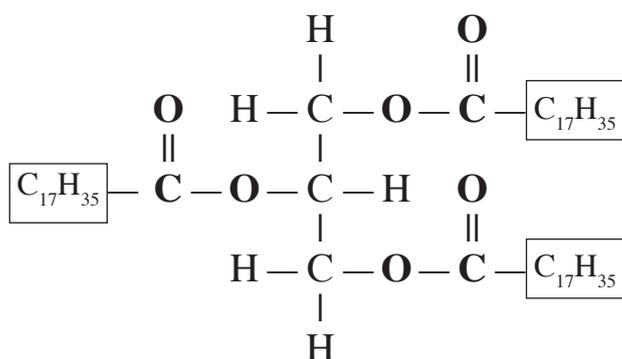
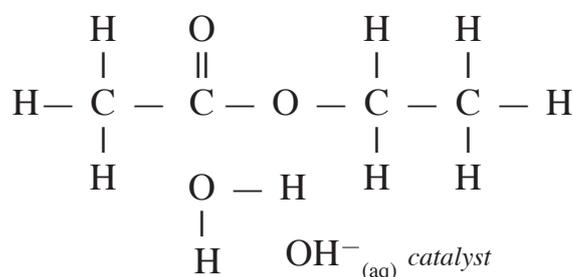


Earlier in this section you learnt that **water** can be used to **break an ester** apart to reform the **parent acid** and **parent alcohol**.

The reaction benefits from the presence of $\text{H}^+_{(\text{aq})}$ ions provided by **dilute sulphuric acid** $\text{H}_2\text{SO}_{4(\text{l})}$. This is often called **acid hydrolysis**.

Alkaline hydrolysis, heating the **ester** with **sodium hydroxide solution**, will also **break apart an ester** to reform the **parent alcohol**,

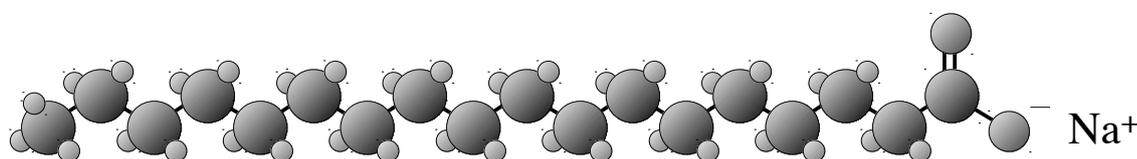
The **acid**, however, goes on to **react** with the **alkali** so the **salt of the acid** is formed instead. In this case **sodium ethanoate** ($\text{CH}_3\text{COO}^- \text{Na}^+$) would be made.



Alkaline hydrolysis of a **glyceride** molecule will yield **glycerol** and the **sodium salts** of its **fatty acids**.

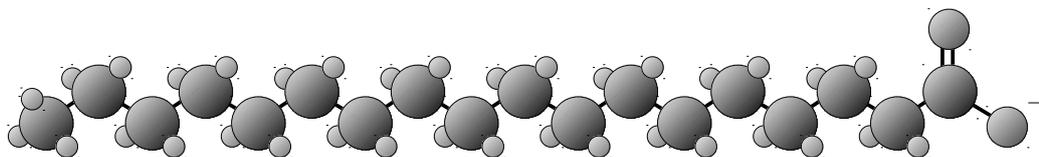
Glyceryl tristearate (found in most animal fats) will produce **sodium stearate** which was one of the earliest and most common **soaps** ever made.

Thousands of years ago, probably by accident, people discovered that boiling animal fat with alkali rocks produced a scum that, when cooled and solidified, could be used as a cleansing agent



Structurally, what makes these molecules capable of acting as a **cleansing agent** is their '**ionic head**' - COO^- , and long '**covalent tail**' - $\text{C}_{17}\text{H}_{35}$.

Covalent Tail - most of the **hydrocarbon tail** is far enough away from the **ionic head** that it maintains the **properties** of a typical **covalent** molecule - namely, **weak London Dispersion forces** between neighbouring molecules.



Ionic Head - the **ionic heads** can set up **strong attractions** - of similar strength to the **hydrogen bonding** between **water** molecules - allowing the **heads to dissolve in water**.

Water, particularly **hot water**, is a very effective **cleansing agent**. The strength of the **hydrogen bonding** between its **polar molecules** is **similar** to the strength of **attractions** in many **ionic substances**, which allows **water** to **dissolve** them.

Many **covalent** substances are **polar enough** to also dissolve in water. The main problem is with **covalent** substances with very **weak polar attractions** or only **Van der Waals attractions**



When mixed with **water**, these **pure covalent liquids** form **separate layers** as the difference in **intermolecular forces** is too **great** to allow **mixing**.

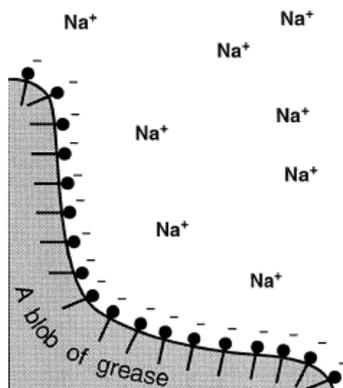
Shaking can form a **temporary emulsion** as small drops of 'oil' float in the water.



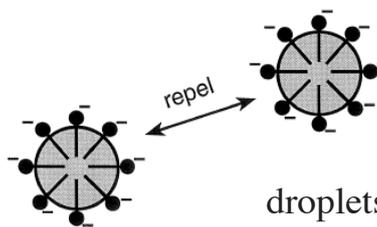
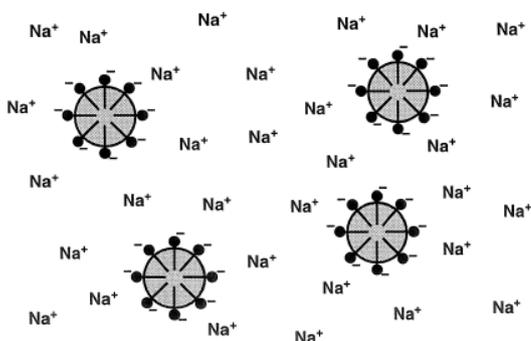
On standing, the two liquids will **separate** out again.

This makes it very difficult for water to remove **greasy** or **oily stains** from **clothes**, **plates** or even **people**.

Soap molecules cannot make grease '**dissolve**' in water, but they can prevent the tiny **droplets of grease** from **reforming** into large **blobs** which would stick to the surface of the **clothes**, **plates** or **skin of a person**.



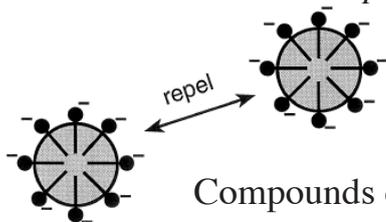
Instead they keep the tiny **globules suspended** in the water so that they can be **rinsed** away with the water.



The **negatively charged ionic heads** set up **strong repulsions** which prevent the droplets reforming into large blobs.

Detergents

This activity considers the problems caused by 'hard' water and the need for soapless detergents



To be effective, it is important that the *soap* molecule is able to *dissolve* and *dissociate* - that the Na^+ ion detaches to leave a *negatively* charged head.

Compounds containing *Group I* metals (The *Alkali* metals) are always *soluble* but water can also contain *Group II* metals (The *Alkali Earth* metals). Scotland gets its water from 'above ground' - water flows over hard *igneous* rocks to collect in *lochs* and *reservoirs*. There are very few Ca^{2+} and Mg^{2+} ions and the water is described as 'soft'.

In much of England, particularly the *South*, water *percolates* through rocks such as *limestone* and collects below ground from where it is *pumped* to the surface or flows out into rivers and lakes. The water has *higher* levels of Ca^{2+} and Mg^{2+} ions and is described as 'hard'.



This causes a number of problems such as *limescale* build up in *water pipes*, *kettles* and *water heaters*. In addition, normal soaps struggle to work as they form an *insoluble scum* with Ca^{2+} and Mg^{2+} ions.

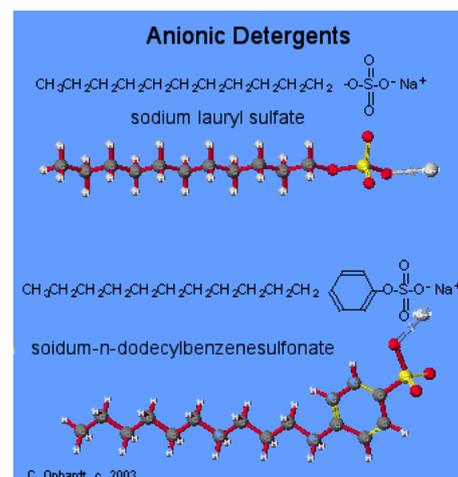
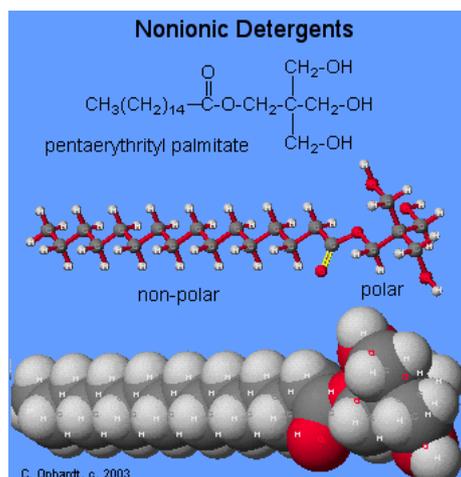


One solution is to *treat* the water by passing it through '*filters*' that absorb the Ca^{2+} and Mg^{2+} ions and replace them with Na^+ and K^+ ions which '*softens*' the water.



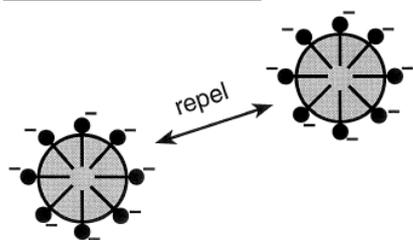
The alternative was to redesign the *soap* molecule to create a molecule that did not form an *insoluble scum* with Ca^{2+} and Mg^{2+} ions. These '*man made*' *soaps* are usually referred to as *detergents*.

There are many variations but they all have the '*covalent tail*' with '*polar/ionic head*' required.

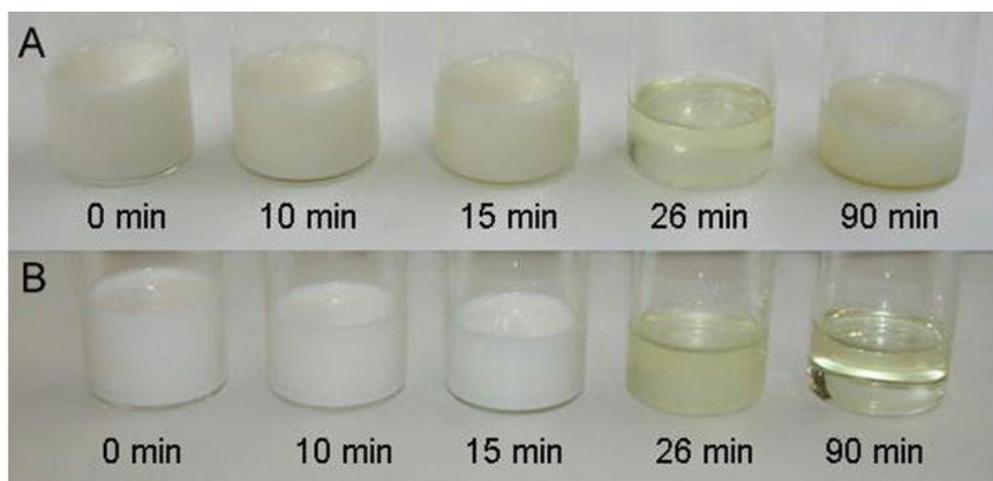


Emulsions

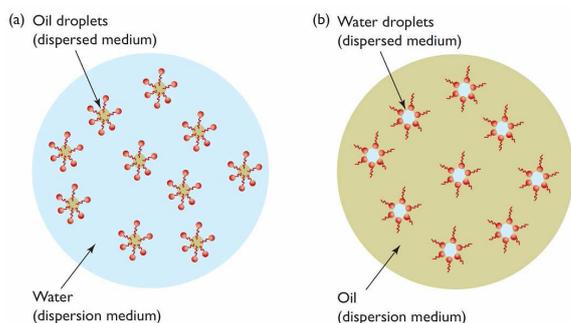
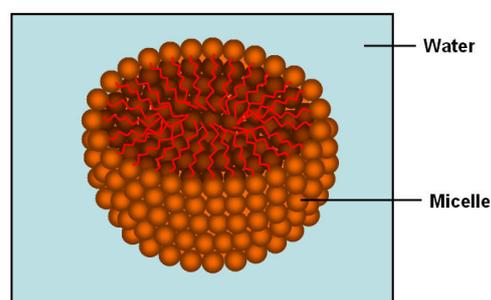
This activity considers the use of emulsifying agents to form stable emulsions.



Oil and **water** will *mix*, but they form an **emulsion** with droplets *suspended*. This is **unstable** and the liquids will *separate* (as shown in **B**).



Soap molecules can be very effective at **stabilising emulsions** (as shown in **A**), by positioning themselves at the **surface** of a droplet of oil with **covalent tails** in the oil and **ionic heads** in the water. These droplets are referred to as **micelles** and molecules like **soap** are described as **surfactants**. The **negative charges** on the **surface** prevent/slow down the **separation** of the liquids and help form a **stable emulsion**. **Emulsion paints** are good examples of **stable** oil/water mixtures.



Cream: Oil-in-Water

Butter: Water-in-Oil

Oil in Water

egg yolk

milk

ice cream

salad cream

mayonnaise

Water in Oil

margarine

butter

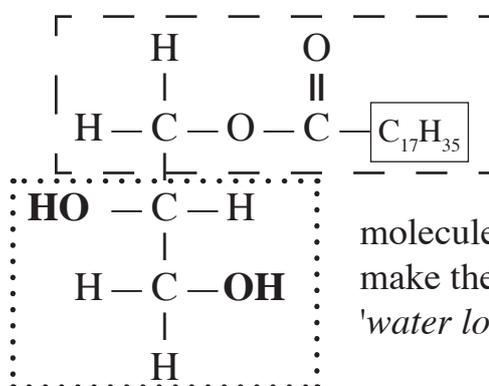
skin cream

moisturiser

There are very few foods that do not contain both **oils/fats** and **water**. Therefore **emulsifying agents** are widely used in the food industry.

Traditional **soap molecules** can have a negative impact on **taste** so other, very similar, molecules have been developed.

A convenient way to make an **emulsifying agent** is to react a **glycerol** molecule with a **single fatty acid** (instead of the 'normal' 3 used to make an edible fat or oil).



Though **not ionic** (like the head of a **soap**

molecule) the **hydroxyl** groups make the head **hydroscopic** - 'water loving' - **water soluble**.



The **largely hydrocarbon fatty acid**, as well as the **ester link**, make the tail **hydrophobic** - 'water hating' - **oil soluble**.

SELF CHECK

6.5

Q1. Soaps are produced from fats and oils by

- A alkaline hydrolysis
- B catalytic cracking
- C acidic condensation
- D dehydrogenation

Q2. Which of the following is a typical soap molecule?

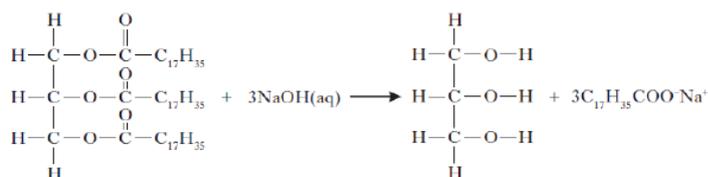
<p>A</p> $\text{CH}_3 - \overset{\text{O}}{\parallel} \text{C} - \text{O}^- \text{Na}^+$	<p>B</p> $\text{C}_{17}\text{H}_{35} - \overset{\text{O}}{\parallel} \text{C} - \text{O}^+ \text{Na}^-$
<p>C</p> $\text{C}_{17}\text{H}_{35} - \overset{\text{O}}{\parallel} \text{C} - \text{O}^- \text{Na}^+$	<p>D</p> $\text{C}_{17}\text{H}_{35} - \overset{\text{O}}{\parallel} \text{C}^- \text{Na}^+$

Q3. Soap molecules are effective cleansing agents because they have

- A grease-soluble hydrocarbon chains joined to water-soluble ionic ends
- B water-soluble hydrocarbon chains joined to grease-soluble ionic ends
- C grease-soluble acidic parts joined to water-soluble alkaline ends
- D water-soluble acidic parts joined to grease-soluble alkaline ends

Q4. Which of following statements about detergents is *incorrect* ?

- A soap detergents can be manufactured from fat
- B soapless detergents do not form foam in water
- C the hydrophilic 'head' of a detergent can be anionic or cationic
- D the hydrophilic 'head' of a detergent can be polar



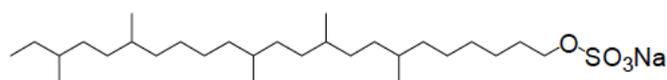
Q5. Soaps are produced by the above reaction. This reaction is an example of

- A condensation
- B esterification
- C hydrolysis
- D oxidation

Q6. When a detergent is added to a beaker of oil and water and then the solution is shaken, an emulsion is formed. The emulsion consists of

- A oil and detergent
- B oil droplets and water
- C water, oil and oil droplets
- D water, oil, detergent and oil droplets

Q7. Consider the following detergent particle



Which part of this structure makes the detergent particle hydrophilic?

- A
- B
- C $-\text{OSO}_3^-$
- D Na^+

Q8. Shaking a soap with grease produces tiny globules which cannot rejoin because

- A they are no longer soluble in water
- B the grease molecules have hydrolysed
- C the molecules in them have changed shape
- D charges on their surfaces make them repel

6.6 Terpenes - Fragrance Molecules

This lesson topic introduces terpenes which are a major component of essential oils.

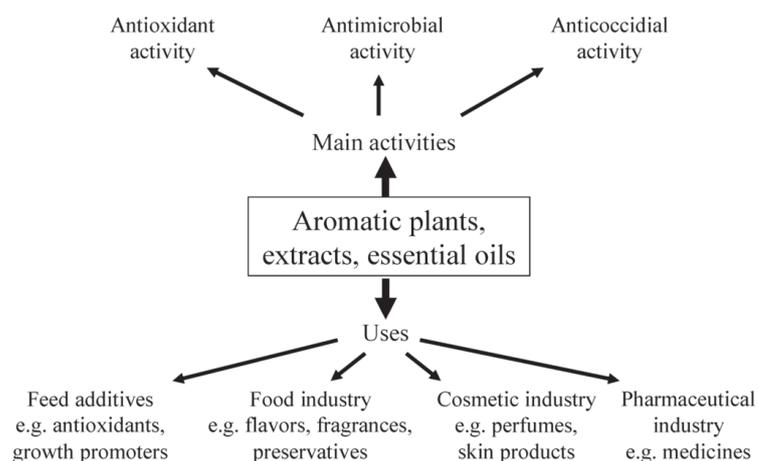
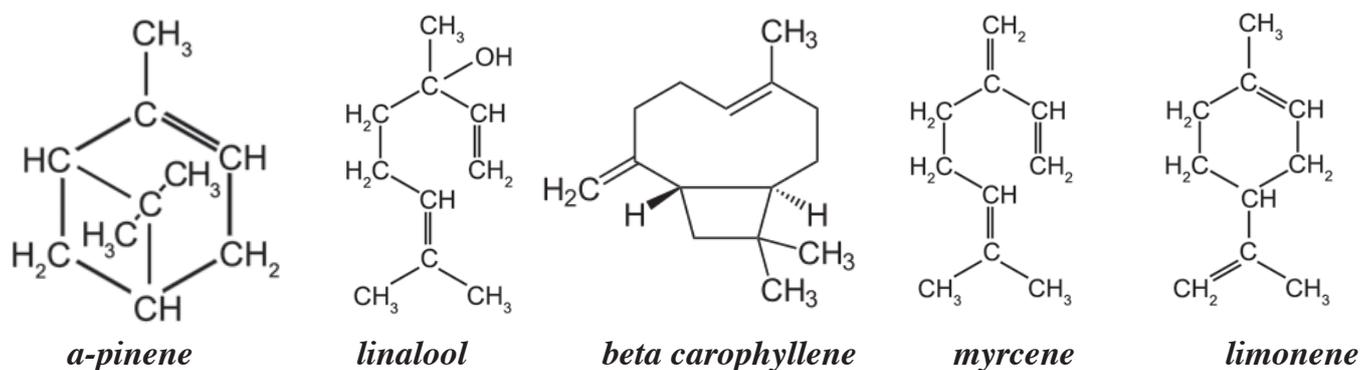
Essential Oils

This activity considers the role of essential oils.

A-PINENE	LINALOOL	BETA CARYOPHYLLENE	MYRCENE	LIMONENE
ANTI-INFLAMMATORY BRONCHODILATOR AIDS MEMORY ANTI-BACTERIAL	ANESTHETIC ANTI-CONVULSANT ANALGESIC ANTI-ANXIETY	ANTI-INFLAMMATORY ANALGESIC PROTECTS CELLS LINING THE DIGESTIVE TRACT	CONTRIBUTES TO SEDATIVE EFFECT OF STRONG INDICAS SLEEP AID MUSCLE RELAXANT	TREATS ACID REFLUX ANTI-ANXIETY ANTIDEPRESSANT
also found in pine needles	also found in lavender	also found in black pepper	also found in hops	also found in citrus
				

Essential oils are the **concentrated** extracts of **volatile**, **non-water-soluble** **aroma** compounds from plants. 'Essential' refers to the fact that the oil carries the **distinctive essence** (*scent*) of the plant. **Essential oils** are mixtures of organic compound, though **Terpenes** are the key components in most essential oils.

The **distinctive** character of an essential oil can be attributed to the **functional group** present in its key molecule. **Esters**, **aldehydes**, **ketones** and **alcohols** are all found in **essential oils** but many are **terpenes**.



As well as their role as **fragrance molecules**, there are 3 other reasons why essential oils might be used:

antioxidant activity
antimicrobial activity
anticoccidial activity

Coccidia are tiny single-celled **parasites** that live in the wall of an animal's intestine.

Terpenes

This activity considers the structure of terpenes

Terpenes are **unsaturated** compounds formed by joining together **isoprene** units.

Isoprene is the common name for **2-methylbuta-1,3-diene**

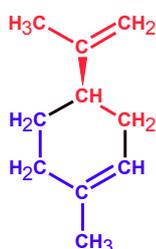
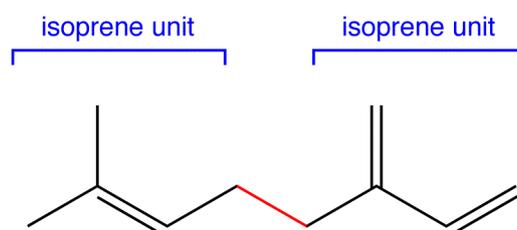
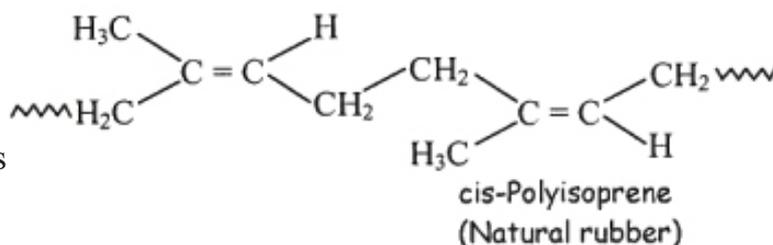
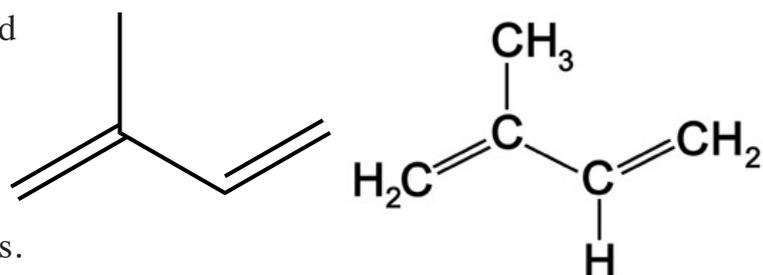
Isoprene is made in the chloroplasts of plants.

Isoprene is produced and **emitted** by many species of trees into the **atmosphere**, where they can be converted by **free radicals** (like the **hydroxyl** (OH) radical) into various species, such as **aldehydes**, **hydroperoxides**, **organic nitrates**, and **epoxides**.

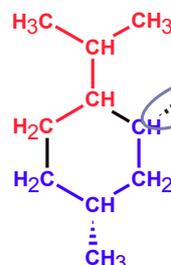
It has been proposed that **isoprene** emission is specifically used by plants to protect against large fluctuations in leaf **temperature**. **Isoprene** also confers **resistance** to **reactive oxygen species**.

The **isoprene** skeleton can be found in naturally occurring compounds called **terpenes**. They can join 'head to tail' to form **linear terpenes**. An example is **myrcene**, a component of plants, including bay, ylang-ylang and thyme

Other **terpenes** are **cyclic**, whilst other have been **oxidised** to form **hydroxyl** or **carbonyl** groups:

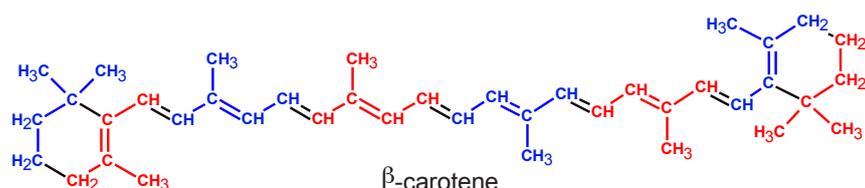


Limonene
(skin of citrus fruits)



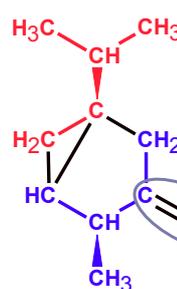
Menthol
(peppermint)

This terpene has been oxidised to a terpenoid



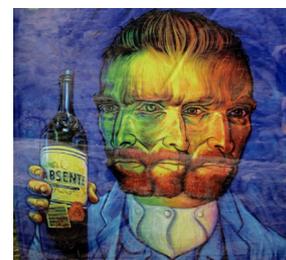
8 isoprene units

40 carbon atoms



Thujone
(Absinthe)

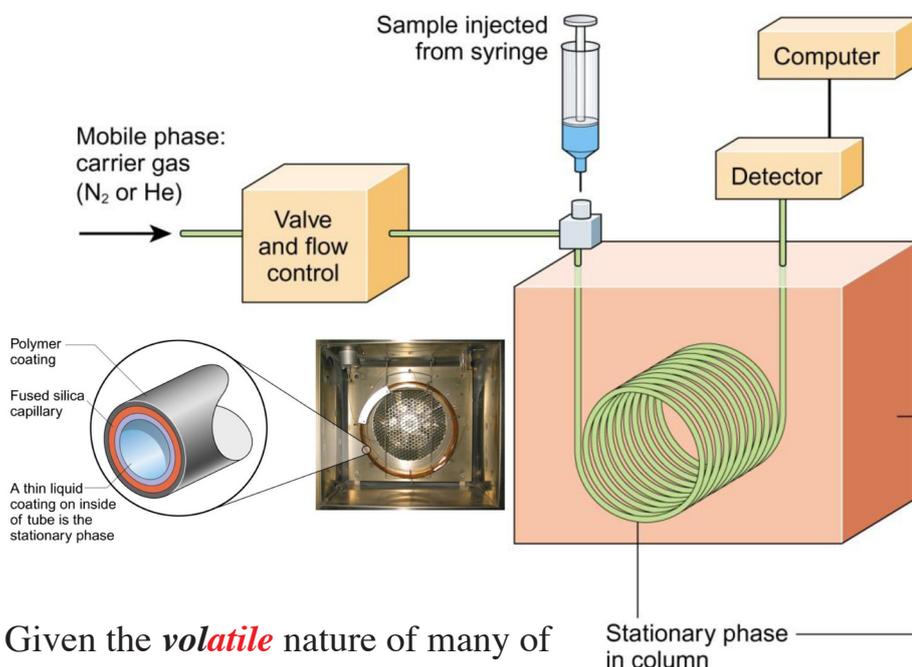
This terpene has been oxidised to a terpenoid



Gas Chromatography

This activity considers the technique of gas chromatography as a means of separating and identifying volatile molecules.

As with *Paper Chromatography*, met in an earlier topic, *molecules* move in a *mobile phase* but can also spend some of their time 'trapped' in a *stationary phase*.



GC door closed

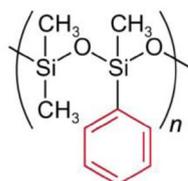
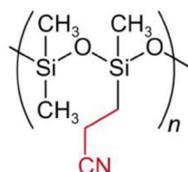
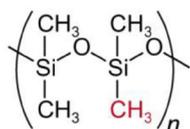
Sample injector

Detector

GC door open

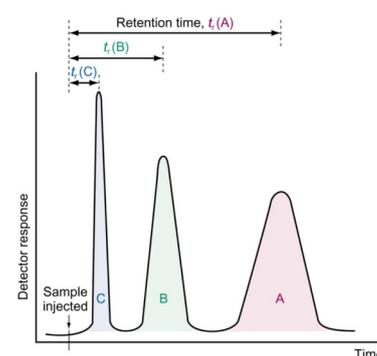
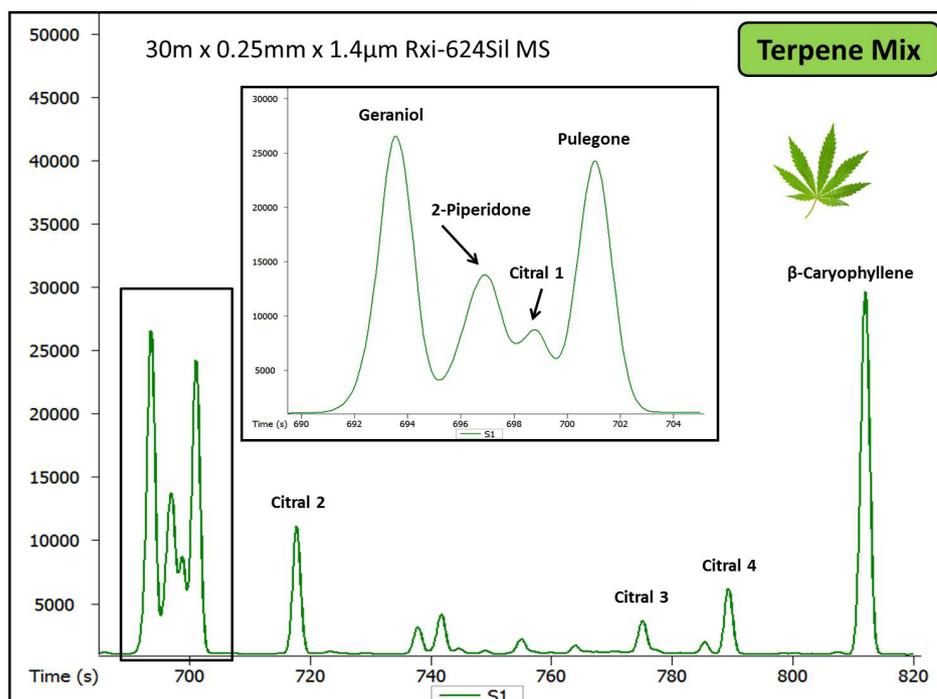
Oven

Given the *volatile* nature of many of the molecules studied in this topic, an *inert carrier gas* is chosen to be the *mobile phase*. Thin metal coils are either packed with suitable solid *powders* (*Gas Solid Chromatography (GSC)*) or *liquids* are *absorbed* onto a polymer coating (*Gas Liquid Chromatography (CLC)*).



Siloxane polymers

Polar or *non-polar* materials can be used depending on the molecules being identified.

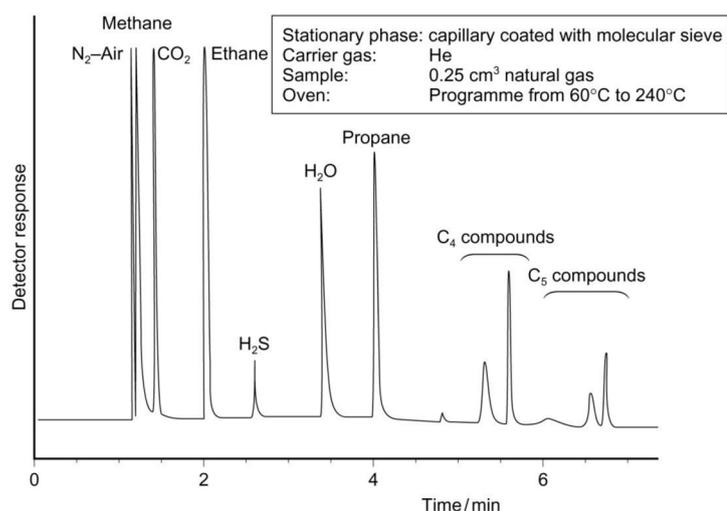


Different molecules take *different* times to move through the coil depending on factors such as *mass*, *shape* and *polarity*.

The *area* under each signal represents the *amount* present.

Mineral Oils

This activity examines the chromatograms of natural gas and unleaded petrol and establishes some of the factors that effect retention times.



Whilst **methane** is the main component (off the scale) in **natural gas**, there are many other molecules present.

molecule	mass	time/min
methane	16	1.5
ethane	30	2.0
propane	44	4.0

Since these molecules should have the **same polarity** we see that:

as mass increases, the time increases

Comparing molecules of **similar mass** like **methane** with **water** and **ethane** with **hydrogen sulphide** shows the effect of **polarity**:

as polaity increases, the time increases

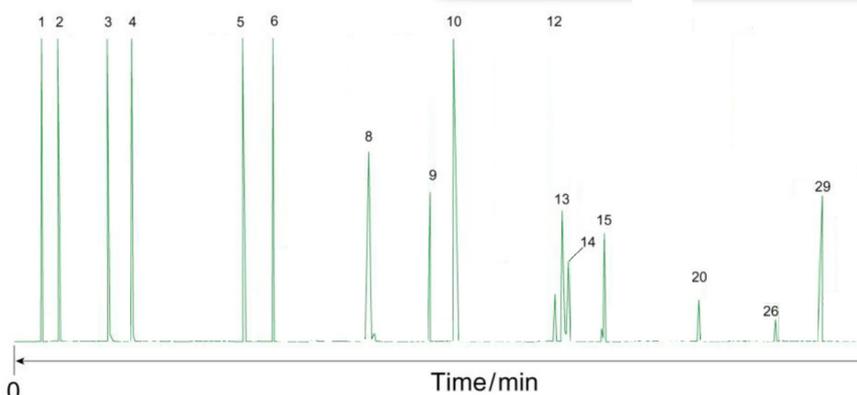
Propane and **carbon dioxide** have **similar mass** and **similar polarity**. **Carbon dioxide** is, however, a more **compact** molecule and moves through the coil **more quickly**.

molecule	mass	polar	time/min
methane	16	no	1.25
water	18	yes	3.5
ethane	30	no	2.0
hydrogen sulphide	34	yes	2.6
propane	44	no	4.0
carbon dioxide	44	no	1.5

Unleaded petrol

Stationary phase: Poly(dimethylsiloxane)
 100 m x 0.25 mm capillary
 Carrier gas: He
 Sample: 1 microlitre of unleaded petrol
 Oven: Programme from 0°C to 180°C

- methane
- butane
- 2-methylbutane
- pentane
- hexane
- methylcyclopentane
- cyclohexane
- 2,2,4-trimethylpentane
- heptane
- nonane
- decane
- 2,3,3-trimethylpentane
- 2-methylheptane
- 4-methylheptane
- octane
- undecane



Examination of the simplified **chromatogram** of **unleaded petrol** confirms that:

as mass ↗, the time ↗

Comparing **branched isomers** with the **equivalent chain** molecule:

as no. of branches ↗, the time ↗

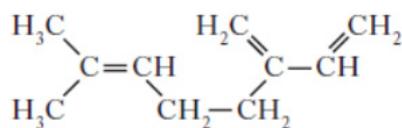
and looking at the **positions** of **branches**, 2-methyl to 4-methyl:

*branch position ↗,
the time ↗*

SELF CHECK

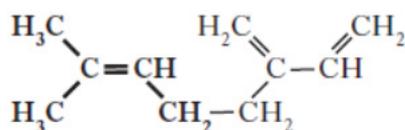
6.6

Q1. Myrcene is a simple terpene.

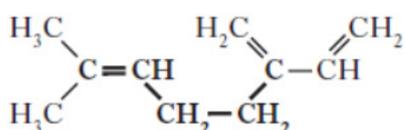


Terpenes contain at least one isoprene unit. Which of the following shows a correctly highlighted isoprene unit?

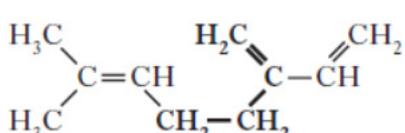
A



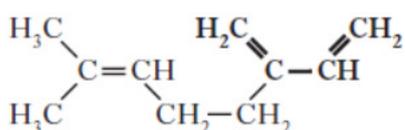
B



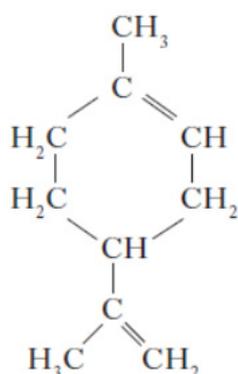
C



D



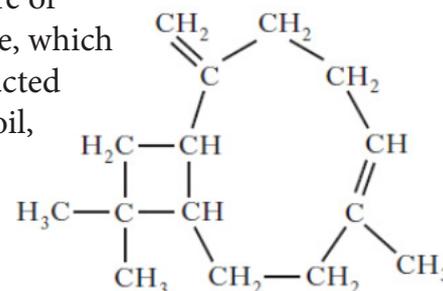
Q2. Limonene is one of the terpene molecules responsible for the flavour of lemons.



How many isoprene molecules are used in the production of one limonene molecule?

- A** 1
B 2
C 3
D 4

Q3. The structure of carophyllene, which can be extracted from clove oil, is



Which of the following would be the best solvent for extracting carophyllene?

- A** hexane
B hexanol
C hexanal
D hexanone

Q4. Which of the following is the most suitable gas to use as a carrier gas in a gas chromatogram?

- A** oxygen
B carbon dioxide
C helium
D methane

Q5. Acetone is an organic molecule with a formula of CH_3COCH_3 . A student runs a sample of acetone through a GLC at 50°C .

The acetone produces a peak after 4.2 mins.

The student then injects a mixture of unknown organic substances into the same column at the same temperature.

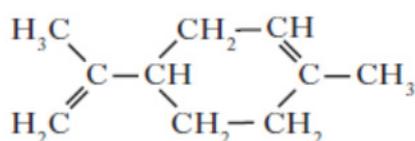
There are peaks after 3.1, 4.2 and 7.4 mins. From this information, it can be concluded that ...

- A** the mixture has 3 components, but acetone is not one of them.
B the mixture has at least 3 components, one of which might be acetone.
C the mixture has 3 components, one of which must be acetone.
D the mixture has at least 3 components, one of which must be acetone

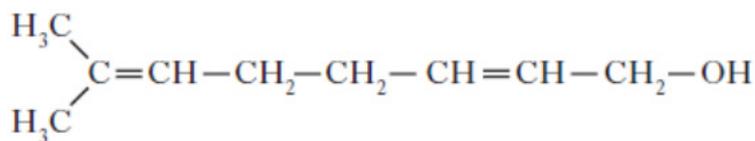
HOME PRACTICE

6.6

Q1. Two typical compounds that are present in many perfumes are shown.



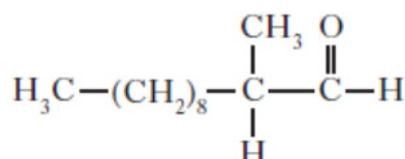
$C_{10}H_{16}$
limonene



$C_9H_{16}O$
geraniol

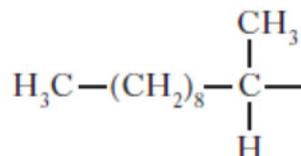
a) Why does geraniol evaporate more slowly than limonene? 1

The structure of one of the first synthetic scents used in perfume is shown opposite.



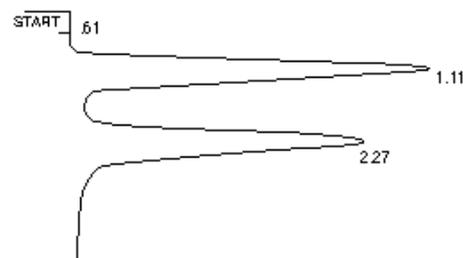
b) Name the family of carbonyl compounds to which this synthetic scent belongs. 1

c) Copy and complete the structure opposite to show the product formed when this scent is oxidised.



d) Suggest what effect the oxidation would have on the fragrance of the molecule. 1

Q2. Study the chromatograph of a mixture of hydrocarbons A and B. Compound A has the shorter retention time.



a) What is the retention time of compound A? 1
Compound B?

RT	AREA	TYPE	AR/HT	AREA%
0.61	XXXX	XX	XX	0.009
1.11	XXXX	XX	XX	55.874
2.27	XXXX	XX	XX	44.117

b) Which compound is present in a larger amount? 1

c) Which compound has the lower boiling point? 1

d) You suspect that compound B is octane. What can you do to provide supporting evidence for this hypothesis? 1

Q3. Traces of a liquid were discovered in a bottle believed to contain perfume belonging to Queen Hatshepsut, ruler of Egypt over 3500 years ago. Perfumes were made by dissolving plant extracts containing pleasant smelling terpenes and esters in an edible oil. A little ethanol and water may also have been added.

Using your knowledge of chemistry, comment on the possible smell(s) when such a bottle is opened after being stored for thousands of years. 3

Total (11)

6.7 Amines, Amides & Amino Acids

This topic introduces the amine functional groups and, in particular, its ability to form a link with other groups - the amide or peptide link.

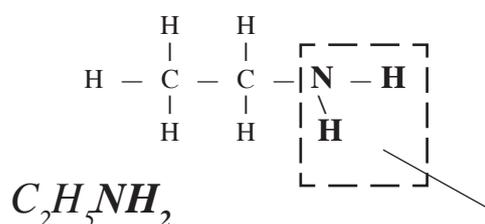
Amines & Amides

This activity deals with how to recognise and identify amines and introduces a few of their properties.



Amines are the **organic** relatives of **ammonia**, NH_3 , and like **ammonia**, their properties are mainly due to the small, highly **electronegative** Nitrogen atom.

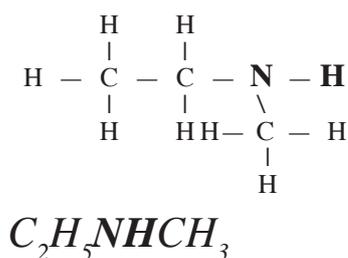
The simplest amines are called **primary amines** and will have a **carbon** chain (*alkyl group*, R—) in place of *one* of the **hydrogen** atoms, $\text{R}-\text{NH}_2$



This amine is usually called simply *ethylamine*, though its more formal name is *aminoethane*.

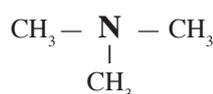
You do not need to learn the formal systematic naming of amines.

This is the **amino functional group**



Replacing *two* of the **hydrogens** with carbon chains produces what is called a **secondary amine**.

Simple naming is used whenever possible, so this molecule is called *ethylmethanamine*.



Replacing *all* of the **hydrogens** with three carbon chains produces what is called a **tertiary amine**.

Simple naming is used whenever possible, so this molecule is called *trimethanamine*.



We will concentrate mainly on **primary amines** and molecules containing the **amino group**, $-\text{NH}_2$, though you are expected to at least recognise other amines when you meet them.

Properties of Amines

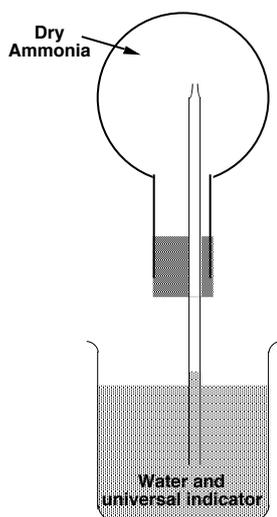
This activity will remind you of some of the properties that *ammonia* has and which are shared by the amines



Ammonia is a gas with a very **pungent** smell. Similarly the smaller amines are **gases** or **volatile liquids** with very **unpleasant** smells similar to rotten fish or.....



~~F P~~



Last year you were shown that **ammonia** is an *extremely soluble* gas - the **fountain** experiment.

This is due to the fact that **ammonia** molecules are like **water** molecules and have *very strong hydrogen bonding* between them. *Small primary amines* will also have *strong hydrogen bonding* and will also be *very soluble*. **Secondary amines** are *less soluble* and **tertiary amines** even *less soluble*.

Ammonia was the *only alkali* gas met during Standard grade. **Amines** are *organic alkalis* and will **dissolve** in water to produce **hydroxide** ions, OH⁻.

Many ions of **Transition metals** are **coloured** but only when *hydrated* - surrounded by **water** molecules. Cu²⁺ ions - or more accurately [Cu(H₂O)₄]²⁺ ions - are **blue** in colour. **Ammonia** and the smaller **amines** can replace these **water** molecules but this will change the **colour** of these ions:



These colour changes are sometimes used as a 'Test' for ammonia/amines

Reactions of Amines

This activity introduce you to the reactions of ammonia/ amines

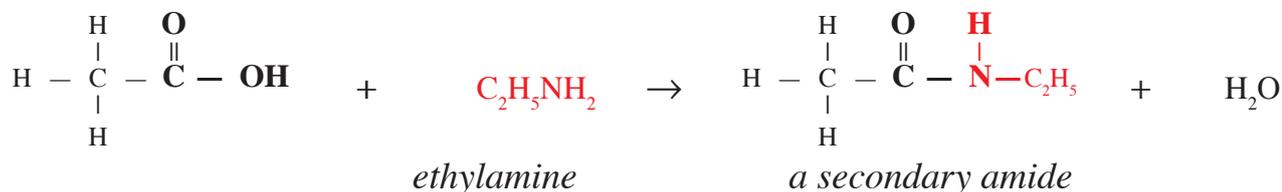
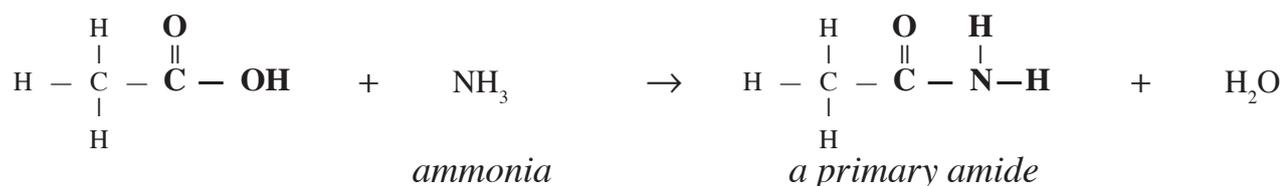
- ① *Dissolve in water to produce Alkalis-* solutions containing the **hydroxide** ion.



- ② *React with acids to produce Salts-*



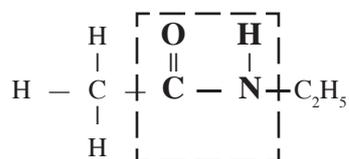
③ React with organic acids to produce Amides-



Amides are not an 'important' family and you are not required to learn how to name them.

The reactions above are very similar to *esterification* - **acid** groups can react with **amine** groups in a **condensation** reaction that allows the two molecules to *join together*.

This is an important reaction in nature, and you will need to learn to recognise the **amide link** (the **peptide link**)



Proteins

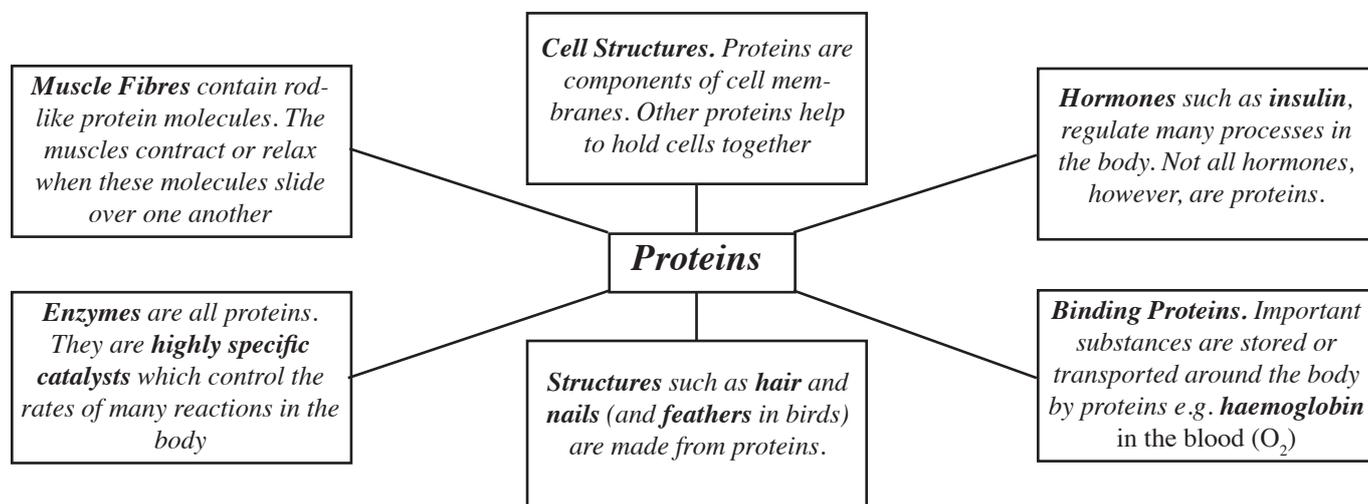
This activity will introduce you to some proteins and the roles played by proteins in living organisms

- **peas and beans**
- **meat**
- **fish**
- **cheese**
- **eggs**
- **hide & skin**
- **wool & silk**

When **protein** is mentioned, most people will think of *foodstuffs* that contain **protein** and make up a very important part of our *diet*.

When we eat **proteins** they are **digested** (*broken down*) into simpler molecules called **amino acids**.

These **amino acids** are then reconstituted as **proteins** that fulfil a large number of important roles in living organisms.



Amino Acids

This activity will introduce you to the structure of amino acids and explains why some are labelled as essential amino acids

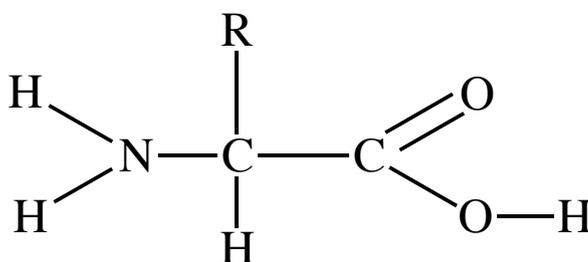
All the **proteins** in the world are made from about 20 **amino acids**.

glycine	Gly	—H
alanine	Ala	—CH ₃
valine	Val	—CH(CH ₃) ₂
leucine	Leu	—CH ₂ —CH(CH ₃) ₂
isoleucine	Ile	—CH(CH ₃)—CH ₂ —CH ₃
phenylalanine	Phe	—CH ₂ —C ₆ H ₅
proline	Pro	
tryptophan	Trp	
serine	Ser	—CH ₂ —OH
threonine	Thr	—CH(CH ₃)—OH

These 20 **amino acids** have a common structure. (one exception)

A central **carbon** atom has an acid group (**carboxyl**), an **amino** group and a **hydrogen** atom attached.

The final group attached to the carbon is *different* for each **amino acid** and is usually represented by —R.



cysteine	Cys	—CH ₂ —SH
methionine	Met	—CH ₂ —CH ₂ —S—CH ₃
aspartic acid	Asp	—CH ₂ —C(=O)OH
glutamic acid	Glu	—CH ₂ —CH ₂ —C(=O)OH
asparagine	Asn	—CH ₂ —C(=O)NH ₂
glutamine	Gln	—CH ₂ —CH ₂ —C(=O)NH ₂
lysine	Tyr	—CH ₂ —C ₆ H ₄ —OH
histidine	His	—CH ₂ —C ₄ H ₃ N ₂
lysine	Lys	—CH ₂ —CH ₂ —CH ₂ —CH ₂ —NH ₂
arginine	Arg	—CH ₂ —CH ₂ —CH ₂ —NH—C(=NH)NH ₂

When we **digest proteins** we break them down into **amino acids** which we then use to build new **proteins**. Most **amino acids** (12) can be made from **carbohydrates** and other **amino acids** so it is not crucial that we eat **foods** containing these amino acids.

The remaining amino acids (8), however, *cannot be made* and therefore must be part of our food intake. These are classified as **essential amino acids** (though we need all 20 to remain healthy).

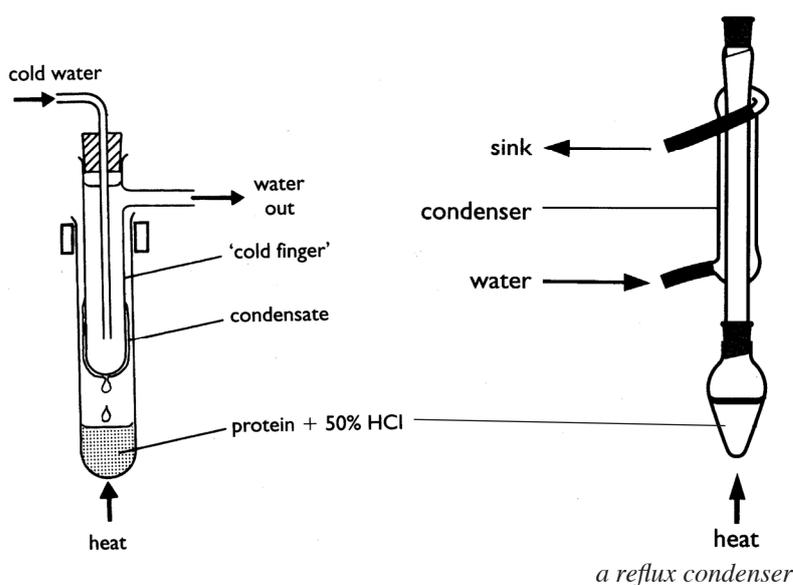
Proteins can be broken down in the lab by **heating** them for several hours with dilute acid.

This reaction is called **hydrolysis**,

hydro = **water**

lysis = **splitting apart**

To prevent the water **evaporating** away before the **reaction** has finished we can use apparatus like these.



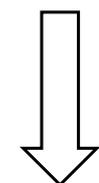
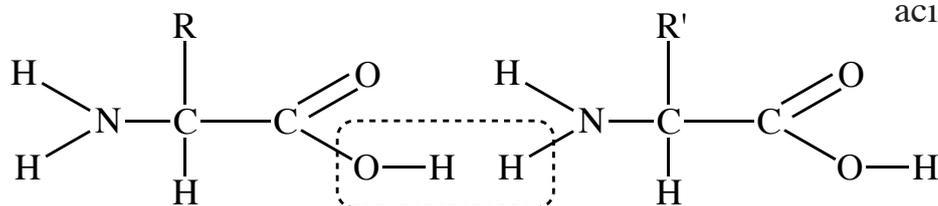
Having **hydrolysed** a **protein**, we will often attempt to **identify** the **amino acids** that made up the **protein** using **paper chromatography**. They all have the **amine** group and **carboxyl** group but differences in the **polarity** of their **side-groups** will effect their **retention** in the **stationary phase**.

Making Peptides

two **amino acids**

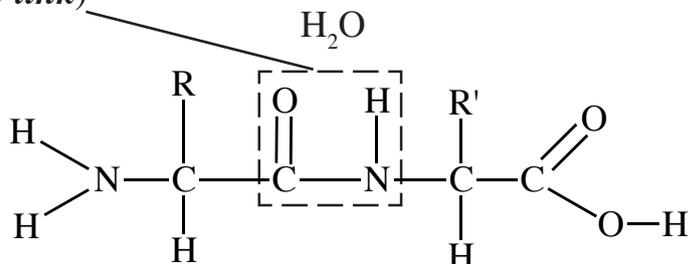
(*R* and *R'* represent different side chains)

link together to form a **dipeptide**



Water is also produced and this is a **condensation** reaction

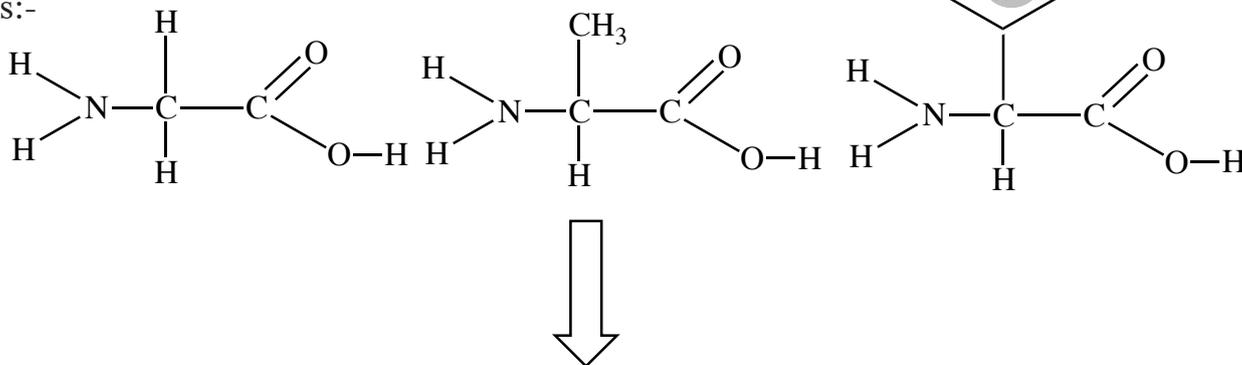
This is known as the **peptide link** (or **amide link**)



The **carboxylic** group from one amino acid links with the **amino** group of another amino acid

Tripeptides are made by joining **three** amino acids together. **Peptides** are 'named' using the accepted **abbreviations** of the **amino acids** they contain.

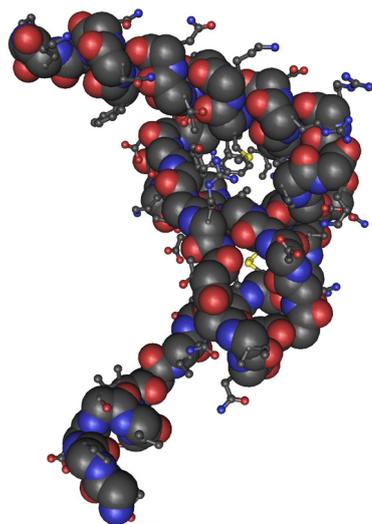
For example, the peptide made from *Glycine*, *Alanine* and *Phenylalanine* would be labelled *GlyAlaPhe* and would look like this:-



By convention, the **amino acids** are always drawn with their **amino** groups to the left, in the order they appear in the **peptide** name - a different **peptide** would be formed if we'd lined the 3 **amino acids** up facing the 'wrong way'.

Polypeptides can contain up to about 40 **amino acids**; more than 40 and we tend to call it a **protein**, though the distinction between a **polypeptide** and a **protein** is an *arbitrary* one.

Making Proteins



This activity looks at how proteins are made

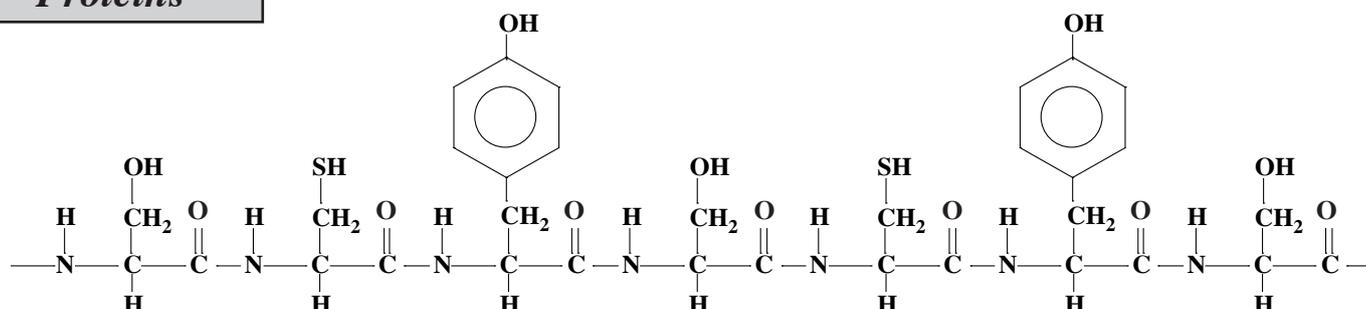
Proteins are **condensation polymers** which can contain several thousand amino acids. A massive variety of **proteins** can be made by arranging up to 26 **amino acids** in varying numbers and varying orders.

Proteins comprise a large part of an animal's **diet**. During **digestion** the **animal** and **vegetable** proteins are **hydrolysed** into their component **amino acids**. Some **amino acids** can be synthesised in the body, but others (the **essential amino acids**) have to be present in the diet.

Proteins required for the body's **specific** needs are built up from **amino acids** in the body cells according to information supplied by DNA in the cell **nuclei**.

Breakdown of Proteins

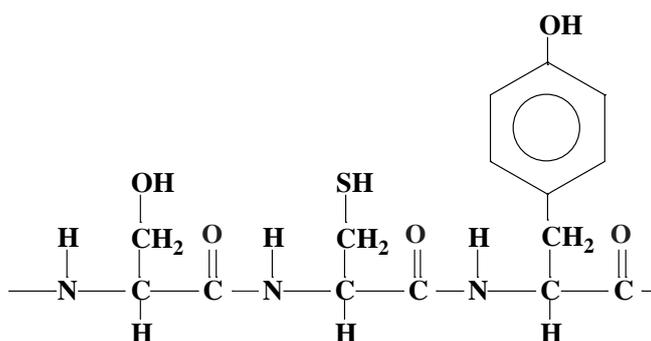
This activity looks at the hydrolysis of proteins to recreate the original amino acids used to form them.



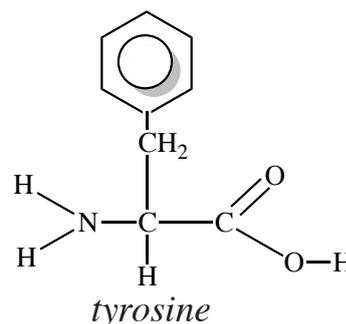
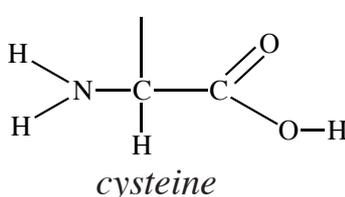
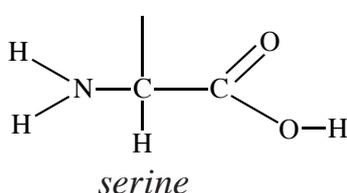
You will only ever see a **fragment** of a **protein chain** but it will be enough to allow you to recognise the “**repeating pattern**” and identify how many **different amino acids** are being used to make this protein.

In this case there are **three** amino acids in the “**repeating pattern**”.

The recognisable **peptide link** is used to show where one amino acid ends and the next one begins.



The **original amino acids** can then be drawn - remembering to replace the —H atoms and —OH groups lost when they joined together. In other words, the original **carboxyl** and **amino** groups are reformed.



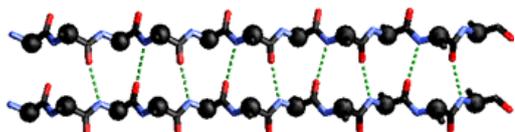
Structure of Proteins

This activity looks at how different structures for proteins depend on their constituent amino acids and affect their role

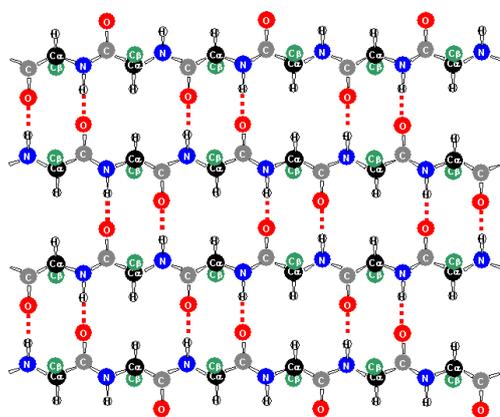


The *primary structure* of all **proteins** is long chains of **amino acids**. However, all along these chains are **polar groups** such as $-\text{N}^{\delta-}-\text{H}^{\delta+}$ and $-\text{C}=\text{O}^{\delta-}$ as well as **polar** and **non-polar** groups ($-\text{R}$) on each amino acid. A lot of **attractions** (and **repulsions**) are set up *within* and *between* chains, plus some reactions that lead to *permanent* bonds.

As a result of these extra bonds, *secondary structures* will be formed.

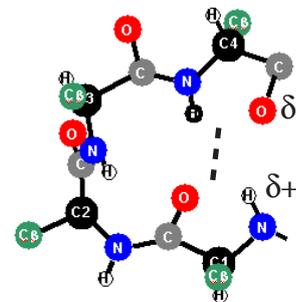


Chains can become linked together by **strong hydrogen bonds** between the chains.

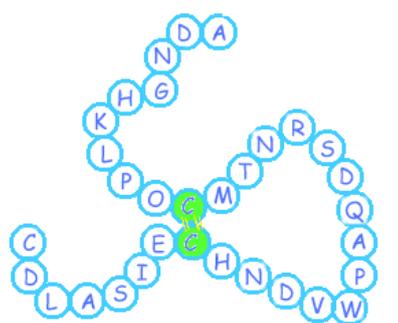


Many chains can link in this way to form a **sheet**.

Often the chains will **twist** around to form **strong hydrogen bonds** within the chain. The length of **4 amino acids** is usually enough to allow this to happen

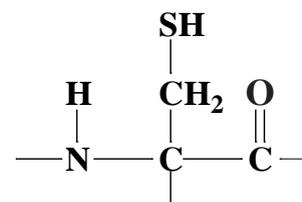


As a result of this *twisting* a **helix** chain will form.



Note: two cysteines form a disulphide bridge.

More **permanent bonds** can also be formed. For example, two *cysteine* side groups can be oxidised and lose hydrogen atoms to form a 'disulphide bridge'

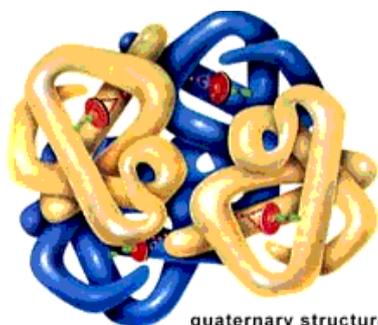


The **folding** of chains and **helixes** is what gives **proteins** their individual **shapes**.

Proteins which remain more elongated are referred to as **FIBROUS** proteins. These make up most *animal tissue* such as *muscles*. Other examples include *Keratin* found in *horns, hoof* and *hair*, and *Collagen* found in *tendons*.



tertiary structure
(folded individual peptide)



quaternary structure
(aggregation of two or more peptides)

Even more complicated structures called **GLOBULAR** proteins can result when a number of *peptide chains* join together. These **Globular** proteins are involved in the *maintenance and regulation of life processes*.

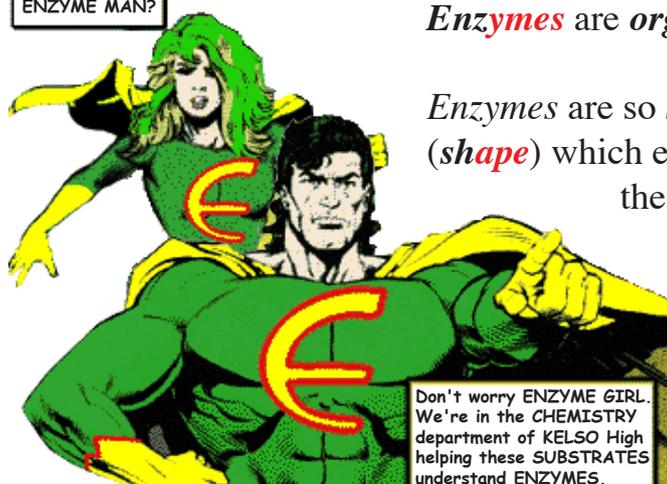
Examples include **hormones** e.g insulin, and **enzymes**.

Shape, and their ability to form various types of bonds to bind to other substances, are crucial to a proteins role.

Enzyme Shape

This activity looks at the importance of molecular shape to the way an enzyme functions

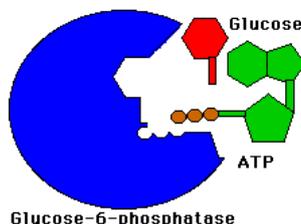
Where ARE we ENZYME MAN?



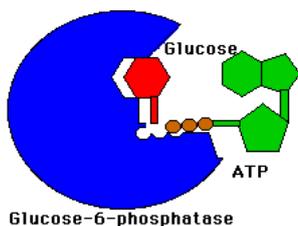
Enzymes are **organic catalysts** and all contain **protein**.

Enzymes are so **specific** because they have a precise structure (**shape**) which exactly matches the structure of the **substrate** - the molecule(s) which is/are reacting.

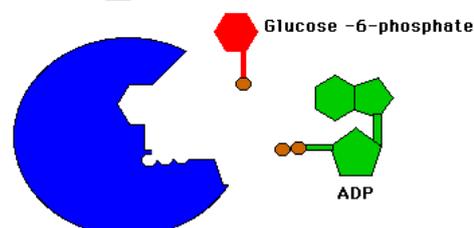
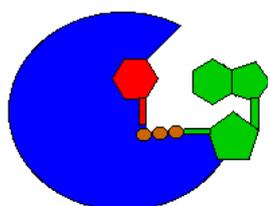
Enzymes will have an **active site** where reaction takes place. Within the **active site** chemical groups (some of the side chains on the **amino acids**) will form **bonds** with the **substrate** molecule



The **bonds** which bind the **substrate** to the **active site** have to be **weak** so that the products can easily leave the **active site** after the reaction. The bonds are usually **hydrogen bonds** or **interactions** between **ionic** groups.



While attached to the **active site**, bonds *within* the **substrate** molecule(s) will be **weakened** making it **easier** for the **substrate** to react - **lowering the activation energy** of the reaction as a **catalyst** should.



Sometimes, being attached to the **active site** will change the **shape** of the substrate bringing atoms or groups that need to react into **closer contact**. This helps overcome awkward '**collision geometry**'.

When talking about **enzymes** we often use the phrase '**lock and key**' to cover the importance of **correct shape** and the fact that each enzyme is likely to only work with one **specific** substrate.

The example on the left is a good illustration of how an enzyme works.

Denaturing

This activity looks at the factors which can change the shape of an enzyme and prevent it functioning.

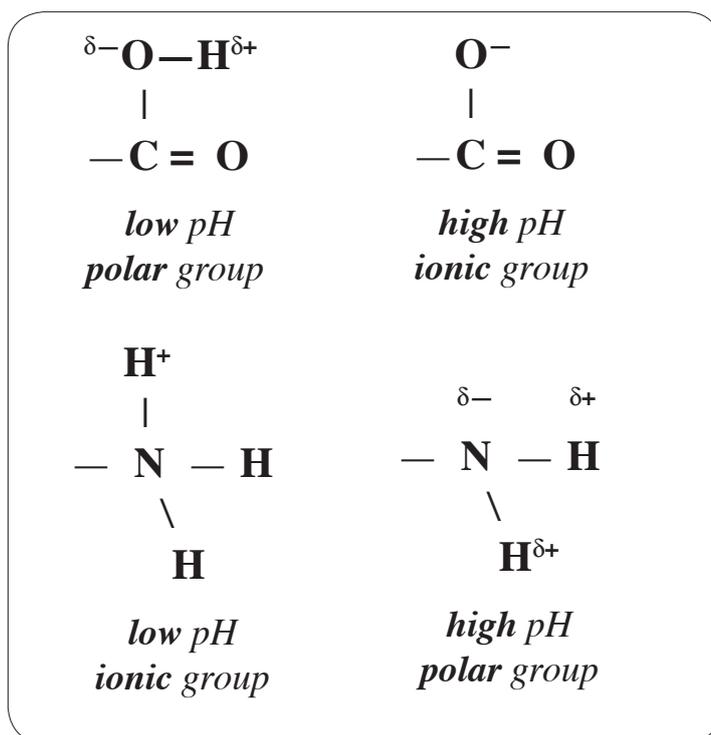
Many of the **groups** found on the **side chains** of **amino acids** (see page 43) are **ionisable** and will be affected by a change in **pH**

Groups such as $-\text{COOH}$ and $-\text{NH}_2$ are **polar** but can become **ionic** as the **pH** changes.

This can change the **active site** so that the **substrate** molecule will be unable to **bond** with the **enzyme**.

Changing the nature of some of these **groups** can also change the **shape** of the **enzyme**, as **folding** of the **peptide chain** may no longer happen at the same points.

If the **active site** is destroyed the enzyme is said to be **DENATURED**.



The structure of an **enzyme** is often held together by **weak polar**—**polar bonds** and **hydrogen bonds**. These can easily be **broken** by raising the **temperature**, which causes them to **vibrate** more vigorously. *So enzymes are sensitive to small changes in temperature or pH.*

Cooking Meat

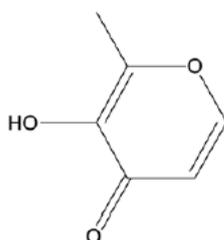
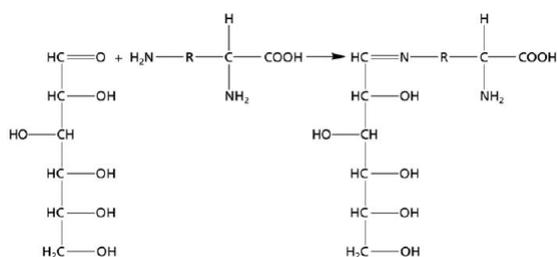
This activity looks at some of the reasons why we cook meat and some of the chemical changes that take place.



Whilst killing bacteria and other microorganisms to make meat safe to eat is a major reason for cooking meat, most of the reasons are to do with increasing the acceptability of the food.

Texture - *tough meat can be softened* - collagen is a **protein** that forms the **connective material** in meat. Too much makes meat **tough**. At **temperatures** above $60\text{ }^\circ\text{C}$, the **spiral protein** **unwinds** (**denatures**) and turns into **soft gelatine**.

Flavour- *browning improves appearance, flavour and aroma* - At **temperatures** above $140\text{ }^\circ\text{C}$ **reactions** between $-\text{NH}_2$ groups and **carbohydrates** produce **polymers** (**brown**) as well as **volatile flavour** and **aroma** molecules. **Maillard Reactions**.



maltol - a flavour molecule

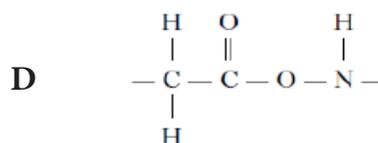
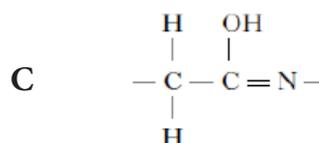
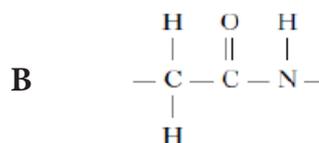
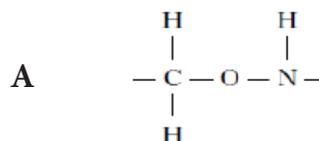
Other **less desirable** changes can happen during the **cooking** of meat - other **protein** molecules begin to change (denature) at about $40\text{ }^\circ\text{C}$ and cause the meat to **harden**, for example. Chefs often **brown** meat with a blow torch and then cook at **low temperatures**.

SELF CHECK

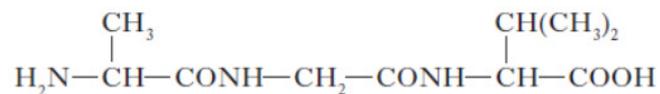
6.7

- Q1.** When a protein is denatured
- its overall shape is distorted
 - its amide links are hydrolysed
 - it is broken into separate peptide fragments
 - it decomposes into amino acids
- Q2.** Which of the following compounds can be classified as proteins?
- fats
 - oils
 - enzymes
 - amino acids
- Q3.** Which of the following is an amine?
- $$\begin{array}{c} \text{H} \quad \text{H} \quad \text{O} \\ | \quad | \quad || \\ \text{H}-\text{C}-\text{N}-\text{C}-\text{H} \\ | \\ \text{H} \end{array}$$
 - $$\begin{array}{c} \text{H} \quad \text{H} \\ | \quad | \\ \text{H}-\text{C}-\text{N}-\text{H} \\ | \\ \text{H} \end{array}$$
 - $$\begin{array}{c} \text{H} \\ | \\ \text{H}-\text{C}-\text{N}=\text{C}=\text{O} \\ | \\ \text{H} \end{array}$$
 - $$\begin{array}{c} \text{H} \quad \text{H} \\ | \quad | \\ \text{H}-\text{C}=\text{C}-\text{C}\equiv\text{N} \end{array}$$
- Q4.** The arrangement of amino acids in a peptide is Z-X-W-V-Y where the letters V, W, X, Y and Z represent amino acids. On partial hydrolysis of the peptide, which of the following sets of dipeptides is possible?
- V-Y, Z-X, W-Y, X-W
 - Z-X, V-Y, W-V, X-W
 - Z-X, X-V, W-V, V-Y
 - X-W, X-Z, Z-W, Y-V

- Q5.** Which of the following arrangements of atoms shows a peptide link?

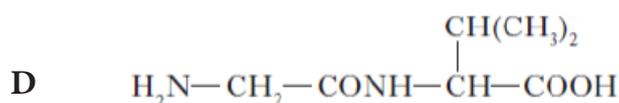
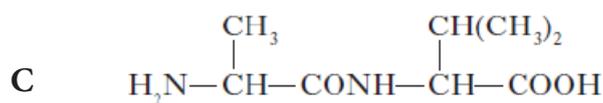
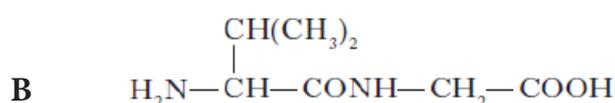
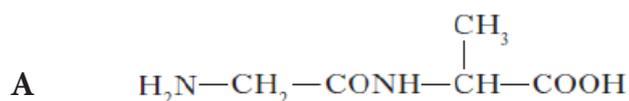


- Q6.** A tripeptide X has the structure



Partial hydrolysis of X yields a mixture of dipeptides.

Which of the following dipeptides could be produced on hydrolysing X?



HOME PRACTICE

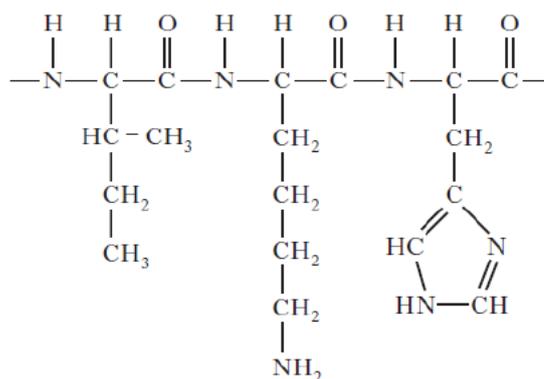
6.7

Q1. Chemists have developed cheeses specifically for use in cheeseburgers. When ordinary cheddar cheese is grilled the shapes of the protein molecules change and the proteins and fats separate leaving a chewy solid and an oily liquid.

- a)** What name is given to the change in protein structure which occurs when ordinary cheddar is grilled? 1

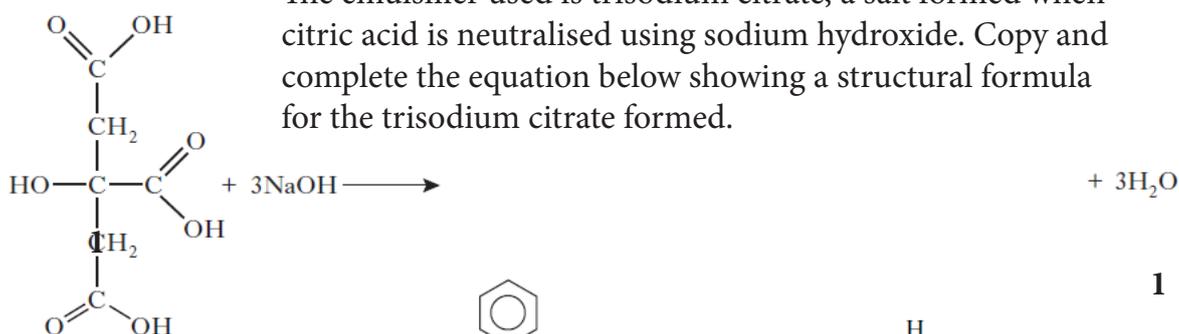
To make cheese for burgers, grated cheddar cheese, soluble milk proteins and some water are mixed and heated to no more than 82 °C. As the cheese begins to melt an emulsifying agent is added and the mixture is stirred.

- b)** Why would a water bath be used to heat the mixture? 1
- c)** A section of the structure of a soluble milk protein is shown.



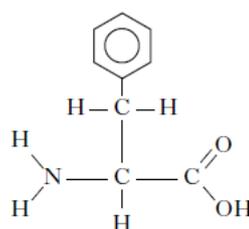
Draw a structural formula for any one of the amino acids formed when this section of protein is hydrolysed.

- d)** The emulsifier used is trisodium citrate, a salt formed when citric acid is neutralised using sodium hydroxide. Copy and complete the equation below showing a structural formula for the trisodium citrate formed. 1

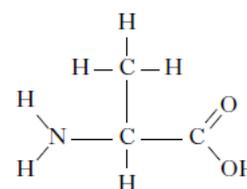


Q2. Phenylalanine and alanine are both amino acids.

Phenylalanine is an essential amino acid.

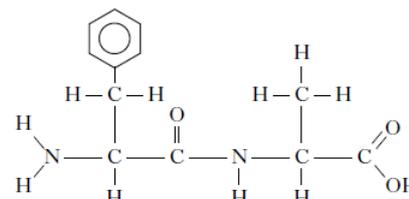


phenylalanine



alanine

- a)** What is meant by an essential amino acid? 1
- b)** How many hydrogen atoms are present in a molecule of phenylalanine? 1
- c)** Phenylalanine and alanine can react to form the dipeptide shown.
- Copy and circle the peptide link in this molecule. 1
- d)** Draw a structural formula for the other dipeptide that could be formed. 1



Total (8)

Context - Kitchen Chemistry

This activity is to remind you how much of the Organic Chemistry met in this Unit has been taught within the context of Kitchen Chemistry

- Flavour** Most of our 'tasting' is done through our noses, so most flavour molecules are *volatile (weak intermolecular forces)* such as
- esters** - many have sweet fruity smells - e.g. 'pear drops' - *pentyl ethanoate*
 - aldehydes** - can however be **oxidised** over time resulting in (sometimes unpleasant) changes in flavour
 - terpenes** - 'essential oils' are responsible for many distinctive flavours such as *cinnamon* and *ginger* as well as *oranges* and *lemons*.
 - acids** - such as *vinegar*, *ethanoic acid* are used to introduce 'sour' flavours
- Cooking** Cooking can *dissolve* out the flavour molecules so whilst many foods can be safely *cooked in water*, others are better *cooked in oil*.
- Cooking also effects the structure of, in particular, **proteins** which can result in significant change in the *texture* of certain foods during cooking.
- Texture** **Protein** structure has an important impact on the *texture* of food and can be effected by *changes in temperature (cooking)* and *changes in pH (marinading in acids such as vinegar, ethanoic acid)*.
- Foods, such as curries, often 'separate' into 'water layer' and 'oil layer' as a result of *differences in solubility (different intermolecular forces)*.
- Colour** Colour can arise in many ways but larger **terpenes** are often responsible for the *yellow, orange or red* colour of so many foods - e.g. *carotene* in carrots.
- Energy** Whilst **carbohydrates**, our main 'energy food', are not covered in this course, **proteins** also contribute energy whilst *fats & oils* are our most concentrated source of energy.
- Additives** **antioxidants** - can be added to food, though many foods already contain *Vitamin C* - a natural *antioxidant*
- emulsifying agents** - are used to prevent 'water layers' and 'oily layers' from separating. These molecules are often made from *fats & oils* and behave in exactly the same way as *soap molecules*. *Milk* contains natural **emulsifiers**.

In addition, many reactions triggered by *UV light*, such as the **substitution** reaction between *alkanes* and *halogens*, involve the production of **free radicals**. **Free radicals** start **chain reactions** which are responsible for the ageing of your skin. *Sun cream* contains chemicals designed to absorb UV before it reaches your skin.

Vitamin E and *melatonin* are natural **free radical scavengers** that can help counter the effect of free radicals. Many **cosmetic** products contain *free radical scavengers* which react with **free radicals** to form stable molecules and prevent **chain-reactions** starting.