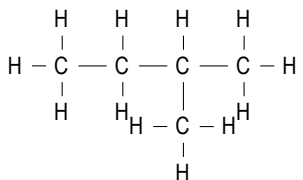
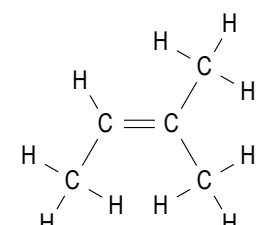
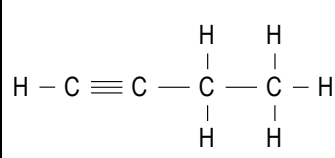
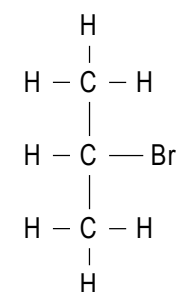
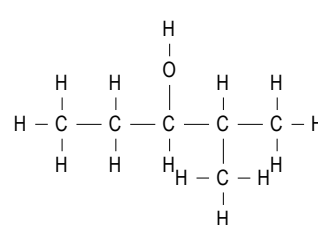
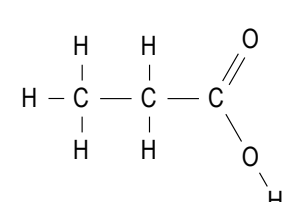
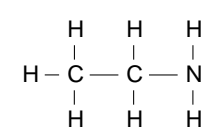


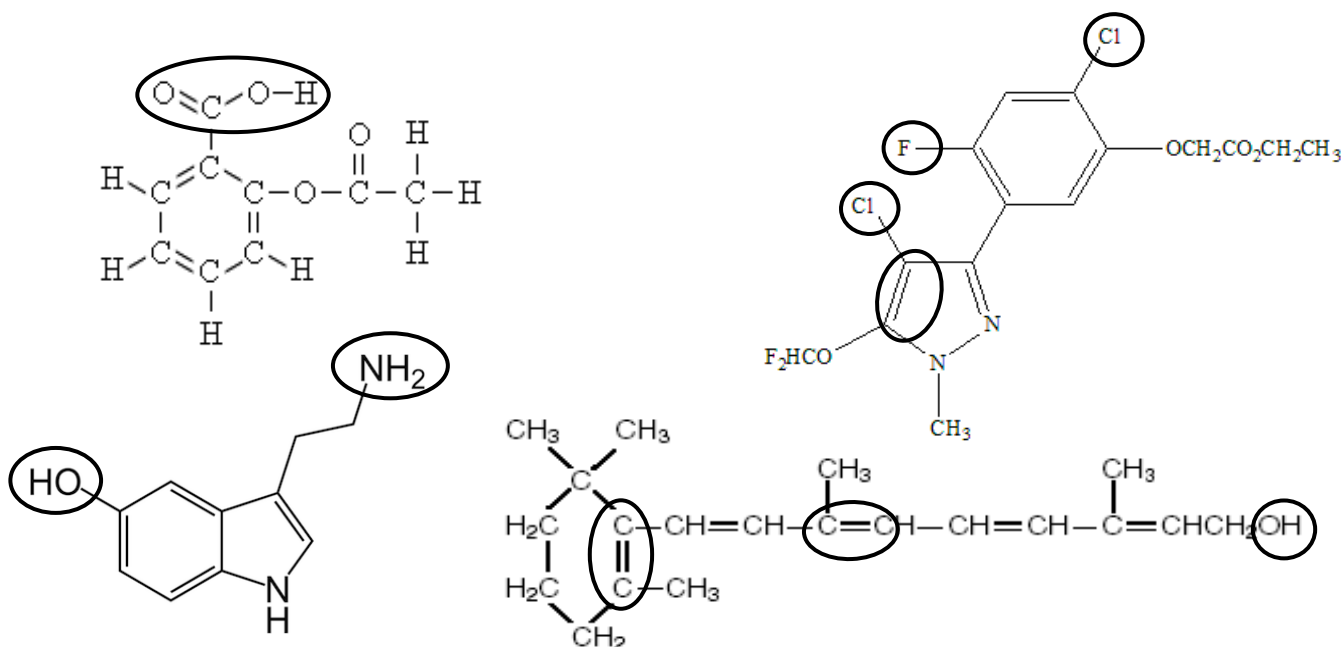
## LEVEL 2 ORGANIC CHEMISTRY

### Naming (IUPAC nomenclature)

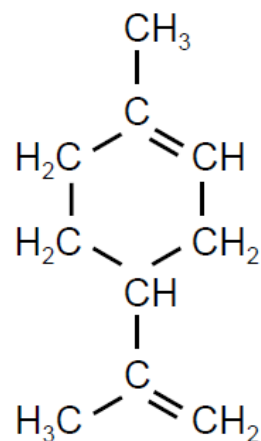
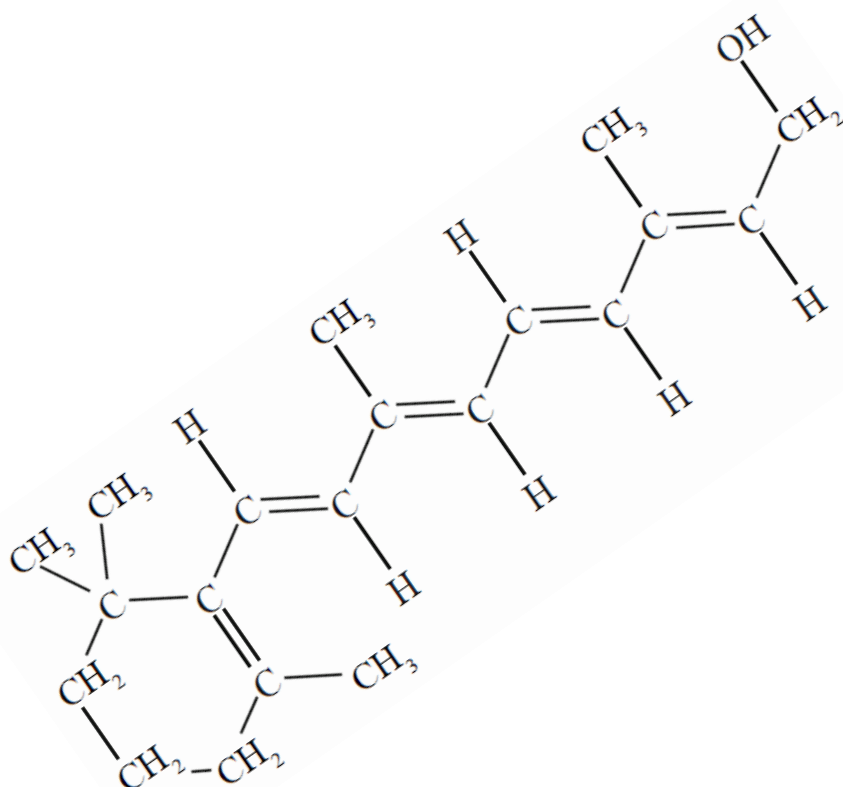
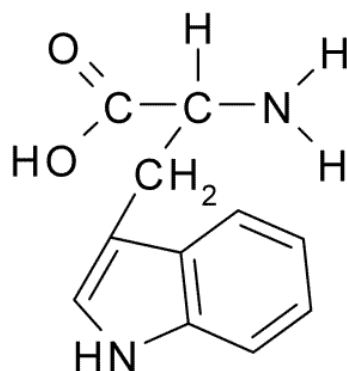
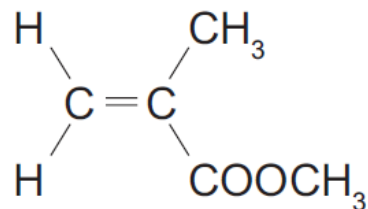
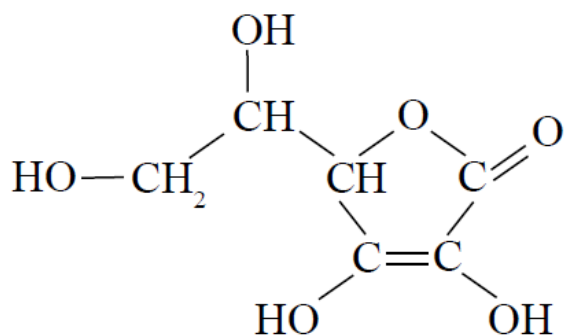
Place in a molecule where a reaction takes place. Compounds with the same functional group – homologous series - have similar properties.

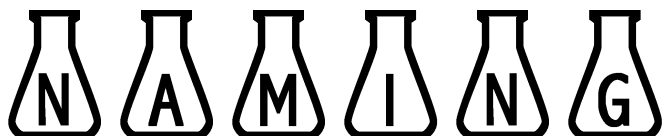
First the functional groups...

alkane -ane	alkene -ene	alkyne -yne	haloalkane bromo, chloro etc
			
alcohol -anol	carboxylic acid -anoic acid	amine amino	C-C C=C C≡C R-X R-OH R-COOH R-NH <sub>2</sub>
			

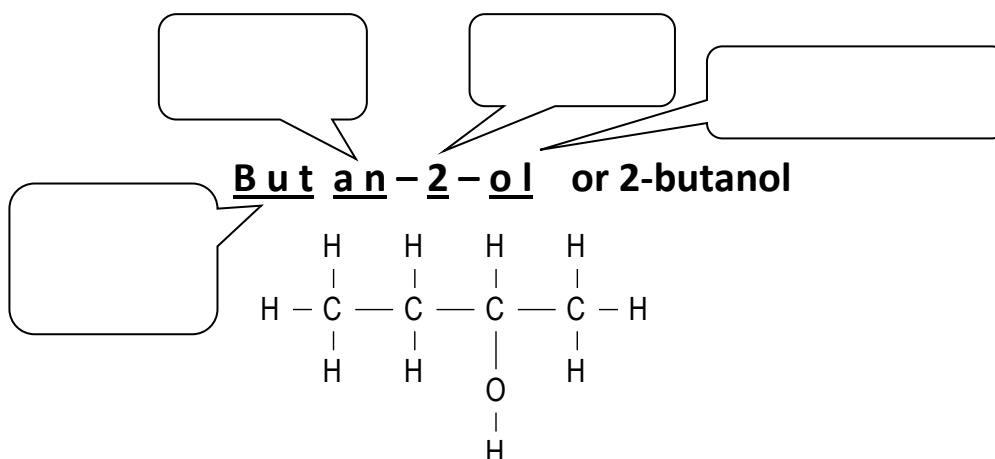


What functional groups can you identify here?

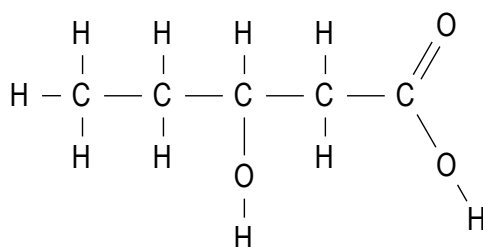




No. of carbons	Alkane name	Prefix	Side chain
1	methane	meth-	methyl-
2	ethane	eth-	ethyl-
3	propane	prop-	propyl-
4	butane	but-	butyl-
5	pentane	pent-	
6	hexane	hex-	
7	heptane	hept-	
8	octane	oct-	



- Find longest carbon chain – base name on the parent alkane
- Number the C atoms in the chain to indicate position of any side chains or functional groups (number from end to give the lowest numbers)
- Give names and positions of functional groups eg 3-bromo, and 4-chloro (alphabetically)
- Where there's more than one functional group there is priority order for naming - ol > halo- -anoic acid > halo -anoic acid > hydroxy (-OH)



\_\_\_ - hydroxyl \_\_\_\_\_anoic acid

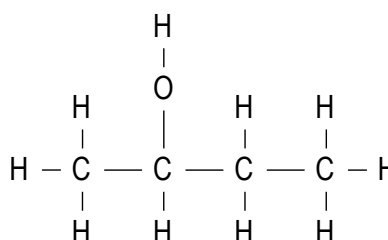
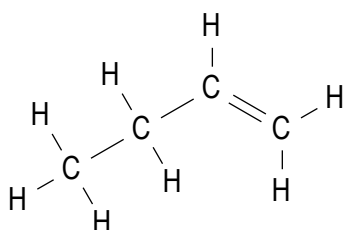
Name me...	Draw me...
$\text{CH}_3\text{CH}_2\text{C}\equiv\text{CCH}_3$	<p style="text-align: center;"><b>3-chlorobutan-1-ol</b></p>
$\begin{array}{cccc} \text{H} & \text{H} & \text{H} & \text{H} \\   &   &   &   \\ \text{H}-\text{C} & -\text{C} & -\text{C} & -\text{N} \\   &   &   &   \\ \text{H} & \text{H} & \text{H} & \text{H} \end{array}$	<p style="text-align: center;"><b>2-chloro-3-methylbutanoic acid</b></p>
$\begin{array}{ccccccc} \text{CH}_3 & -\text{CH}_2 & -\text{CH} & -\text{CH}_2 & -\text{Cl} \\ & &   & & \\ & & \text{Cl} & & \end{array}$	<p style="text-align: center;"><b>2-methyl propan-1,2-diol</b></p>
$\begin{array}{ccccccc} \text{H} & \text{H} & \text{H} & \text{H} & & & \\   &   &   &   & & & \\ \text{N} & -\text{C} & -\text{C} & -\text{C} & -\text{H} & & \\   &   &   &   & & & \\ \text{H} & \text{H} & \text{H} & \text{H} & & & \end{array}$	<p style="text-align: center;"><b>pentan-2-ol</b></p>
$\begin{array}{ccccccc} \text{CH}_3 & -\text{CH}_2 & -\text{CH} & -\text{CH}_2 \\ & &   &   \\ & & \text{Br} & \text{Br} \end{array}$	<p style="text-align: center;"><b>4-methylpent-2-yne</b></p>
$\begin{array}{ccc} \text{CH}_3 & -\text{C} & =\text{CH}_2 \\ &   & \\ & \text{Br} & \end{array}$	<p style="text-align: center;"><b>3-chloropropan-1-ol</b></p>
$\begin{array}{cccc} \text{H} & \text{H} & \text{H} & \text{H} \\   &   &   &   \\ \text{H}-\text{C} & -\text{C} & -\text{C} & -\text{N} \\   &   &   &   \\ \text{H} & \text{H} & \text{H} & \text{H} \end{array}$	<p style="text-align: center;"><b>2-amino-2-methyl propane</b></p>
$\begin{array}{cccc} \text{H} & \text{H} & \text{H} & \text{H} \\   &   &   &   \\ \text{H}-\text{C} & -\text{C} & -\text{C} & -\text{C}-\text{H} \\   &   &   &   \\ \text{H} & \text{H} & \text{H}-\text{C}-\text{H} & \text{H} \\ & &   & \\ & & \text{H} & \end{array}$	<p style="text-align: center;"><b>pent-2-ene</b></p>

# FORMULAE

General Formula  
 $C_nH_{2n+2}$  alkanes  
 $C_nH_{2n}$  alkenes  
 $C_nH_{2n-2}$  alkynes  
 $C_nH_{2n+1}OH$  alkanols (alcohols)  
 $C_nH_{2n+1}Cl$  chloroalkanes  
 $C_nH_{2n+1}NH_2$  amines  
 $C_nH_{2n+1}COOH$  carboxylic acids

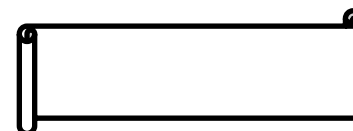
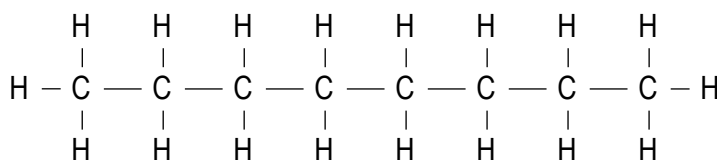
**Molecular formula** – gives the number of each type of atom in a molecule

**Structural formula** – gives the arrangement of atoms in a molecule, indicating how atoms are bonded together.

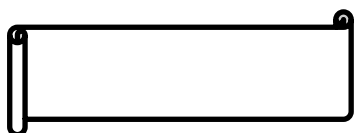


• Expanded

• Condensed



is  $CH_3CH_2CH_2CH_2CH_2CH_2CH_2CH_3$  or  $CH_3(CH_2)_6CH_3$





## Structural isomers

- **Same number and type of atoms but arranged in different ways**

*Structural isomers have the same molecular formula but they differ in the sequence in which the atoms are joined together.*

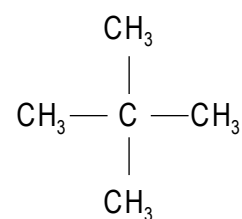
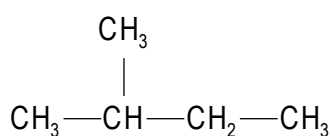
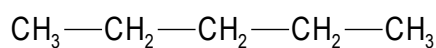
### Structural

- Chain/branched chain
- Positional
- Functional group

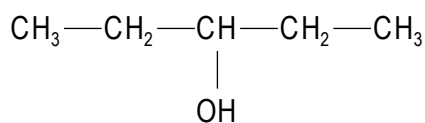
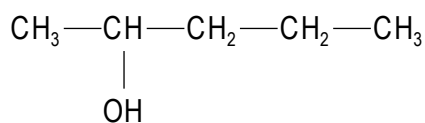
### Geometric

- Cis
- Trans

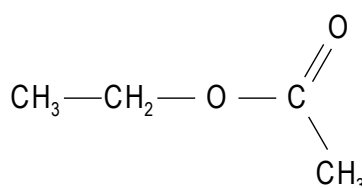
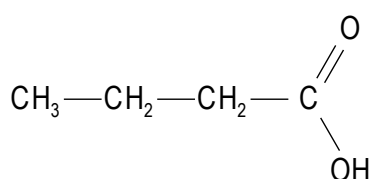
### *Chain/branched chain*



### *Positional*



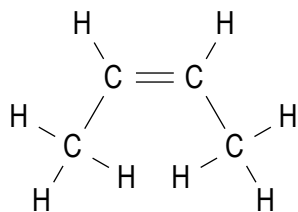
### *Functional group*



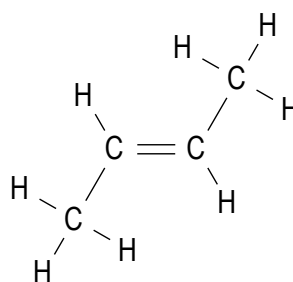
- **Geometric**

C=C double bond in alkenes is fixed and cannot be rotated (“is no free rotation about the C=C double bond”). This allows for different arrangements of the atoms/groups of atoms **in space**.

- groups on same side, cis-
- groups on opposite sides, trans

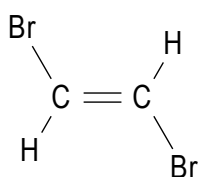


*cis-*

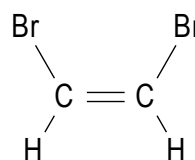


and *trans-*

*Cis–trans* (geometric) isomers exist where there is a C=C which **cannot freely rotate**. If there are two different groups bonded to the Cs of the double bond, two arrangements are possible.



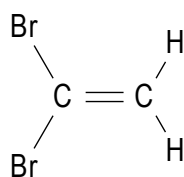
*trans-*



and *cis-*

1,2–dibromoethene meets these requirements since each C of the double bond has –H and –Br, i.e. different groups.

However, 1,1–dibromoethene does not meet these requirements since the two groups on the Cs of the double bond are the same, ie one C has two –H and the other two –Br.



### Example Question

But–2–ene can exist as geometric (*cis–trans*) isomers, whereas but–1–ene cannot. Explain this difference, using structural formulae to illustrate your answer.

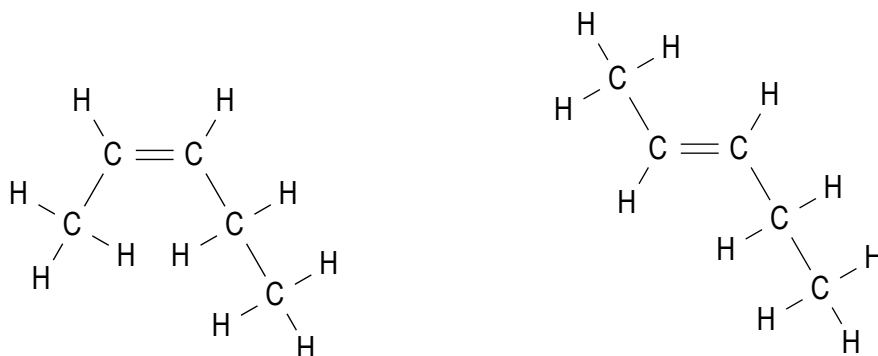
*Cis–trans* isomers can occur in molecules that have double bonds, because rotation of the atoms about the axis of the carbon to carbon double bond is restricted. They must also have two different groups attached to each of the carbons involved in the double bond.

Cis-but-2-ene	Trans-but-2-ene	But-1-ene

# Drawing cis- and trans- isomers

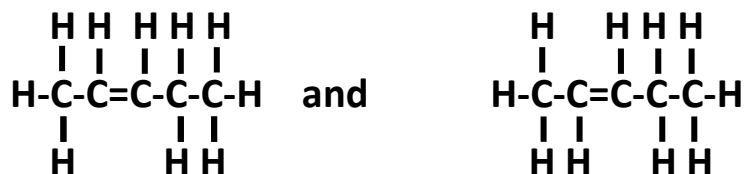
Always start from a  $\begin{array}{c} \diagup \\ \text{C} \\ \diagdown \end{array} = \begin{array}{c} \diagdown \\ \text{C} \\ \diagup \end{array}$  shape!!

Right 😊



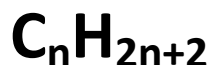
and

Wrong 😞



Using the top way, you will clearly see the effect on the shape of the molecule.

## ALKANES



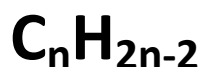
- single C-C bonds
- are saturated (no more H atoms can be added to their molecules)

## ALKENES



- contain a C=C double bond
- are unsaturated (more H atoms (or other atoms) can be added to their molecules)

## ALKYNES



- contain a C≡C triple bond
- are unsaturated (more H atoms (or other atoms) can be added to their molecules)

## *Physical properties*

- as no. of C↑ go from being gas to liquid to solid at room temperature
- smell – have weak intermolecular forces
- low m.pt & b.pt – have weak intermolecular forces
- insoluble in water – hydrocarbons are non-polar
- good solvents for fats & oils – non-polar substances dissolve other non-polar substances
- don't conduct electricity – no electrons that are free to move

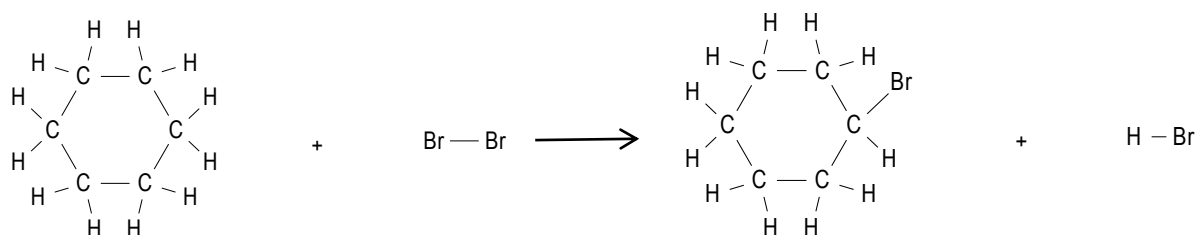


# ALKANES

- Fairly unreactive (only C-C); reactions need heat and/or UV light.
- But they do BURN well – used as fuels ☺ see below

*Substitution reactions:* chlorination / bromination.

- React in presence of UV light and/or heat.



- The reaction is slow. Orange bromine is slowly decolourised.
- Chlorine reacts in a similar way.
- The reaction continues but normally we only write equations for “monosubstitution”.
- It is called a SUBSTITUTION reaction because one of the hydrogen atoms in the molecule is replaced by a bromine atom.

Molecules that undergo substitution reactions have carbon to carbon single bonds and form molecules with carbon to carbon single bonds. In a substitution reaction an atom or group of atoms is replaced by another atom or group of atoms.

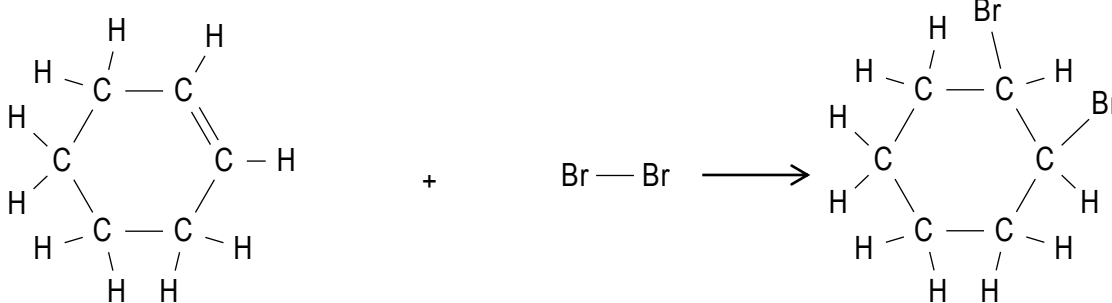
# Combustion

- Complete; plentiful O<sub>2</sub>. Products CO<sub>2</sub> & H<sub>2</sub>O and lots of energy
- Incomplete; limited O<sub>2</sub>. Products C (soot), CO & CO<sub>2</sub> & H<sub>2</sub>O and less energy

# ALKENES...

- More reactive than alkanes because of C=C
- Undergo ADDITION reactions
- C=C double bond replaced with C-C bond & 2 new bonds are made.

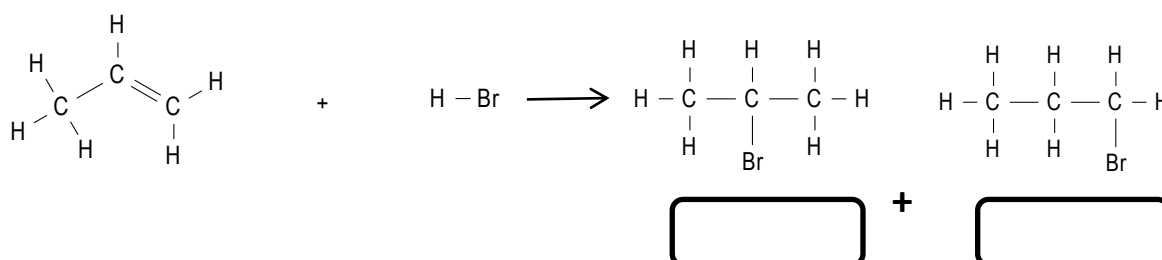
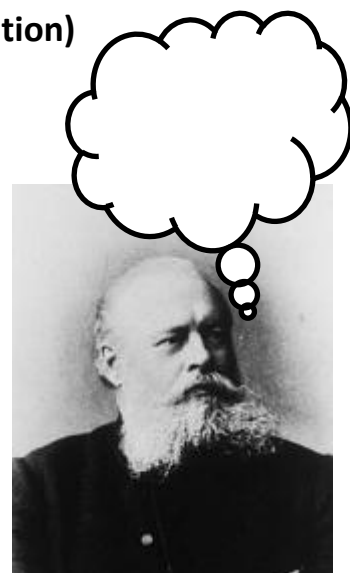
Molecules that undergo **addition** reactions have carbon to carbon double bonds and form molecules with carbon to carbon single bonds. In an addition reaction the reaction involves breaking a double bond between the carbon atoms and forming a single bond in its place as well as forming two new single bonds.



- Addition reactions can occur with
  - Hydrogen, H<sub>2</sub>, Pt catalyst – hydrogenation
  - Water, conc H<sub>2</sub>SO<sub>4</sub> then water & heat – hydration
  - Halogens, Cl<sub>2</sub> & Br<sub>2</sub> – halogenation (chlorination, bromination)
  - Hydrogen halides, HBr – hydrohalogenation
  - Themselves, monomer → polymer, polymerisation

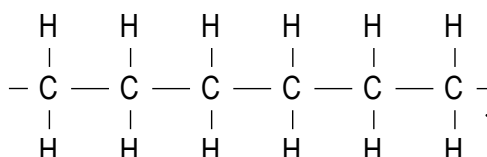
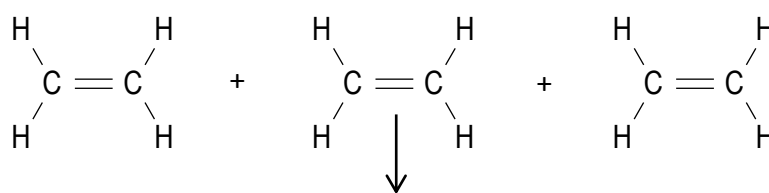
## Markownikoff's rule

- Addition of an unsymmetrical reagent (eg HCl, HBr, H<sub>2</sub>O (H-OH)) to an unsymmetrical alkene eg propene or but-1-ene
- 2 possible products, one major one minor
- Predict using "the rich get richer"
- H atom adds to the C of the C=C that already has most H atoms

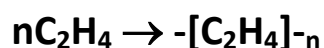


# POLYMERISATION

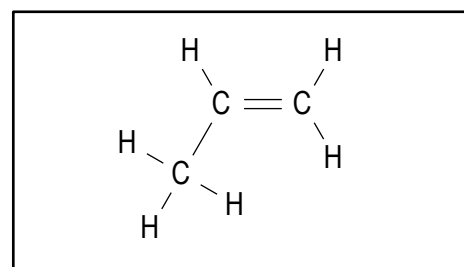
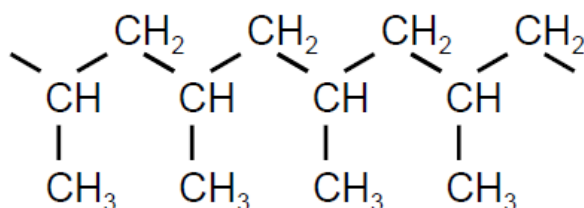
- Need high temperatures, pressure & catalyst
- Many monomer molecules → polymer molecule
- Addition reaction – called “addition polymerisation”
- E.g. ethene → polyethene, propene → polypropene
- Feature that allows this is the C=C double bond



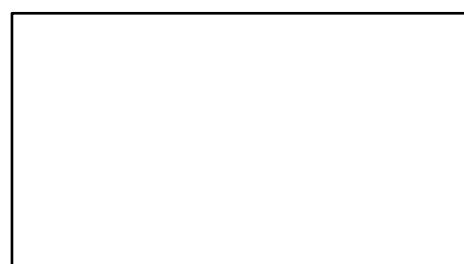
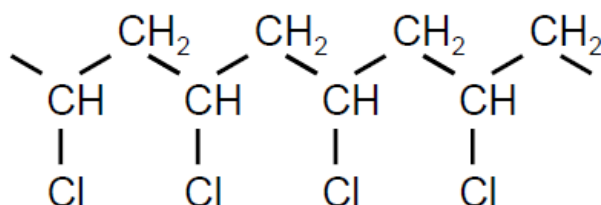
Now a saturated molecule but still called polyethene



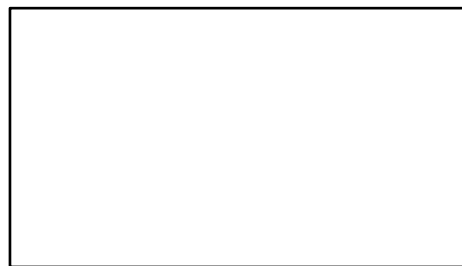
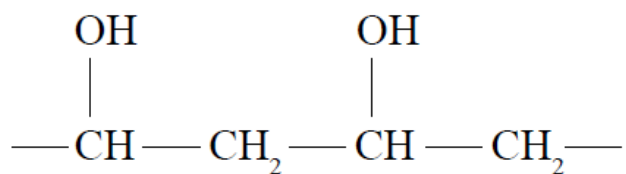
## Polypropene



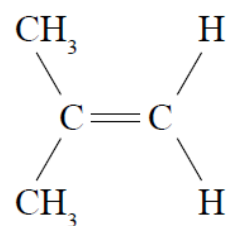
## Polyvinyl chloride (PVC)



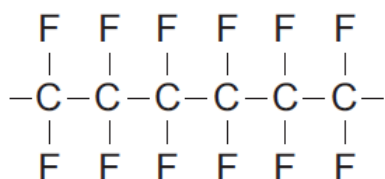
### Polyvinyl alcohol



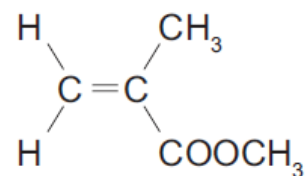
### Polymethylpropene



### Polytetrafluoroethene (PTFE)



### Perspex





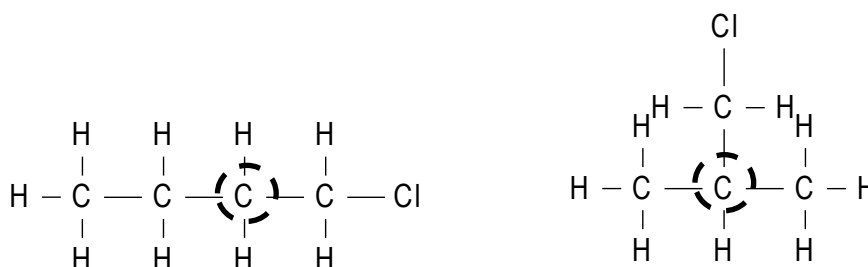
- R-X
- $C_nH_{2n+1}X$
- X – is F, Cl, Br or I (halogen)

### Formed by

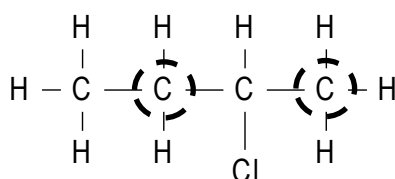
- Substitution of an alkane (needs *uv* light and or heat)
- Addition of HX to an alkene
- Addition of  $X_2$  to an alkene
- Reaction of an alcohol with  $PCl_3$ ,  $PCl_5$  or  $SOCl_2$

### Classification – based on position of the halogen group

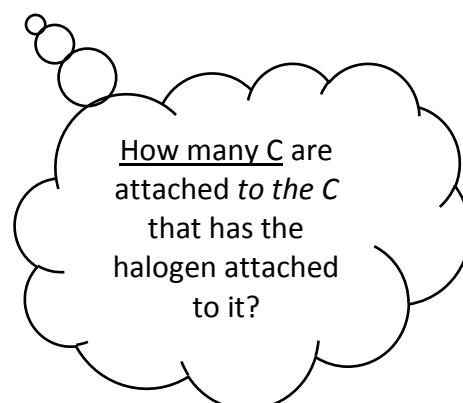
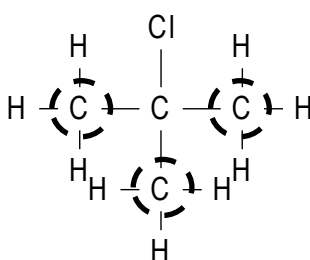
- Primary



- Secondary



- Tertiary



# Reactions of haloalkanes

## Substitution reactions

$\text{NH}_3$ : haloalkane  $\rightarrow$  amine

$\text{R-X(l)} + \text{NH}_3(\text{alc}) \rightarrow \text{R-NH}_2 + \text{HX(alc)}$ ; they must be dissolved in alcohol, not in  $\text{H}_2\text{O}$  (which would produce  $\text{NH}_4^+$  and  $\text{OH}^-$ :  $\text{OH}^-$  would react with the  $\text{R-X}$  instead of the  $\text{NH}_3$ ).

$\text{KOH}$  or  $\text{NaOH(aq)}$ : haloalkane  $\rightarrow$  alcohol

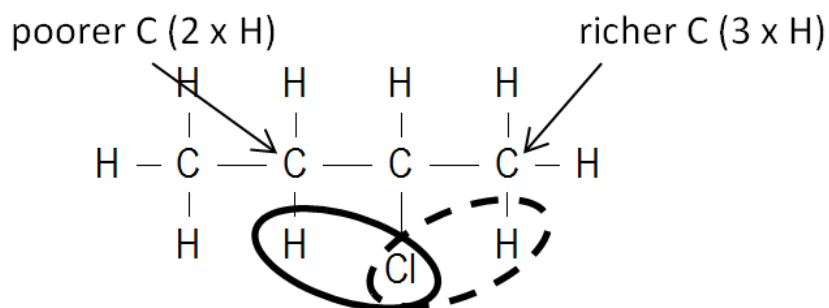
$\text{R-X(l)} + \text{NaOH(aq)} \rightarrow \text{R-OH} + \text{NaX(aq)}$

## Elimination reactions:

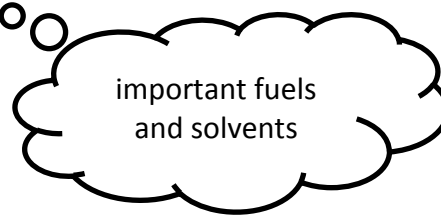
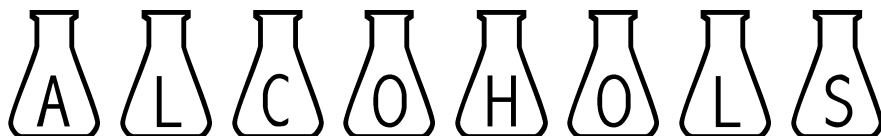
Haloalkanes  $\rightarrow$  alkenes,  $\text{R-X} \rightarrow \text{R=C} + \text{HX}$

This uses  $\text{KOH}$  dissolved in alcohol, usually ethanol which we write as  $\text{KOH(alc)}$  or  $\text{OH}^-(\text{alc})$

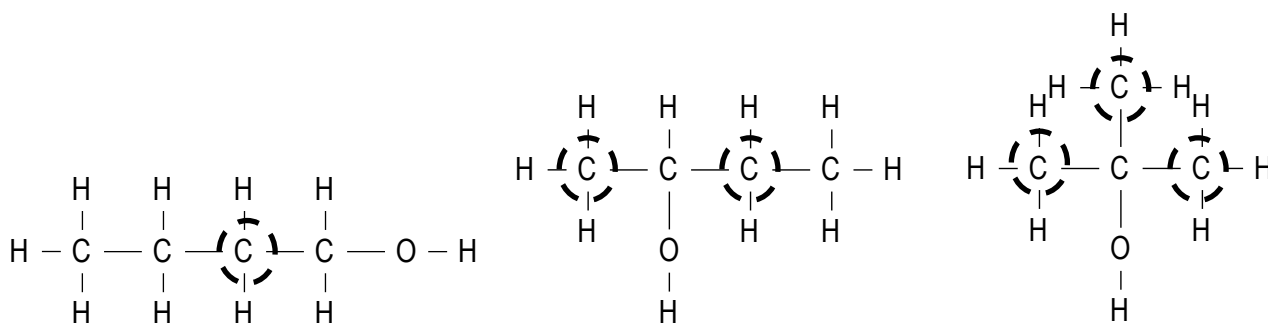
- $\text{KOH(alc)}$  or  $\text{NaOH(alc)}$  - dissolved in alcohol to prevent substitution by  $\text{OH}^-$  (forming the alcohol)
- $3^\circ$  haloalkanes  $>$   $2^\circ$  haloalkanes  $>$   $1^\circ$  haloalkanes to undergo elimination
- **Saytzeff's rule** applies: *the poor get poorer*. (reverse of Markovnikov's rule)
- Will get a major & minor products if unsymmetrical/asymmetric haloalkane



Molecules that undergo elimination reactions have carbon to carbon single bonds and form molecules with carbon to carbon double bonds. In an elimination reaction two atoms or small groups are removed from a molecule forming a carbon to carbon to double bond.

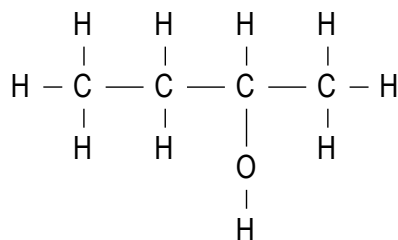


- -OH functional group
- $C_nH_{2n+1}OH$  or R-OH
- Position of -OH group decides primary, secondary or tertiary

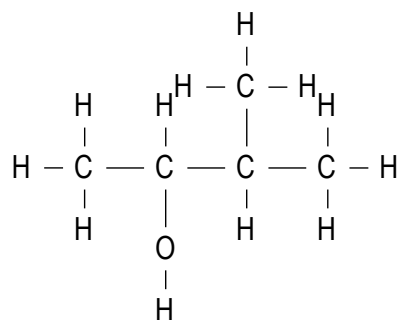


- Polar molecules due to -OH group; small ones (C 1 -3) are water soluble, larger ones not.

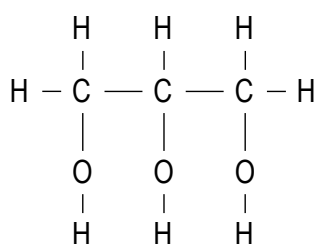
### Naming - name or draw



propan-2-ol

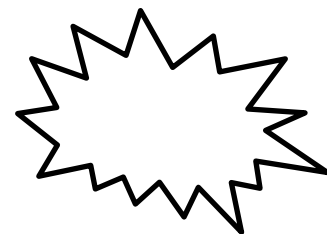


ethan-1,2-diol



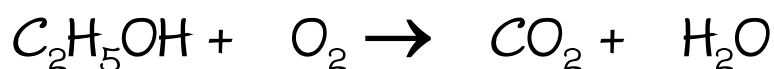
2-chloro-butan-2-ol

# Reactions of alcohols



## Combustion

Burn with a clean almost colourless flame, good fuels



## OXIDATION

Primary alcohols can be oxidised by heating them with either:

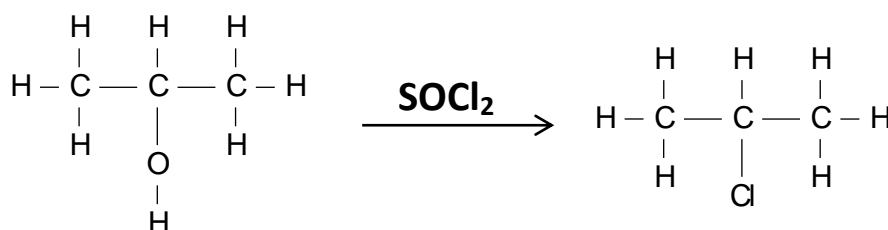
- Acidified dichromate ( $H^+/Cr_2O_7^{2-}$ ):  
colour change  $Cr_2O_7^{2-}$  \_\_\_\_\_ to  $Cr^{3+}$  \_\_\_\_\_
- Acidified permanganate ( $H^+/MnO_4^-$ ):  
colour change  $MnO_4^-$  \_\_\_\_\_ to  $Mn^{2+}$  \_\_\_\_\_

The alcohols are oxidised to carboxylic acid ( $-COOH$  functional group)

## SUBSTITUTION REACTIONS – to form chloroalkanes

Chloroalkanes are important for the formation of many other organic chemicals.

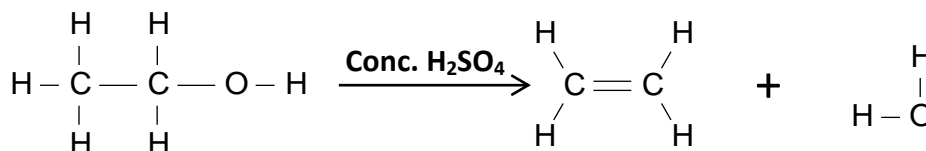
- Can use  $PCl_3$ ,  $PCl_5$  or  $SOCl_2$  ( $SOCl_2$  being the most effective for primary, secondary & tertiary alcohols).



(+  $SO_2$  +  $HCl$ )

# Elimination of water – essentially dehydration where –OH is removed with –H from an adjacent C atom – to produce a C=C (alkene)

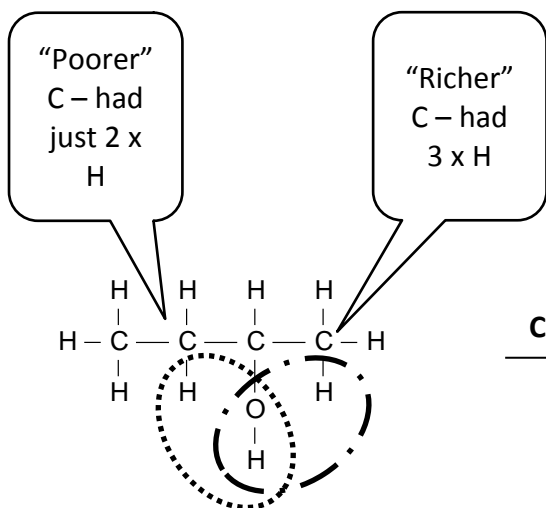
Reagent: conc H<sub>2</sub>SO<sub>4</sub>



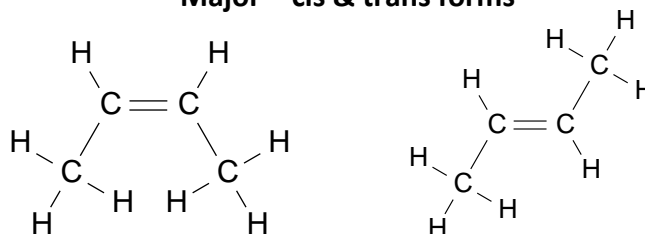
Here, we may need to apply the **Saytzeff's rule**:

the poor get poorer if the alcohol is **ASSYMETRIC**. The double bond starts on the carbon which had the –OH, but the other end of that double bond goes to the carbon atom which had the fewer hydrogen atoms.

e.g. butan-2-ol

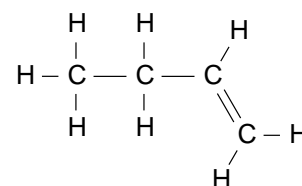


Major – cis & trans forms



Conc. H<sub>2</sub>SO<sub>4</sub>

Minor



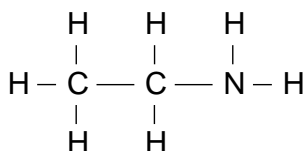
Pentan-3-ol  $\begin{array}{c} \text{H} & \text{H} & \text{H} & \text{H} & \text{H} \\ | & | & | & | & | \\ \text{H}-\text{C} & -\text{C} & -\text{C} & -\text{C} & -\text{C}-\text{H} \\ | & | & | & | & | \\ \text{H} & \text{H} & \text{O} & \text{H} & \text{H} \\ & & | & & \\ & & \text{H} & & \end{array}$  would only form one product, pent-2-ene



# AMINES –NH<sub>2</sub> amino group

Essentially NH<sub>3</sub> with 1, 2 or 3 H atoms replaced with alkyl group(s)

- naming based on alkane so C<sub>2</sub>H<sub>5</sub>NH<sub>2</sub> is aminoethane (IUPAC) but also often called ethylamine

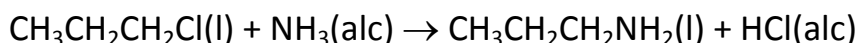


## Physical properties

- Methylamine and ethylamine are gases; others low m.pt. volatile liquids; C > 5 are usually solids
- Characteristic “fishy” smell (actually they stink!!)  
Low mass amines are soluble in water – as form hydrogen bonds with water; larger ones insoluble.




## Preparation

Primary amines can be prepared by the substitution of a haloalkane by alcoholic ammonia.



## Acid-base reactions of amines.

- Amines are bases - like ammonia.
- Amines are **weak bases** since proton acceptors,  $\text{R-NH}_2 + \text{H}_2\text{O} \rightleftharpoons \text{R-NH}_3^+ + \text{OH}^-$ 
  - $\text{NH}_3 + \text{H}_2\text{O} \rightleftharpoons \text{NH}_4^+ + \text{OH}^-$
  - $\text{R-NH}_2 + \text{H}_2\text{O} \rightleftharpoons \text{R-NH}_3^+ + \text{OH}^-$

They turn damp  litmus paper   
and green universal indicator solution/paper blue. 

### Reaction with HCl

As bases, amines react with acids such as HCl to form salts.

- $\text{NH}_3(\text{g}) + \text{HCl}(\text{g}) \rightarrow \text{NH}_4^+\text{Cl}^-(\text{s})$
- $\text{CH}_3\text{NH}_2(\text{g}) + \text{HCl}(\text{g}) \rightarrow \text{CH}_3\text{NH}_3^+\text{Cl}^-(\text{s})$  methyl ammonium chloride

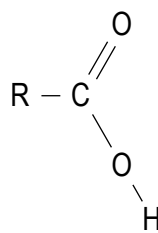
These salts are colourless, crystalline solids, soluble in water and have no smell.

## Carboxylic acids – count the “C”s ☺

HCOOH methanoic acid





CH<sub>3</sub>COOH ethanoic acid

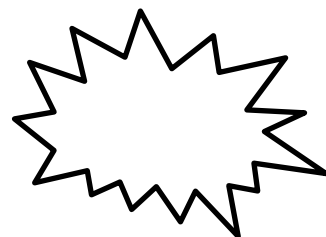
C<sub>2</sub>H<sub>5</sub>COOH propanoic acid



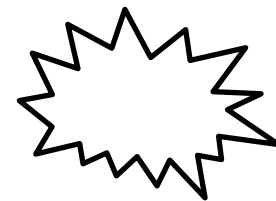
- contain the –COOH functional group
- the -COOH group is very polar; C1-3 are soluble in water, C4 and above not (due to long hydrocarbon portion)
- Higher m.pt. and b.pt. than alcohols of similar mass due to stronger intermolecular attractions
- are weak acids – only partly ionised when placed in water
 
$$\text{CH}_3\text{COOH} + \text{H}_2\text{O} \rightleftharpoons \text{CH}_3\text{COO}^- + \text{H}_3\text{O}^+$$
- are poor conductors of electricity as aq. solutions (as weak acids)

## Reactions as acids

- turn  blue litmus  red
- turn  green UI paper  orange pH 3-4
- $\text{RCOOH} + \text{H}_2\text{O} \rightleftharpoons \text{RCOO}^- + \text{H}_3\text{O}^+$  ie release  $\text{H}_3\text{O}^+$  ions
- “Typical acids reactions”
  - acid + metal → salt + hydrogen eg Mg or Zn to produce bubbles of H<sub>2</sub> gas
  - acid + base → salt + water eg NaOH
  - acid + carbonate → salt + water + carbon dioxide eg Na<sub>2</sub>CO<sub>3</sub> and CaCO<sub>3</sub> & hydrogen carbonates NaHCO<sub>3</sub> to produce bubbles of CO<sub>2</sub> gas



# TYPES OF REACTIONS



The list may look long & bewildering but that's just because some reactions can be called a number of things.....

All of this has been previously covered in separate sections. The "must knows" are highlighted

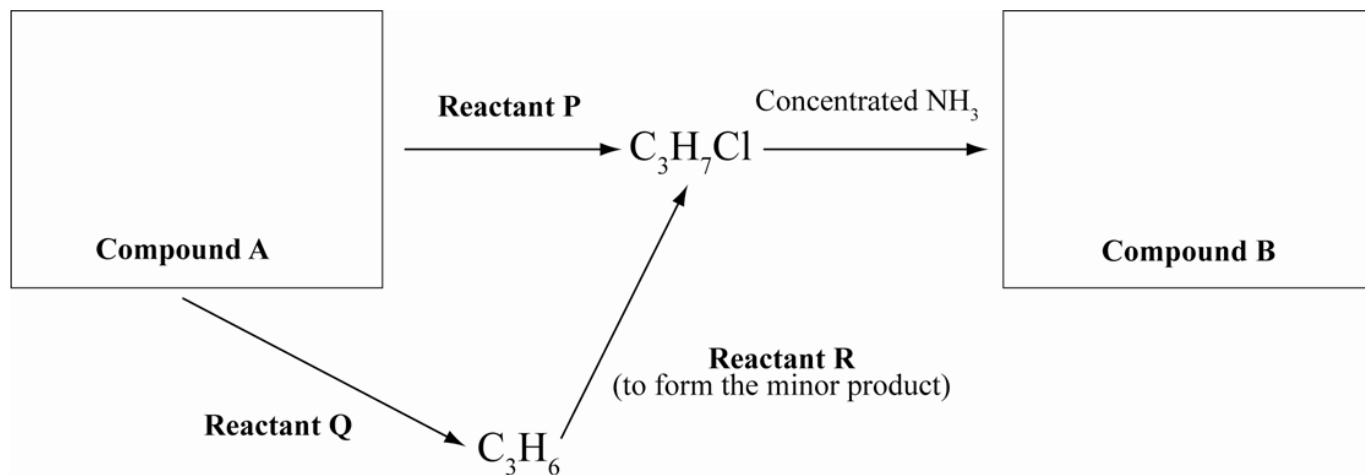
- **acid-base**
  - reaction involving a carboxylic acid & a base e.g. NaOH OR
  - reaction involving a carboxylic acid & a carbonate/hydrogen carbonate e.g. NaHCO<sub>3</sub>
  - reaction involving an amine & an acid
  - these are neutralisation reactions
  
- **addition**
  - involves a small molecule joining across the C=C double bond of an unsaturated molecule (alkene/yne). The molecule becomes saturated.
  - no other product is made
  
- **bromination**
  - addition of bromine
  
- **chlorination**
  - addition of chlorine
  
- **dehydration**
  - the removal of water (H and OH on adjacent C atoms) to form a C=C bond
  - heat with conc. H<sub>2</sub>SO<sub>4</sub> or pass over Al<sub>2</sub>O<sub>3</sub> catalyst
  - since it involves removal it is also an elimination reaction
  
- **elimination**
  - -H & -OH / water / removed from neighbouring C atoms in an alcohol. Also known as dehydration
  - H-X / hydrogen halide removed from neighbouring C atoms in a haloalkane.

A C=C double bond forms / forms an alkene / the molecule becomes unsaturated.

- **halogenation** (halogens are  $\text{Cl}_2$ ,  $\text{Br}_2$  etc)
  - halogen added across the double bond of alkene (or  $\text{C}\equiv\text{C}$  of alkyne)
  - since it involves adding atoms it is also an addition reaction
  - chlorination, bromination & iodination are all halogenation reactions**OR**
  - halogen swapped for an H on an alkane (alkane & halogen & UV light and/or heat)
  - since it involves replacing atoms it is also a substitution reaction
- **hydration**
  - water added across the double bond of alkene (alkene  $\rightarrow$  alcohol)
  - since it involves adding atoms H and OH it is also an addition reaction
- **hydrogenation**
  - hydrogen added across the double bond of alkene (or  $\text{C}\equiv\text{C}$  of alkyne)
  - needs a Pt or Ni catalyst
  - since it involves adding atoms it is also an addition reaction
- **oxidation**
  - conversion of primary alcohol to carboxylic acid using heat and  $\text{H}^+/\text{Cr}_2\text{O}_7^{2-}$  or  $\text{H}^+/\text{MnO}_4^-$  (both oxidising agents)
  - conversion of an alkene to a diol using  $\text{H}^+/\text{MnO}_4^-$  (no heat needed)
- **polymerisation** (limited to addition polymerisation @ level 2)
  - unsaturated monomers joined to make a polymer (saturated)
  - it is an addition reaction as the monomers add (and no other product made)
- **substitution** – one atom is removed and replaced with another atom
  - e.g.  $\text{Cl}_2$  and an alkane: one hydrogen atom will be removed from the molecule and one chlorine atom will take its place. UV light is required for the process. HCl also formed.

# REACTION FLOW CHARTS

The secret is that you don't have start at the beginning! Looking for clues and working backwards often works very well.



**Compound A** has a molecular formula  $\text{C}_3\text{H}_8\text{O}$ . It reacts readily with acidified potassium dichromate solution.

## Clues!

A had the molecular formula  $\text{C}_3\text{H}_8\text{O}$  – only 1 x O so an alcohol with a  $-\text{OH}$  group.

A reacts with acidified dichromate – more evidence for it being an alcohol.

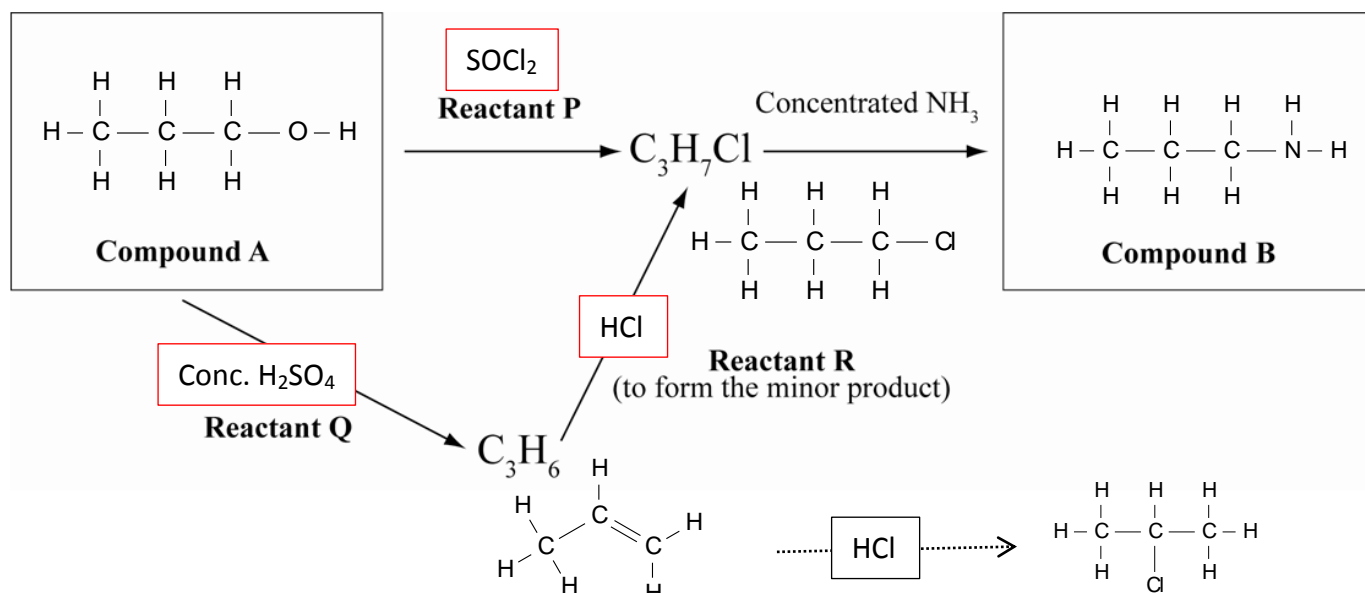
$\text{C}_3\text{H}_7\text{Cl}$  reacts with ammonia – B must be an amine (a substitution reaction) with  $-\text{Cl}$  off and  $-\text{NH}_2$  on.

$\text{C}_3\text{H}_6 \rightarrow \text{C}_3\text{H}_7\text{Cl}$ ; must be the addition of HCl; so  $\text{C}_3\text{H}_6$  had to be an alkene ( $\text{C}=\text{C}$ ), and an unsymmetrical one since there is a reference to the "minor" product.

A,  $\text{C}_3\text{H}_8\text{O} \rightarrow \text{C}_3\text{H}_6$  involves the loss of 2 x H and 1 x O, i.e.  $\text{H}_2\text{O}$ . This is an elimination or dehydration reaction, and needs conc.  $\text{H}_2\text{SO}_4$ .

A,  $\text{C}_3\text{H}_8\text{O} \rightarrow \text{C}_3\text{H}_7\text{Cl}$  needs substitution of  $-\text{OH}$  for  $-\text{Cl}$ , best done using  $\text{SOCl}_2$ .

Since the  $\text{C}_3\text{H}_7\text{Cl}$  is the minor product with the  $-\text{Cl}$  on the end, the  $-\text{OH}$  and  $-\text{NH}_2$  must also be there.



This would be the major product