

Surname	Centre Number	Candidate Number
First name(s)		2



GCE A LEVEL

A410U20-1



MONDAY, 20 JUNE 2022 – MORNING

CHEMISTRY – A level component 2

Organic Chemistry and Analysis

2 hours 30 minutes

ADDITIONAL MATERIALS

In addition to this examination paper, you will need a:

- calculator;
- **Data Booklet** supplied by WJEC.

INSTRUCTIONS TO CANDIDATES

Use black ink or black ball-point pen. Do not use gel pen or correction fluid.

You may use a pencil for graphs and diagrams only.

Write your name, centre number and candidate number in the spaces at the top of this page.

Section A Answer **all** questions.

Section B Answer **all** questions.

Write your answers in the spaces provided in this booklet. If you run out of space, use the additional page(s) at the back of the booklet, taking care to number the question(s) correctly.

Candidates are advised to allocate their time appropriately between **Section A (15 marks)** and **Section B (105 marks)**.

INFORMATION FOR CANDIDATES

The number of marks is given in brackets at the end of each question or part-question.

The maximum mark for this paper is 120.

Your answers must be relevant and must make full use of the information given to be awarded full marks for a question.

The assessment of the quality of extended response (QER) will take place in **Q.8(e)** and **Q.10(b)(ii)**.

Section A

Section B

For Examiner's use only		
Question	Maximum Mark	Mark Awarded
1. to 4.	15	
5.	16	
6.	16	
7.	15	
8.	18	
9.	20	
10.	20	
Total	120	

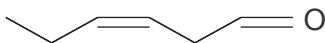
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SECTION AAnswer **all** questions.

1. The smell of freshly cut grass is partly due to (Z)-hex-3-enal.



- (a) Give the **displayed** formula of this compound. [1]

- (b) (Z)-hex-3-enal and cyclopentanal have the same molecular formula.

- (i) Calculate the percentage of oxygen by mass in these compounds. Give your answer to **three** significant figures. [2]

Percentage by mass = %

- (ii) State a chemical test that will give a positive result for (Z)-hex-3-enal but not for cyclopentanal. Give the result of the test. [1]

.....



2. (a) At a certain pH the amino acid norleucine, $\text{CH}_3(\text{CH}_2)_3\text{CH}(\text{NH}_2)\text{COOH}$, exists as its zwitterion form.

Draw the structure of this zwitterion.

[1]

- (b) A student was provided with a solution containing a mixture of the amino acids norleucine and isoleucine.

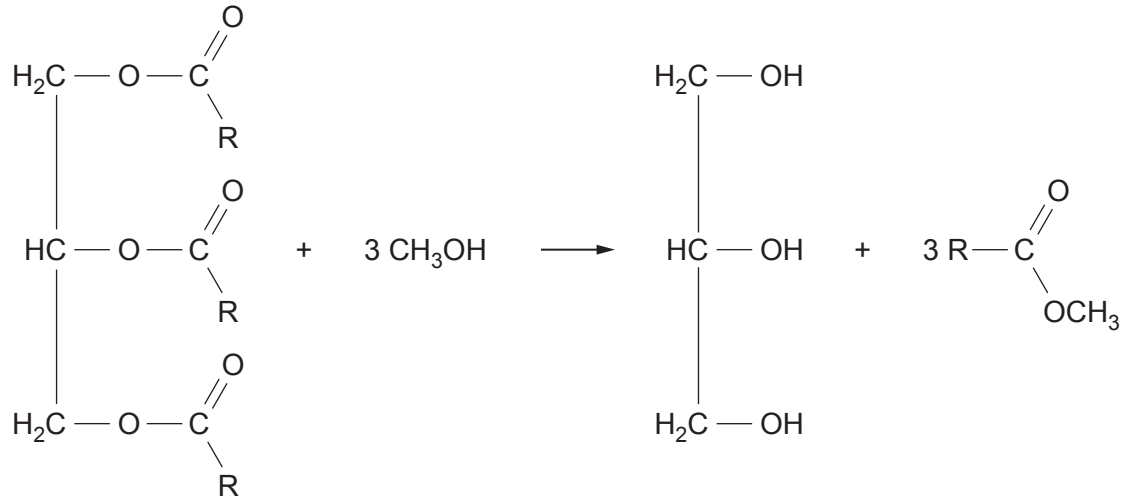
He obtained a thin layer chromatogram of the mixture but found that both norleucine and isoleucine had the same R_f value, using a particular solvent.

Suggest how he might obtain a thin layer chromatogram where these two amino acids have different R_f values.

[1]



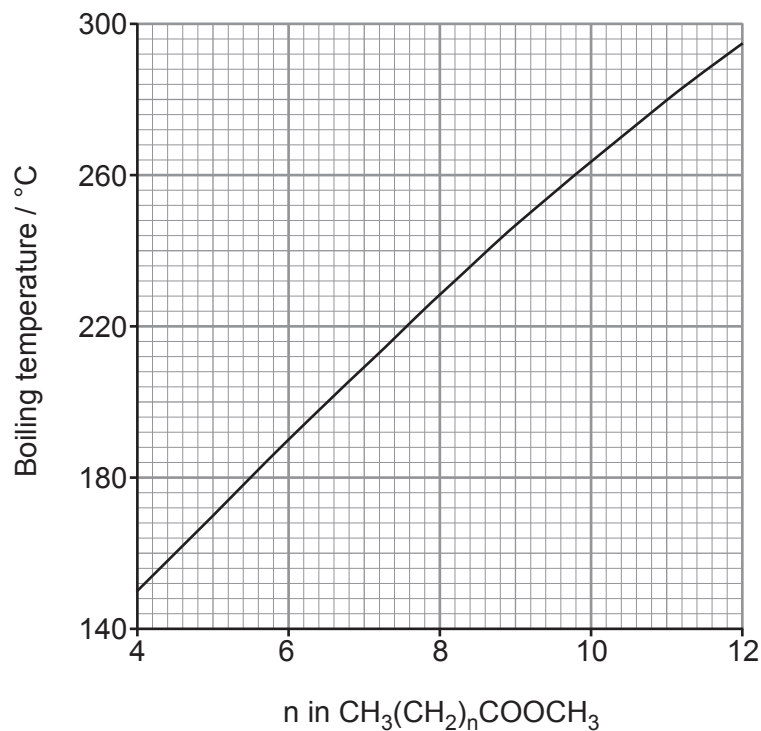
3. Biodiesel is an increasingly important fuel that is made from fats and oils. This fuel is often the methyl ester of a long chain fatty acid, produced by reacting the starting fat or oil with methanol.



- (a) The alkyl group in this ester is generally a linear chain of between 9 and 18 carbon atoms, depending on the source of the fat or oil.

The boiling temperature of these methyl esters increases as the length of the carbon chain increases.

Use the graph to deduce the boiling temperature of methyl decanoate, $C_{11}H_{22}O_2$. [1]



Boiling temperature °C

- (b) Comment on how the shape of the graph supports the statement that 'as the length of the alkyl chain increases, the relative effect of the polar group on the boiling temperature becomes less important'. [1]

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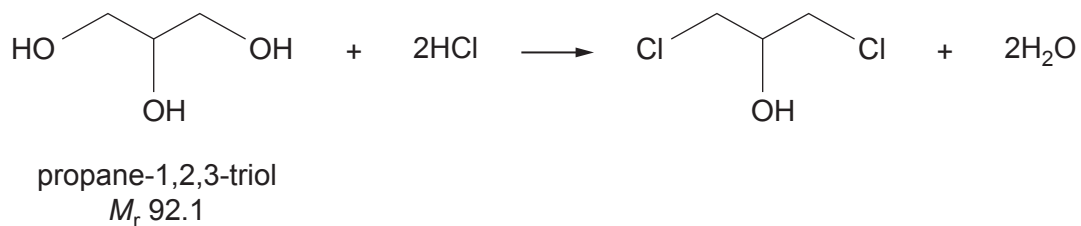


- (c) One problem with producing methyl esters for use as biofuels is the co-product, propane-1,2,3-triol, for which there is a limited market.

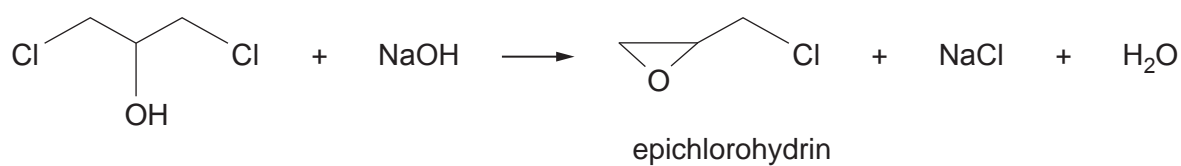
One use for this triol is in the manufacture of epichlorohydrin (C_3H_5ClO) [M_r 92.6].

This is made from propane-1,2,3-triol in a two-stage process.

Stage 1



Stage 2



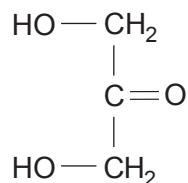
Calculate the atom economy of this two-stage reaction.

You should only consider propane-1,2,3-triol, hydrogen chloride and sodium hydroxide as reactants and epichlorohydrin as the required product in your calculation. [1]

Atom economy = %



- (d) Another product made from propane-1,2,3-triol is 1,3-dihydroxypropanone.



- (i) The traditional method of manufacture is via a biochemical route using propane-1,2,3-triol and appropriate bacteria.

A more modern method uses the aerial oxidation of this triol, in the presence of a catalyst.

Suggest **two** factors, apart from cost, that should be considered when choosing which of these two methods should be used. [2]

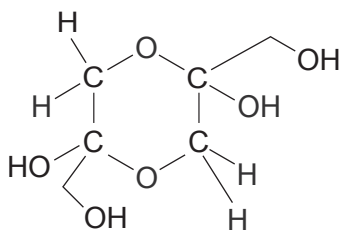
1.

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2.

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- (ii) 1,3-Dihydroxypropanone generally occurs as its dimer, from which the monomer is obtained by dissolving it in water.

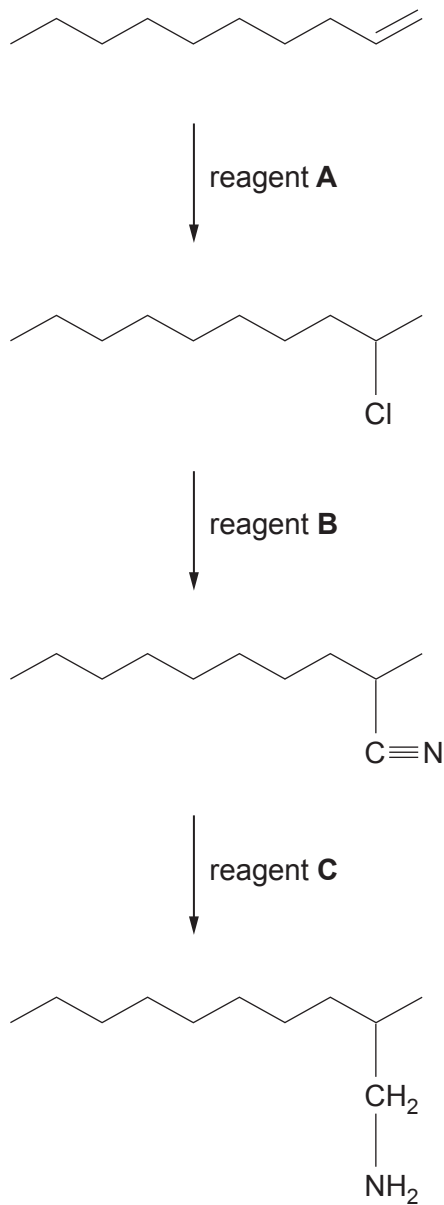


Identify a bond that is present in the monomer which is absent in the dimer, suggesting an infrared absorption value for this bond. [1]

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4. The sequence below shows a method for producing 2-methyldecylamine from dec-1-ene. State the reagent used in each of the three stages. [3]



Reagent A

Reagent B

Reagent C



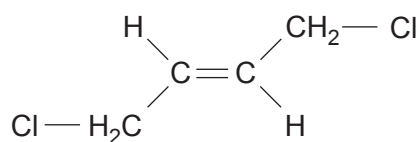
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SECTION BAnswer **all** questions.

5. (a) 1,4-Dichlorobut-2-ene is an important intermediate in the production of the synthetic rubber Neoprene. It exists as (*E*) and (*Z*) isomers.

*(E)*-1,4-dichlorobut-2-ene

- (i) Explain why 1,4-dichlorobut-2-ene exists as (*E*) and (*Z*) isomers. [2]

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- (ii) Describe the ^1H NMR spectrum of 1,4-dichlorobut-2-ene by completing the table. [2]

Group	Chemical shift δ /ppm	Splitting pattern
ClCH_2		
$\text{--- HC} = \text{C}$		



- (iii) The mass spectrum of 1,4-dichlorobut-2-ene shows a strong fragmentation signal at m/z 75.

The two common isotopes of chlorine are ^{35}Cl and ^{37}Cl in the ratio of 3:1.

Suggest a formula for this fragmentation signal at m/z 75. Show your working. [2]

- (iv) I. 1,4-Dichlorobut-2-ene is made from butadiene by chlorination. This then undergoes a rearrangement to give 3,4-dichlorobut-1-ene.

Give the **skeletal** formula of 3,4-dichlorobut-1-ene. [1]

- II. 3,4-Dichlorobut-1-ene is then further reacted to give 2-chlorobuta-1,3-diene, $\text{H}_2\text{C} = \text{C}(\text{Cl}) - \text{CH} = \text{CH}_2$.

Suggest a reagent that can be used for this stage. [1]

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- III. Radical polymerisation of 2-chlorobuta-1,3-diene (chloroprene) gives the medically important material poly(chloroprene) or Neoprene.

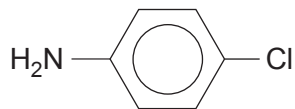
This polymer has important uses in gloves and facemasks that help to prevent transmission of the Covid-19 virus.

State what is meant by the term 'radical' and give the formula of a radical of your own choice. [2]

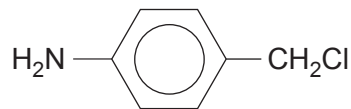
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- (b) (i) Explain why 4-chlorophenylamine does not react readily with aqueous sodium hydroxide but (4-chloromethyl)phenylamine produces (4-hydroxymethyl)phenylamine when treated with the same reagent.



4-chlorophenylamine



(4-chloromethyl)phenylamine

You should refer to both compounds in your answer.

[2]

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- (ii) 4-Chlorophenylamine reacts with nitric(III) acid (produced from sodium nitrate(III) and hydrochloric acid) to give a diazonium compound.

This can then react with phenol to give an azo dye.

- I. State the temperature necessary to produce a diazonium compound. [1]

..... °C

- II. Give the structure of the azo dye produced in this reaction. [1]

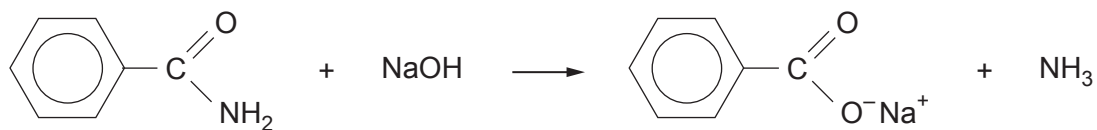
- III. Another azo dye, Solvent Yellow 7, has a maximum absorption in its UV-visible spectrum at a wavelength of 347 nm.

Calculate the frequency of this maximum absorption. [2]

Frequency = Hz



6. (a) (i) Benzamide reacts with aqueous sodium hydroxide to produce sodium benzoate and ammonia.



6.30 g of a damp solid sample of benzamide reacted with an excess of aqueous sodium hydroxide to give 0.0500 mol of ammonia.

Calculate the percentage purity of this sample of benzamide. [2]

Purity = %

- (ii) Benzamide starts to decompose at 100 °C into benzonitrile and water.

Suggest how this damp sample should be treated to give a dry sample. [1]

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- (b) (i) The addition of a solute to a solvent gives a solution that has a lower freezing temperature than the pure solvent. The freezing temperature obtained can be used to find the relative molecular mass of the solute.

In a modification to this method 0.698 g of a substituted amide, $\mathbf{R}-\text{CONH}(\text{C}_6\text{H}_5)$, was mixed with 5.00 g of camphor and the freezing temperature of the mixture found.

Pure camphor freezes at 179°C and the freezing temperature of the mixture was 145°C .

Use the formula below to work out the relative molecular mass (M_r) of the amide. [1]

$$\Delta T = \frac{1000 \times w \times k}{W \times M_r}$$

where ΔT is the lowering of the freezing temperature
 w is the mass of the substituted amide
 W is the mass of camphor
 k is 39.7

$M_r = \dots\dots\dots$

- (ii) Use the answer to part (i) to show that ' M_r ' for the \mathbf{R} group is 43. [1]

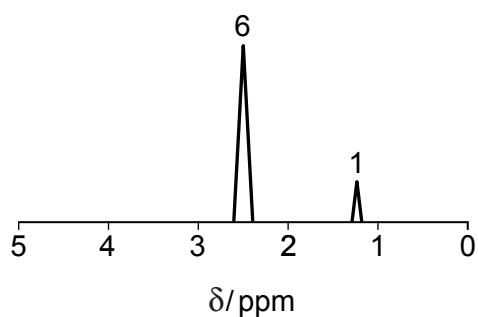
- (iii) \mathbf{R} represents the formula of a saturated hydrocarbon chain.

Deduce a molecular formula for the \mathbf{R} group. [1]

\mathbf{R} is



(iv) The diagram shows the low resolution ^1H NMR spectrum of the **R** group.



Use the spectrum to deduce the structure of the **R** group. Explain your answer. [2]

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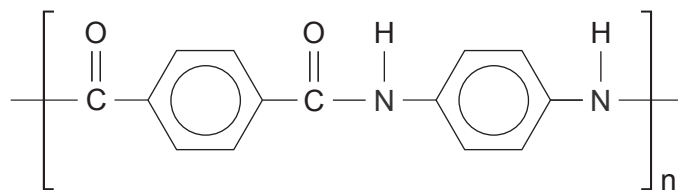
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- (c) The structure of the repeating unit of the condensation polymer Nomex[®] is shown below.



One way of producing this polymer is from benzene-1,3-dicarbonyl dichloride and benzene-1,3-diamine.

- (i) Give the **empirical** formula of benzene-1,3-diamine. [1]

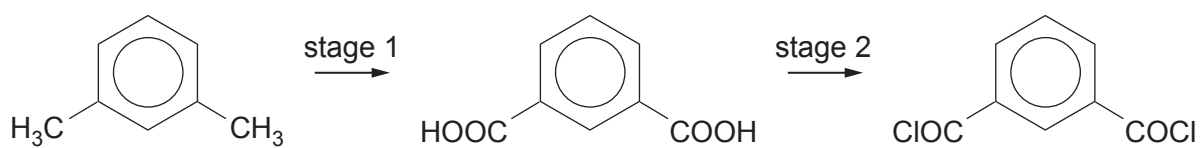
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- (ii) Nomex[®] is formed by condensation polymerisation.
State the meaning of 'condensation polymerisation'. [1]

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(iii) Benzene-1,3-dicarbonyl dichloride is produced from 1,3-dimethylbenzene.



I. State a reagent that can be used in the laboratory for stage 1. [1]

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II. State a reagent that can be used in the laboratory for stage 2. [1]

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(iv) Polyamides such as Nomex[®] are very slowly decomposed by heating with aqueous sodium hydroxide.

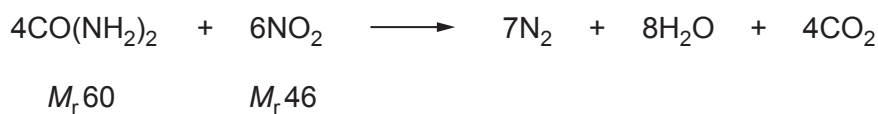
Suggest how the rate of this slow reaction can be increased. [1]

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- (d) Urea, $\text{CO}(\text{NH}_2)_2$, has an important use in reducing atmospheric pollution by reacting with nitrogen oxides in diesel exhaust fumes.



For this purpose, it is supplied as an aqueous solution containing 480g dm^{-3} of urea.

- (i) Calculate the mass of nitrogen(IV) oxide that can be removed from diesel exhaust fumes by 5 dm^3 of the urea solution. Give your answer in kg. [2]

Mass = kg

- (ii) This reaction removes toxic nitrogen(IV) oxide from the exhaust fumes.

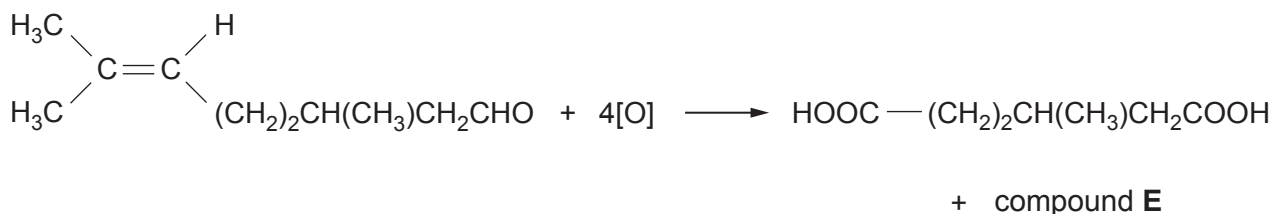
Use the equation to suggest **one** disadvantage of this reaction. Explain your answer. [1]

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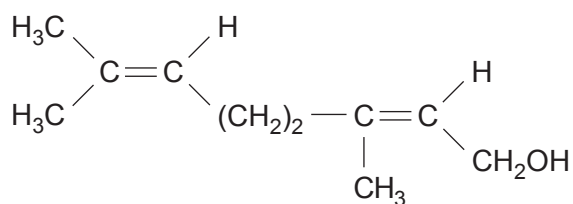


- (iii) Citronellal can be oxidised to 3-methylhexane-1,6-dioic acid and compound **E**.

Use the equation to find the molecular formula for compound **E** and then suggest its possible structure. [2]



- (iv) Citronella oil also contains geraniol.



The structure of geraniol shows that it contains two double bonds between carbon atoms.

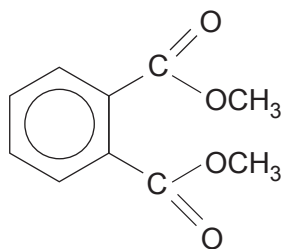
Calculate the volume of bromine that will need to be added to just react with 0.020 mol of geraniol.

The density of bromine is 3.2 g cm^{-3} . [2]

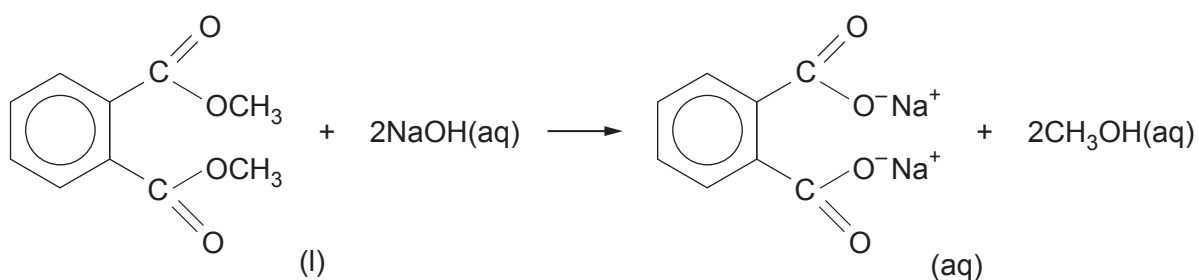
Volume = cm^3



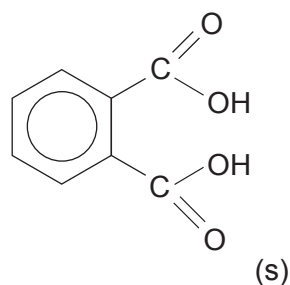
- (b) Dimethyl benzene-1,2-dicarboxylate (DMP) is a liquid ester that has also been used to repel insects.



DMP can be hydrolysed to benzene-1,2-dicarboxylic acid by refluxing it with aqueous sodium hydroxide, followed by acidification.



↓ acidification



- (i) Describe what is meant by the term 'refluxing'.

[1]

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- (ii) DMP and aqueous sodium hydroxide are immiscible.

Use the equation to deduce what will be observed when the refluxing stage has gone to completion. [1]

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- (iii) After refluxing, the mixture is acidified and the solid benzene-1,2-dicarboxylic acid is filtered off. It is then recrystallised from water.

State what should be done to the filtered solid before it is recrystallised. [1]

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- (iv) The solubility of benzene-1,2-dicarboxylic acid in water at two temperatures is shown in the table.

Temperature/°C	Solubility/g per 100g H ₂ O
14	0.7
100	18.0

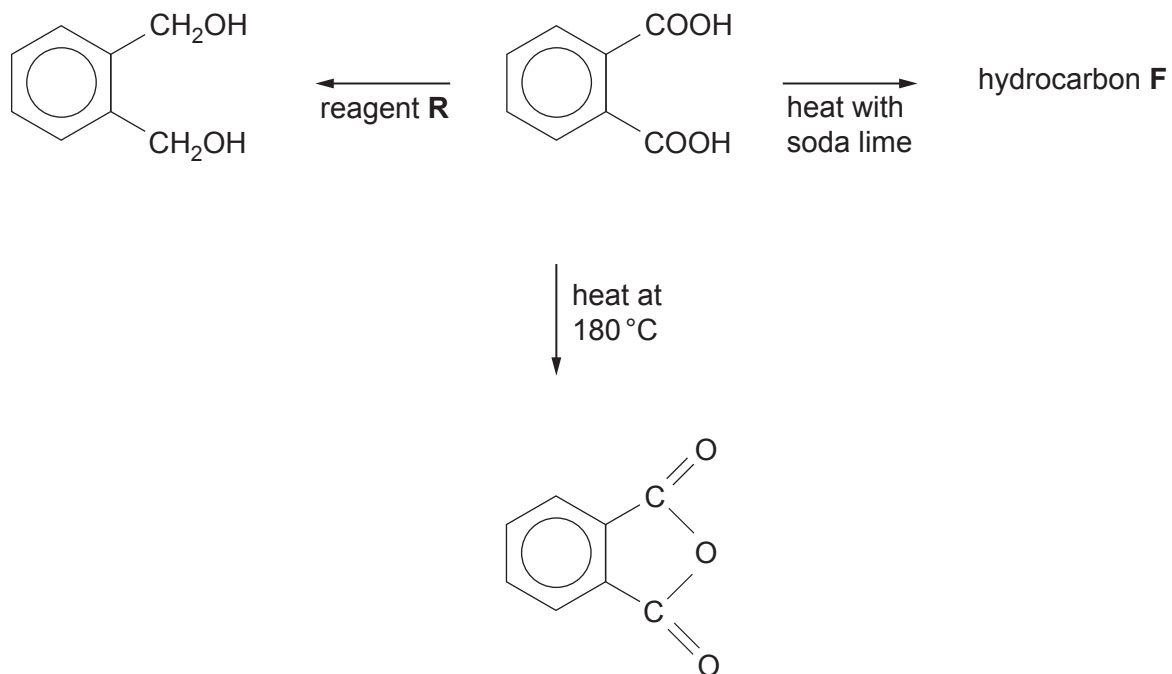
Calculate the mass of acid precipitated if a solution containing 8.0g of the acid in 50g of water is cooled from 100 °C to 14 °C. [1]

Mass = g



(c) Some of the reactions of benzene-1,2-dicarboxylic acid are shown below.

Use this flow chart to answer the questions that follow.



(i) Give the **formula** of reagent **R**. [1]

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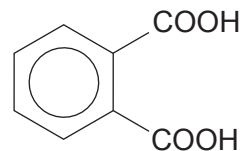
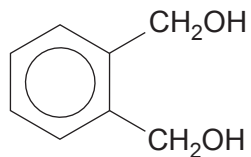
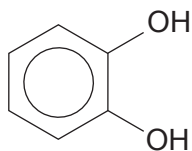
(ii) Give the structure of hydrocarbon **F**. [1]

(iii) State the type of reaction that occurs when benzene-1,2-dicarboxylic acid is heated at 180°C . [1]

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- (d) Describe a chemical test that will identify which of these three compounds, present in separate unlabelled aqueous solutions, is the most acidic. [2]



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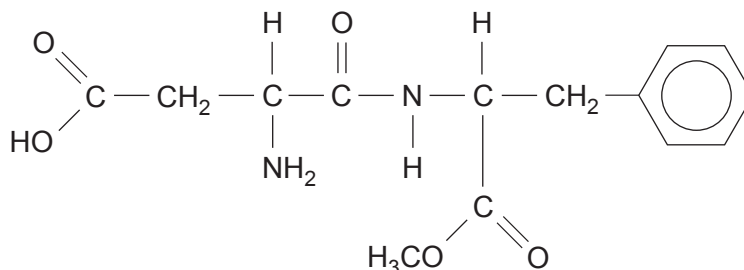
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8. (a) Aspartame is a dipeptide produced from aspartic acid and the methyl ester of phenylalanine.

The structure of α -aspartame is shown, where one COOH group of aspartic acid bonds to the NH_2 group of the methyl ester of phenylalanine.



Write the structure of β -aspartame, in which the other COOH group of aspartic acid bonds to the NH_2 group. [1]

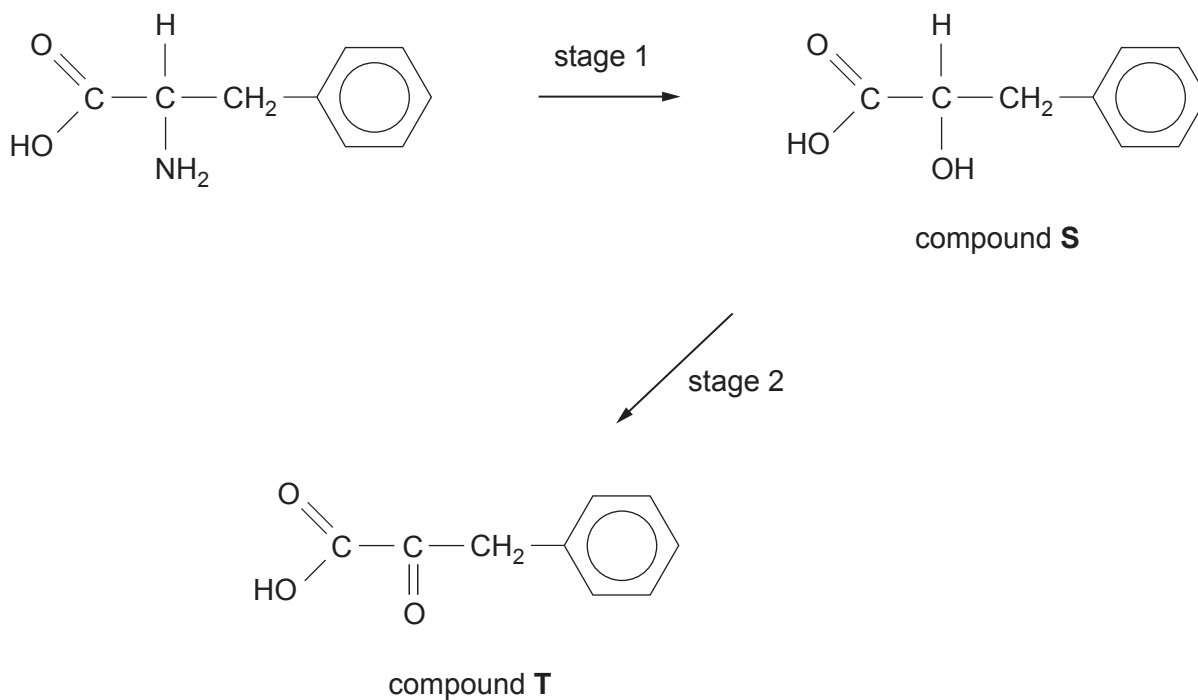
- (b) Both α - and β -aspartame (M_r 294) react with aqueous sodium hydroxide. In this reaction 3 mol of sodium hydroxide react with 1 mol of aspartame.

Calculate the volume of aqueous sodium hydroxide of concentration 4.00 mol dm^{-3} needed to just react with 73.5 g of aspartame. [2]

Volume = cm^3



(c) Phenylalanine can be converted to compound **T** in a two-stage reaction.



(i) Suggest a reagent(s) that can be used in stage 1 to produce compound **S**. [1]

.....

(ii) Explain why stage 2 can be described as an oxidation reaction. [1]

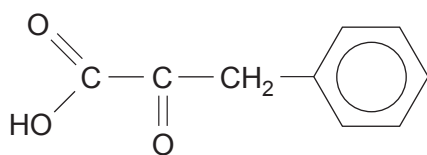
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(iii) Unusually, compound **T** reacts with Tollens' reagent to give a silver mirror.

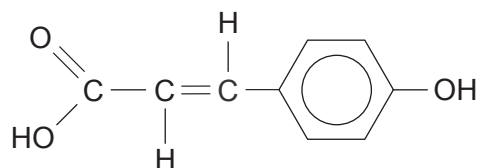
Give the structure of any other compound that will also produce a silver mirror with Tollens' reagent. [1]



(d) Compound **T** and compound **U** are isomers of formula $C_9H_8O_3$.



compound **T**



compound **U**

- (i) State the name of an element that will produce effervescence when added to a solution of either of these two compounds. [1]

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- (ii) State what is seen when aqueous iron(III) chloride is added to a solution of compound **U**. [1]

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(iii) Compound **U** reacts with aqueous bromine to give compound **V**.

Each molecule of compound **V** contains 9 carbon atoms and 3 oxygen atoms, as well as hydrogen and bromine.

Its mass spectrum shows a molecular ion at m/z 482.

Use this information to deduce a possible structure for compound **V**. Show your reasoning. [4]



- (e) Many characteristic reactions of benzene involve electrophilic substitution.

Discuss this statement, illustrating your answer by the bromination of benzene.

Your answer should include the mechanism for this reaction and any necessary conditions.

[6 QER]

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9. (a) The solubilities of some straight chain aliphatic carboxylic acids, $\text{R}-\text{COOH}$, in water at 20°C are shown in the table.

Number of carbon atoms in the R group	Solubility/g per 100 g H_2O
4	5.00
5	1.10
6	0.50
7	0.07
8	0.03

- (i) State the name of the carboxylic acid that has 7 carbon atoms in its **R** group. [1]
-
- (ii) Explain why the solubility decreases as the number of carbon atoms in the **R** group increases. Include a suitable diagram in your answer. [3]

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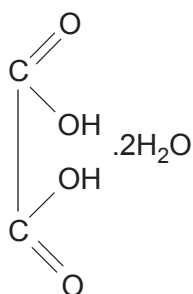


(b) Ethanedioic acid was first made in 1776 by oxidising sucrose with concentrated nitric acid (represented as [O] in the equation below).

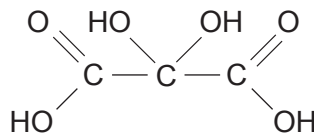
(i) Complete the equation for this reaction. [1]



(ii) Ethanedioic acid is obtained from the aqueous mixture as the dihydrate. A small quantity of hydrated mesoxalic acid is also formed during this reaction.



ethanedioic acid
dihydrate



hydrated
mesoxalic acid

I. If the ethanedioic acid dihydrate is contaminated with a small quantity of hydrated mesoxalic acid, describe how the melting temperature of the dihydrate will be affected. [1]

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II. Explain how the ^{13}C NMR spectra of these hydrated acids would differ.

The position of the signals is **not** required in your answer. [2]

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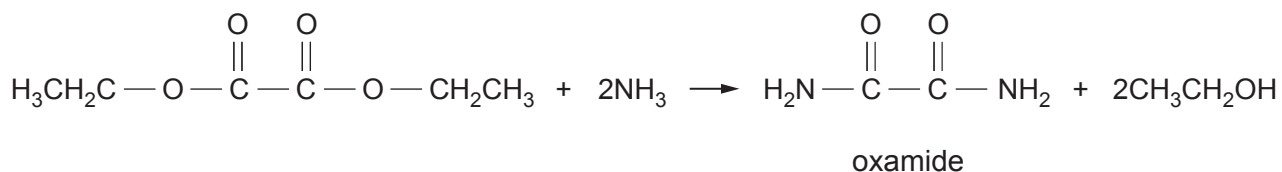
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(c) A student made oxamide by reacting diethyl ethanedioate with aqueous ammonia.



She followed a book method, starting with 13.5 cm³ of diethyl ethanedioate.

(i) Calculate the number of moles of diethyl ethanedioate used.

Its density is 1.08 g cm⁻³.

[2]

Number of moles = mol

(ii) The method stated that she should obtain a yield of 57%.

Calculate the mass that she should obtain.

[1]

Mass = g

(iii) On weighing the product, the student found that she had obtained a yield of 110%.

Apart from incorrect weighing of the product, suggest **two** reasons why this may have happened.

[2]

1.

2.



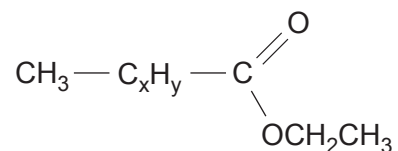
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- (d) There is considerable interest in developing drugs that are used to treat high fat levels in the blood.

One compound being studied is E-EPA, which is the ethyl ester of a linear polyunsaturated carboxylic acid, where C_xH_y represents the number of carbon and hydrogen atoms in a long hydrocarbon chain.



- (i) 0.0600 mol of E-EPA has a mass of 19.8 g.

Calculate the relative molecular mass (M_r) of E-EPA.

[1]

$M_r = \dots\dots\dots$

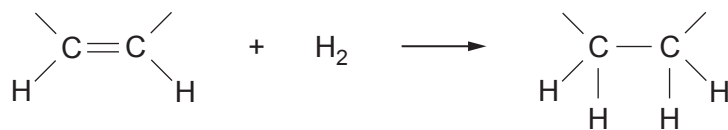
- (ii) Use the formula of E-EPA and your answer to part (i) to calculate the 'relative molecular mass' of the C_xH_y part of the molecule.

[2]

$M_r = \dots\dots\dots$



- (iii) 0.0600 mol of E-EPA reacts with 7.35 dm³ of hydrogen gas, measured at 298 K and 1 atm, to convert the compound to a saturated ester.



Use this information to find the number of — CH = CH — groups present in each molecule of E-EPA. [2]

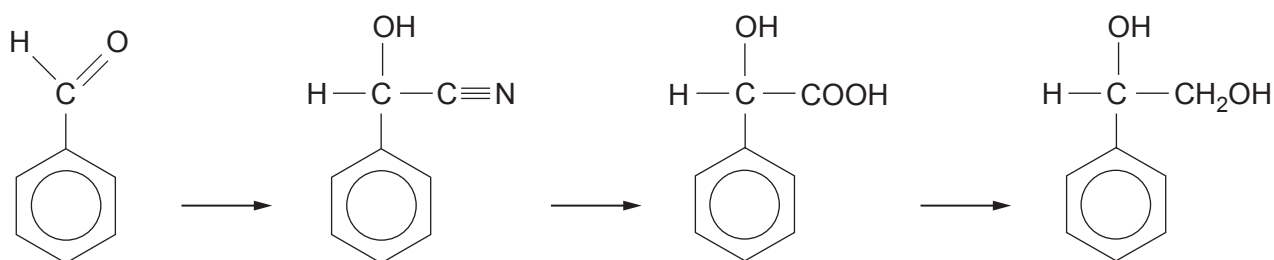
Number of — CH = CH — groups =

- (iv) Use your answers to parts (ii) and (iii) to calculate the number of CH₂ groups present in each molecule of E-EPA. [2]

Number of CH₂ groups =



10. (a) The scheme below shows a sequence of reactions starting from benzaldehyde.



(i) One way of increasing the length of a carbon chain is by reacting an aldehyde with hydrogen cyanide.

I. Show the mechanism for the reaction of benzaldehyde with hydrogen cyanide to produce 2-hydroxy-2-phenylethanenitrile.

Include appropriate charges and curly arrows.

[3]

II. State the name of the reaction mechanism that you have described in part I. above.

[1]

.....



- (ii) State a reagent that is used in aqueous solution to convert 2-hydroxy-2-phenylethanenitrile to the corresponding carboxylic acid, 2-hydroxy-2-phenylethanoic acid. [1]

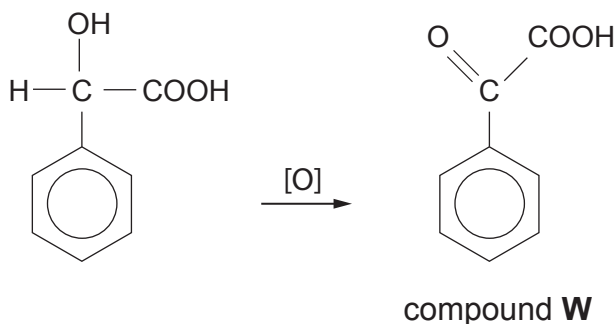
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- (iii) Suggest why a solution of 2-hydroxy-2-phenylethanoic acid, produced in part (ii), does not show any optical activity when placed in a polarimeter. [1]

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- (iv) 2-Hydroxy-2-phenylethanoic acid can be oxidised to the corresponding keto-acid, compound **W**.



Suggest why compound **W** does not show any optical activity when a solution is placed in a polarimeter. [1]

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- (v) I. Explain why 2-hydroxy-2-phenylethanoic acid will react with methanol, in the presence of a catalyst.

Give the formula of the organic compound formed. [2]

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.....

- II. Explain why 2-hydroxy-2-phenylethanoic acid can also react with ethanoic acid, in the presence of a catalyst. [1]

.....
.....



(b) Pentan-2-one can be made by the oxidation of pentan-2-ol using acidified potassium dichromate.

(i) State the colour change that is seen in the reaction flask as acidified dichromate is added to pentan-2-ol. [1]

.....

(ii) After the reaction the mixture is distilled and the fraction boiling between 100 and 120 °C is collected. This distillate is largely pentan-2-one. It also contains a little unreacted pentan-2-ol and water.

Describe how you would obtain a dry sample of pentan-2-one from this distillate.

It is not necessary to redistil your dry sample of pentan-2-one.

You are given the following information to help you in your answer.

- You should use a separating funnel
- Pentan-2-ol is more soluble than pentan-2-one in water
- Pentan-2-one is very soluble in ethoxyethane
- Ethoxyethane boils at 35 °C and is very flammable
- The density of ethoxyethane is 0.71 g cm⁻³
- Solid anhydrous magnesium sulfate is a suitable drying agent for pentan-2-one

[6 QER]

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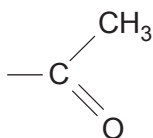


- (iii) Describe how infrared spectroscopy could confirm that there is no longer pentan-2-ol present. [1]

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.....

- (iv) Give the reagent(s) and an observation to show that pentan-2-one contains the following group. [2]



.....

.....

END OF PAPER

20





MONDAY, 20 JUNE 2022 – MORNING

CHEMISTRY – A level component 2
Data Booklet

Avogadro constant	$N_A = 6.02 \times 10^{23} \text{ mol}^{-1}$
molar gas constant	$R = 8.31 \text{ J mol}^{-1} \text{ K}^{-1}$
molar gas volume at 273 K and 1 atm	$V_m = 22.4 \text{ dm}^3 \text{ mol}^{-1}$
molar gas volume at 298 K and 1 atm	$V_m = 24.5 \text{ dm}^3 \text{ mol}^{-1}$
Planck constant	$h = 6.63 \times 10^{-34} \text{ Js}$
speed of light	$c = 3.00 \times 10^8 \text{ ms}^{-1}$
density of water	$d = 1.00 \text{ g cm}^{-3}$
specific heat capacity of water	$c = 4.18 \text{ J g}^{-1} \text{ K}^{-1}$
ionic product of water at 298 K	$K_w = 1.00 \times 10^{-14} \text{ mol}^2 \text{ dm}^{-6}$
fundamental electronic charge	$e = 1.60 \times 10^{-19} \text{ C}$

temperature (K) = temperature (°C) + 273

$$1 \text{ dm}^3 = 1000 \text{ cm}^3$$

$$1 \text{ m}^3 = 1000 \text{ dm}^3$$

$$1 \text{ tonne} = 1000 \text{ kg}$$

$$1 \text{ atm} = 1.01 \times 10^5 \text{ Pa}$$

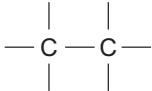
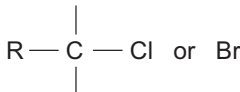
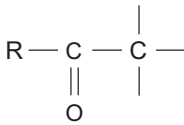
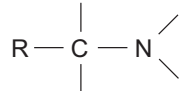
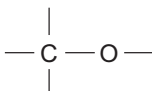
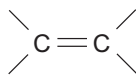
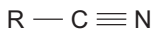
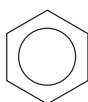
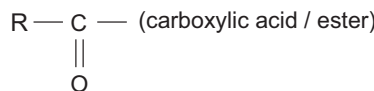
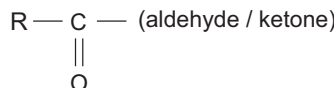
Multiple	Prefix	Symbol
10^{-9}	nano	n
10^{-6}	micro	μ
10^{-3}	milli	m

Multiple	Prefix	Symbol
10^3	kilo	k
10^6	mega	M
10^9	giga	G

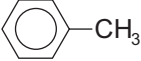
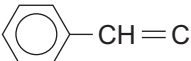
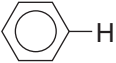
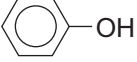
Infrared absorption values

Bond	Wavenumber / cm^{-1}
C—Br	500 to 600
C—Cl	650 to 800
C—O	1000 to 1300
C=C	1620 to 1670
C=O	1650 to 1750
C≡N	2100 to 2250
C—H	2800 to 3100
O—H (carboxylic acid)	2500 to 3200 (very broad)
O—H (alcohol / phenol)	3200 to 3550 (broad)
N—H	3300 to 3500

¹³C NMR chemical shifts relative to TMS = 0

Type of carbon	Chemical shift, δ (ppm)
	5 to 40
	10 to 70
	20 to 50
	25 to 60
	50 to 90
	90 to 150
	110 to 125
	110 to 160
	160 to 185
	190 to 220

¹H NMR chemical shifts relative to TMS = 0

Type of proton	Chemical shift, δ (ppm)
$-\text{CH}_3$	0.1 to 2.0
$\text{R}-\text{CH}_3$	0.9
$\text{R}-\text{CH}_2-\text{R}$	1.3
$\text{CH}_3-\text{C}\equiv\text{N}$	2.0
$\text{CH}_3-\text{C}(=\text{O})$	2.0 to 2.5
$-\text{CH}_2-\text{C}(=\text{O})$	2.0 to 3.0
	2.2 to 2.3
$\text{HC}-\text{Cl}$ or $\text{HC}-\text{Br}$	3.1 to 4.3
$\text{HC}-\text{O}$	3.3 to 4.3
$\text{R}-\text{OH}$	4.5 *
$-\text{C}=\text{CH}$	4.5 to 6.3
$-\text{C}=\text{CH}-\text{CO}$	5.8 to 6.5
	6.5 to 7.5
	6.5 to 8.0
	7.0 *
$\text{R}-\text{C}(=\text{O})\text{H}$	9.8 *
$\text{R}-\text{C}(=\text{O})\text{OH}$	11.0 *

*variable figure dependent on concentration and solvent

THE PERIODIC TABLE

Group

1 2 3 4 5 6 7 0

Period

1	1.01 H Hydrogen 1											4.00 He Helium 2						
2	6.94 Li Lithium 3	9.01 Be Beryllium 4											19.0 F Fluorine 9	20.2 Ne Neon 10				
3	23.0 Na Sodium 11	24.3 Mg Magnesium 12											35.5 Cl Chlorine 17	40.0 Ar Argon 18				
4	39.1 K Potassium 19	40.1 Ca Calcium 20	45.0 Sc Scandium 21	47.9 Ti Titanium 22	50.9 V Vanadium 23	52.0 Cr Chromium 24	54.9 Mn Manganese 25	55.8 Fe Iron 26	58.9 Co Cobalt 27	58.7 Ni Nickel 28	63.5 Cu Copper 29	65.4 Zn Zinc 30	69.7 Ga Gallium 31	72.6 Ge Germanium 32	74.9 As Arsenic 33	79.0 Se Selenium 34	79.9 Br Bromine 35	83.8 Kr Krypton 36
5	85.5 Rb Rubidium 37	87.6 Sr Strontium 38	88.9 Y Yttrium 39	91.2 Zr Zirconium 40	92.9 Nb Niobium 41	95.9 Mo Molybdenum 42	98.9 Tc Technetium 43	101 Ru Ruthenium 44	103 Rh Rhodium 45	106 Pd Palladium 46	108 Ag Silver 47	112 Cd Cadmium 48	115 In Indium 49	119 Sn Tin 50	122 Sb Antimony 51	128 Te Tellurium 52	127 I Iodine 53	131 Xe Xenon 54
6	133 Cs Caesium 55	137 Ba Barium 56	139 La Lanthanum 57	179 Hf Hafnium 72	181 Ta Tantalum 73	184 W Tungsten 74	186 Re Rhenium 75	190 Os Osmium 76	192 Ir Iridium 77	195 Pt Platinum 78	197 Au Gold 79	201 Hg Mercury 80	204 Tl Thallium 81	207 Pb Lead 82	209 Bi Bismuth 83	(210) Po Polonium 84	(210) At Astatine 85	(222) Rn Radon 86
7	(223) Fr Francium 87	(226) Ra Radium 88	(227) Ac Actinium 89															

Key

Ar	Symbol
Name	atomic number
Z	relative atomic mass

f block

140 Ce Cerium 58	141 Pr Praseodymium 59	144 Nd Neodymium 60	(147) Pm Promethium 61	150 Sm Samarium 62	(153) Eu Europium 63	157 Gd Gadolinium 64	159 Tb Terbium 65	163 Dy Dysprosium 66	165 Ho Holmium 67	167 Er Erbium 68	169 Tm Thulium 69	173 Yb Ytterbium 70	175 Lu Lutetium 71
232 Th Thorium 90	(231) Pa Protactinium 91	238 U Uranium 92	(237) Np Neptunium 93	(242) Pu Plutonium 94	(243) Am Americium 95	(247) Cm Curium 96	(245) Bk Berkelium 97	(251) Cf Californium 98	(254) Es Einsteinium 99	(253) Fm Fermium 100	(256) Md Mendelevium 101	(254) No Nobelium 102	(257) Lr Lawrencium 103

► Lanthanoid elements

►► Actinoid elements