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ADVANCED SUBSIDIARY (AS)
General Certificate of Education
2023

Centre Number

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Candidate Number

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Chemistry

Assessment Unit AS 3

assessing

Module 3: Practical Examination

Practical Booklet B (Theory)



[SCH32]

SCH32

THURSDAY 1 JUNE, MORNING

TIME

1 hour 15 minutes.

INSTRUCTIONS TO CANDIDATES

Write your Centre Number and Candidate Number in the spaces provided at the top of this page.

You must answer the questions in the spaces provided.

Do not write outside the boxed area on each page or on blank pages.

Complete in black ink only. **Do not write with a gel pen or a pencil.**

Answer **all four** questions.

INFORMATION FOR CANDIDATES

The total mark for this paper is 55.

Figures in brackets printed down the right-hand side of pages indicate the marks awarded to each question or part question.

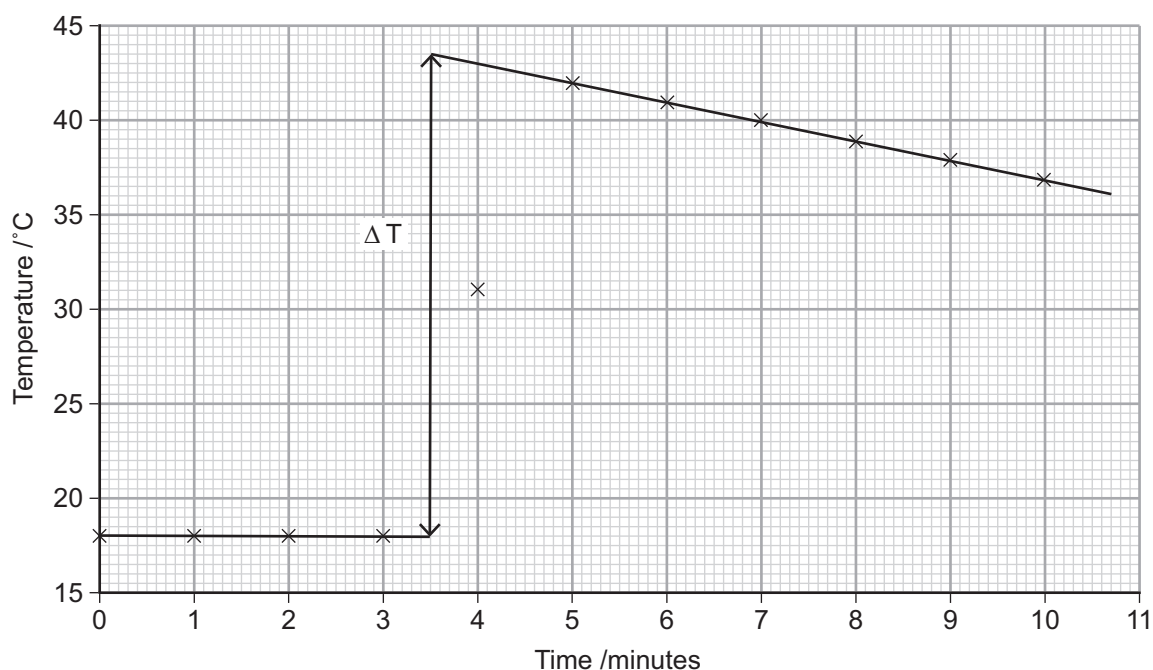
A Periodic Table of Elements (including some data) is provided.



1 The enthalpy change for the reaction between zinc and copper(II) sulfate solution was determined using the method below.

1. Weigh 2.50 g of zinc powder.
2. Measure 25 cm³ of 0.50 mol dm⁻³ copper(II) sulfate solution using a measuring cylinder and transfer the solution into a polystyrene cup.
3. Stir the solution continuously with a thermometer and record the temperature of the solution at 1 minute intervals up to 3 minutes.
4. At exactly 3.5 minutes add the zinc powder to the copper(II) sulfate solution.
5. Continue to stir the solution and record the temperature each minute from 4 to 10 minutes.

The results obtained are shown on the graph below. Two straight lines of best fit have been drawn on the graph, one from 0.0 to 3.5 minutes and the other from 5.0 to 10.0 minutes extrapolating back to 3.5 minutes.



(a) Use the lines of best fit to determine the temperature change, ΔT , at 3.5 minutes.

Answer _____ [2]

(b) (i) Use the answer from (a) to calculate the heat produced in the reaction in kJ. Give your answer to 2 decimal places.

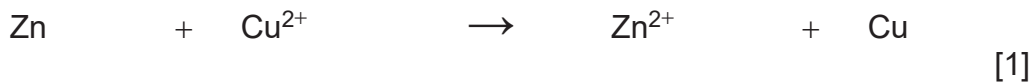
Answer _____ kJ [2]

(ii) State two assumptions you made when carrying out the calculation in (b)(i).

[2]

(c) The 2.50 g of zinc powder is an excess.

(i) The ionic equation for the reaction is given below. Add state symbols to this ionic equation.



(ii) Calculate the number of moles of copper(II) sulfate in 25 cm³ of the solution.

Answer _____ [1]

[Turn over



(iii) Calculate the minimum mass of zinc powder required to react with the copper(II) sulfate solution.

Answer _____ [1]

(iv) Use your answers to (b)(i) and (c)(ii) to calculate the enthalpy change for the reaction in kJ mol^{-1} . Give your answer to 2 significant figures.

Answer _____ kJ mol^{-1} [3]

(v) Describe how the presence of sulfate ions in the solution could be confirmed.

[2]

(d) State the flame test colour for copper(II) ions.

_____ [1]





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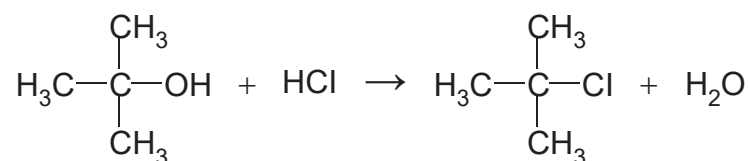
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[Turn over



16SCH3205

- 2 The halogenoalkane 2-chloro-2-methylpropane may be prepared by the reaction between 2-methylpropan-2-ol and concentrated hydrochloric acid.



The steps of a method are:

1. Measure approximately 6.5 cm^3 of 2-methylpropan-2-ol using a measuring cylinder. Record the mass of the measuring cylinder and the alcohol.
2. Transfer the alcohol into a separating funnel and record the mass of the measuring cylinder after transfer.

Mass of measuring cylinder and alcohol = 157.55 g

Mass of measuring cylinder after transfer = 152.45 g

3. Measure approximately 20 cm^3 (an excess) of concentrated hydrochloric acid using a measuring cylinder and gradually add the acid to the 2-methylpropan-2-ol in the separating funnel.
4. Stopper the funnel and shake the mixture periodically over the next 15 minutes.
5. Allow the mixture to stand until two layers appear, remove the stopper and separate the layers. Discard the aqueous layer.
6. Slowly add 10 cm^3 of sodium hydrogencarbonate solution to the crude 2-chloro-2-methylpropane in the separating funnel. Stopper the funnel and shake the contents gently, inverting the funnel and opening the tap periodically.
7. Repeat the washing with sodium hydrogencarbonate solution until no more gas is given off.
8. Allow the layers to separate, run off and discard the aqueous layer and then run the crude 2-chloro-2-methylpropane into a clean conical flask.
9. To the liquid in the flask, add small amounts of anhydrous sodium sulfate and swirl after each addition. Remove the solid sodium sulfate.
10. Transfer the 2-chloro-2-methylpropane into a pear-shaped flask. Distil the 2-chloro-2-methylpropane (boiling point = 51°C) and record the mass of product collected.



(a) (i) Calculate the mass of 2-methylpropan-2-ol used in the preparation.

Answer _____ g [1]

(ii) Suggest why a measuring cylinder is used to measure the volume of concentrated hydrochloric acid rather than a more accurate piece of apparatus.

_____ [1]

(iii) Suggest why the concentrated hydrochloric acid is added **gradually** to the 2-methylpropan-2-ol in Step 3.

_____ [1]

(iv) Explain, without reference to density values, how the aqueous layer could be identified from the organic layer in the separating funnel in Step 5.

_____ [2]

(v) Explain, with an equation, the purpose of adding sodium hydrogencarbonate solution to the crude 2-chloro-2-methylpropane in Step 6.

_____ [2]

[Turn over



(vi) Suggest the purpose of adding anhydrous sodium sulfate in Step 9 and explain how you would know when enough has been added to the crude 2-chloro-2-methylpropane.

_____ [2]

(vii) Describe how the solid is removed in Step 9.

_____ [1]

(viii) What should be added to the contents of the pear-shaped flask in Step 10 to promote smooth boiling?

_____ [1]

(ix) The pear-shaped flask is heated in a water bath in Step 10. Suggest why the pear-shaped flask is not heated directly.

_____ [1]

(x) Suggest the effect that any impurities present in the organic product will have on its boiling point.

_____ [1]

(b) Assuming a 40% yield, use your answer to (a)(i) to calculate the mass of 2-chloro-2-methylpropane obtained.

Answer _____ g [3]



(c) 2-methylpropan-2-ol reacts slowly with sodium.

(i) Write an equation for the reaction between 2-methylpropan-2-ol and sodium.

_____ [1]

(ii) Describe a test for the gas produced in the reaction in (c)(i).

_____ [1]



- 3 The trend in oxidising ability of the halogens can be determined by reacting aqueous solutions of halogens with aqueous solutions of potassium halide salts.

(a) Complete the table.

Aqueous solution	Colour of aqueous solution
Chlorine	
Bromine	
Iodine	
Potassium halide	

[2]

- (b) A 1 cm³ portion of each aqueous halogen solution is added separately to 1 cm³ of potassium chloride solution in a test tube and any observations noted. The procedure is repeated using potassium bromide solution and also using potassium iodide solution.

(i) Complete the following table, using a tick (✓) to indicate that a reaction occurs and a cross (X) to indicate that no reaction occurs.

	Potassium chloride	Potassium bromide	Potassium iodide
Chlorine			✓
Bromine			
Iodine			

[1]

(ii) Write an equation for the reaction of chlorine with potassium iodide.

[1]



(iii) Some hexane was added to the test tube after the reaction of aqueous chlorine with potassium iodide solution was complete. The test tube was stoppered and shaken for one minute. The contents were allowed to settle. What would be observed?

[3]

(c) Chlorine reacts with water and with sodium hydroxide solution.

(i) Write an equation for the reaction of chlorine with water.

[1]

(ii) State the conditions required for the reaction between chlorine and sodium hydroxide solution which yield products containing chlorine in the same oxidation states as those in (c)(i).

[1]

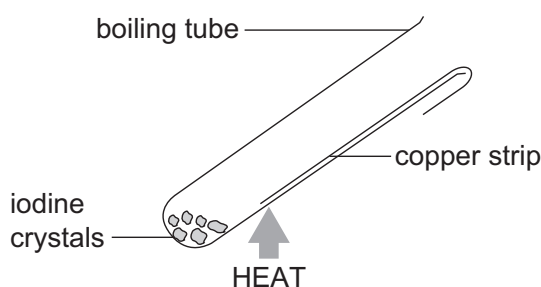
(iii) Suggest why ozone is often preferred to chlorine in water treatment.

[1]

[Turn over



- 4 The empirical formula of copper iodide may be determined using the apparatus below.



A small quantity of iodine crystals is added to the boiling tube. A clean copper strip is placed into the boiling tube and bent at one end so that it fits over the mouth of the boiling tube.

The part of the copper strip nearest the iodine crystals is heated gently in a fume cupboard until no more purple vapour is observed. Once the boiling tube is cool, the copper strip is carefully removed and reweighed. The yellow coating of copper iodide is scraped from the surface of the copper strip and the copper strip reweighed. The following results are obtained.

	Mass /g
Initial mass of copper strip	2.94
Mass of copper strip and copper iodide	3.28
Final mass of copper strip	2.77

- (a) Explain why the iodine crystals are not heated directly.

_____ [1]

- (b) Suggest why the procedure is carried out in a fume cupboard.

_____ [1]



(c) Calculate the mass of iodine that reacted.

Answer _____ g [1]

(d) Calculate the mass of copper that reacted.

Answer _____ g [1]

(e) Calculate the empirical formula of the copper iodide formed.

Answer _____ [3]

[Turn over

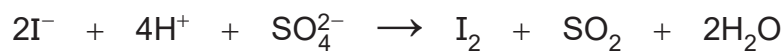
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(f) Some of the copper iodide formed was removed and placed in a test tube. A few drops of concentrated sulfuric acid were added.

(i) The ionic equation for one reaction which occurs when concentrated sulfuric acid is added to the sample of copper iodide is:



Explain, using oxidation numbers, why this is a redox reaction.

[3]

(ii) Describe how you would test for the presence of iodine.

[2]

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Question Number	Marks
1	
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Total Marks	

Examiner Number

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General Information

1 tonne = 10^6 g

1 metre = 10^9 nm

One mole of any gas at 293 K and a pressure of 1 atmosphere (10^5 Pa) occupies a volume of 24 dm³

Avogadro Constant = 6.02×10^{23} mol⁻¹

Planck Constant = 6.63×10^{-34} Js

Specific Heat Capacity of water = $4.2 \text{ J g}^{-1} \text{ K}^{-1}$

Speed of Light = $3 \times 10^8 \text{ ms}^{-1}$



Characteristic absorptions in IR spectroscopy

Wavenumber/cm ⁻¹	Bond	Compound
550–850	C–X (X = Cl, Br, I)	Haloalkanes
750–1100	C–C	Alkanes, alkyl groups
1000–1300	C–O	Alcohols, esters, carboxylic acids
1450–1650	C=C	Arenes
1600–1700	C=C	Alkenes
1650–1800	C=O	Carboxylic acids, esters, aldehydes, ketones, amides, acyl chlorides
2200–2300	C≡N	Nitriles
2500–3200	O–H	Carboxylic acids
2750–2850	C–H	Aldehydes
2850–3000	C–H	Alkanes, alkyl groups, alkenes, arenes
3200–3600	O–H	Alcohols
3300–3500	N–H	Amines, amides

Proton Chemical Shifts in Nuclear Magnetic Resonance Spectroscopy (relative to TMS)

Chemical Shift	Structure	
0.5–2.0	–CH	Saturated alkanes
0.5–5.5	–OH	Alcohols
1.0–3.0	–NH	Amines
2.0–3.0	–CO–CH	Ketones
	–N–CH	Amines
	C ₆ H ₅ –CH	Arene (aliphatic on ring)
2.0–4.0	X–CH	X = Cl or Br (3.0–4.0) X = I (2.0–3.0)
4.5–6.0	–C=CH	Alkenes
5.5–8.5	RCONH	Amides
6.0–8.0	–C ₆ H ₅	Arenes (on ring)
9.0–10.0	–CHO	Aldehydes
10.0–12.0	–COOH	Carboxylic acids

These chemical shifts are concentration and temperature dependent and may be outside the ranges indicated above.

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Data Leaflet

Including the Periodic Table of the Elements

For the use of candidates taking
Advanced Subsidiary and
Advanced Level Examinations

Copies must be free from notes or additions of any kind. No other type of data booklet or information sheet is authorised for use in the examinations

gce a/as examinations

chemistry

THE PERIODIC TABLE OF ELEMENTS

Group

	I	II											III	IV	V	VI	VII	0
	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18
1 H Hydrogen 1													4 He Helium 2					
7 Li Lithium 3	9 Be Beryllium 4											11 B Boron 5	12 C Carbon 6	14 N Nitrogen 7	16 O Oxygen 8	19 F Fluorine 9	20 Ne Neon 10	
23 Na Sodium 11	24 Mg Magnesium 12											27 Al Aluminium 13	28 Si Silicon 14	31 P Phosphorus 15	32 S Sulfur 16	35.5 Cl Chlorine 17	40 Ar Argon 18	
39 K Potassium 19	40 Ca Calcium 20	45 Sc Scandium 21	48 Ti Titanium 22	51 V Vanadium 23	52 Cr Chromium 24	55 Mn Manganese 25	56 Fe Iron 26	59 Co Cobalt 27	59 Ni Nickel 28	64 Cu Copper 29	65 Zn Zinc 30	70 Ga Gallium 31	73 Ge Germanium 32	75 As Arsenic 33	79 Se Selenium 34	80 Br Bromine 35	84 Kr Krypton 36	
85 Rb Rubidium 37	88 Sr Strontium 38	89 Y Yttrium 39	91 Zr Zirconium 40	93 Nb Niobium 41	96 Mo Molybdenum 42	98 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	
133 Cs Caesium 55	137 Ba Barium 56	139 La [*] Lanthanum 57	178 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	
223 Fr Francium 87	226 Ra Radium 88	227 Ac [†] Actinium 89	261 Rf Rutherfordium 104	262 Db Dubnium 105	266 Sg Seaborgium 106	264 Bh Bohrium 107	277 Hs Hassium 108	268 Mt Meitnerium 109	271 Ds Darmstadtium 110	272 Rg Roentgenium 111	285 Cn Copernicium 112							

* 58 – 71 Lanthanum series
† 90 – 103 Actinium series

$\begin{matrix} a \\ X \\ b \end{matrix}$ a = relative atomic mass (approx)
x = atomic symbol
b = atomic number

140 Ce Cerium 58	141 Pr Praseodymium 59	144 Nd Neodymium 60	145 Pm Promethium 61	150 Sm Samarium 62	152 Eu Europium 63	157 Gd Gadolinium 64	159 Tb Terbium 65	162 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