

Please check the examination details below before entering your candidate information

Candidate surname

Other names

Centre Number

Candidate Number

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Pearson Edexcel Level 3 GCE

Friday 21 June 2024

Morning (Time: 2 hours 30 minutes)

**Paper
reference**

9CH0/03



Chemistry

Advanced

PAPER 3: General and Practical Principles in Chemistry

You must have:

Scientific calculator, Data Booklet, ruler

Total Marks

Instructions

- Use **black** ink or ball-point pen.
- If pencil is used for diagrams/sketches/graphs it must be dark (HB or B).
- **Fill in the boxes** at the top of this page with your name, centre number and candidate number.
- Answer **all** questions.
- Answer the questions in the spaces provided
– *there may be more space than you need.*

Information

- The total mark for this paper is 120.
- The marks for **each** question are shown in brackets
– *use this as a guide as to how much time to spend on each question.*
- For the question marked with an **asterisk (*)**, marks will be awarded for your ability to structure your answer logically, showing the points that you make are related or follow on from each other where appropriate.
- A Periodic Table is printed on the back cover of this paper.

Advice

- Read each question carefully before you start to answer it.
- Show all your working in calculations and include units where appropriate.
- Check your answers if you have time at the end.

Turn over ►

P74455A

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Pearson

Answer ALL questions.

Write your answers in the spaces provided.

- 1** Relative isotopic mass is an important concept in chemistry.

- (a) Define relative isotopic mass.

(2)

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- (b) A sample of hydrogen gas is formed from atoms of three isotopes, ^1H , ^2H and ^3H .

Deduce the number of peaks due to molecular ions, H_2^+ , in the mass spectrum of this sample.

(1)

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(Total for Question 1 = 3 marks)

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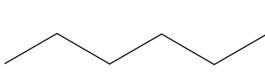
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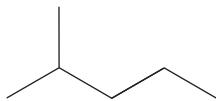
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- 2 This question is about the intermolecular forces in organic compounds.

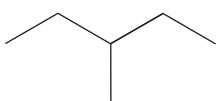
The structures of four isomeric alkanes are shown.



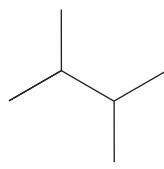
compound A



compound B



compound C



compound D

The only significant intermolecular forces between molecules of these compounds are London forces.

- (a) Explain how London forces arise.

(3)

- (b) Explain which of the compounds A–D will have the highest boiling temperature.

(2)

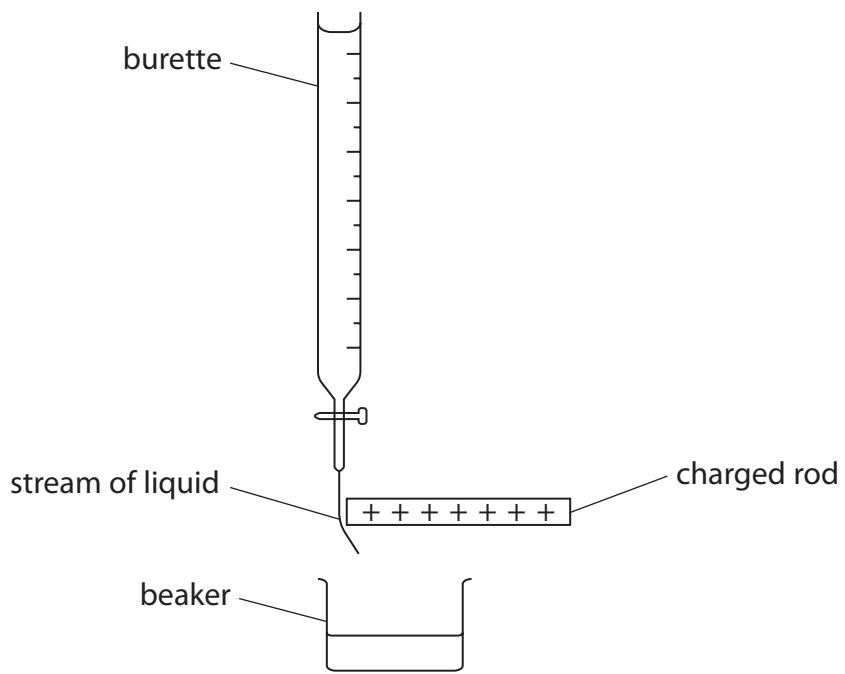


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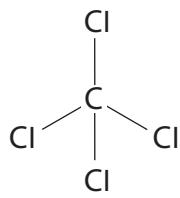
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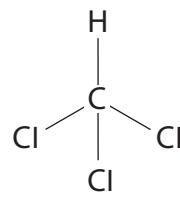
- (c) The apparatus shown may be used in an experiment to compare how far, if at all, a charged rod deflects different liquids.



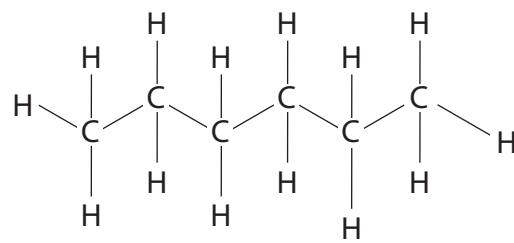
The structures of three liquids are shown.



CCl_4



CHCl_3



C_6H_{14}

Explain whether or not each of these liquids will be deflected in this experiment.

(3)

(Total for Question 2 = 8 marks)



- 3 A student carried out an experiment to identify the Group 1 metal ion, M^+ , in a sulfate, M_2SO_4 .

Procedure

Step 1 A 0.48 g sample of M_2SO_4 was added to about 50 cm^3 of deionised water in a beaker. The mixture was stirred with a glass rod until the sample dissolved completely.

Step 2 An excess of acidified barium chloride solution, $BaCl_2(aq)$, was added very slowly to the solution from Step 1 to ensure the crystals of barium sulfate that formed were as large as possible.

Step 3 The barium sulfate crystals were collected by gravity filtration onto a dry filter paper of known mass.

Step 4 The crystals were rinsed using deionised water.

Step 5 The filter paper containing the crystals was dried in a warm oven for an hour and then weighed.

Results

Mass of dry filter paper = 0.17 g

Mass of dry filter paper and barium sulfate crystals = 1.19 g

- (a) Give a reason, with reference to the filtration in Step 3, why the formation of **extremely** small crystals in Step 2 would lead to a lower mass of dry barium sulfate in Step 5.

(1)

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- (b) Explain why the crystals are rinsed in Step 4.

(2)

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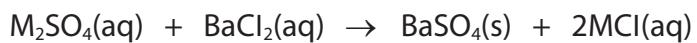
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- (c) Calculate the relative formula mass of M_2SO_4 and hence identify M.



[M_r of $BaSO_4$ = 233.4]

(3)

- (d) Another student carrying out the experiment placed the glass rod on the laboratory bench when the stirring was complete in Step 1.

- (i) Explain what effect this would have on the calculated value of the relative formula mass of M_2SO_4 .

(2)

- (ii) Describe how the student should amend the procedure.

(1)



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(e) Another way to identify a metal ion is a flame test.

(i) Explain what causes the flame colour in such a test.

(2)

(ii) Give one advantage and one disadvantage of using a flame test to identify a metal cation, rather than the method followed by the students in Steps 1–5.

(2)

(Total for Question 3 = 13 marks)

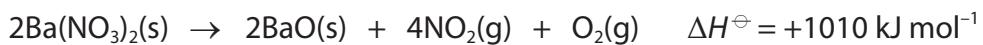
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4 This question is about the white crystalline solid, barium nitrate.

- (a) Barium nitrate decomposes under suitable conditions to form barium oxide, nitrogen dioxide and oxygen.



Standard molar entropy data related to this reaction are shown.

Substance	Standard molar entropy, $S^\ominus / \text{JK}^{-1} \text{ mol}^{-1}$
$\text{Ba}(\text{NO}_3)_2(\text{s})$	213.8
$\text{BaO}(\text{s})$	70.4
$\text{NO}_2(\text{g})$	240.0
$\text{O}_2(\text{g})$	205.0

- (i) Show that barium nitrate is thermally stable at 298 K.

(5)

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- (ii) Calculate the **minimum** temperature, in °C, at which it is thermodynamically feasible for barium nitrate to decompose.

Give your answer to an appropriate number of significant figures.

(3)

- (b) Explain why calcium nitrate is less thermally stable than barium nitrate.

(3)

- (c) A qualitative test for nitrate ions involves heating an alkaline solution of the suspected nitrate with aluminium.

- (i) Complete the ionic equation for this redox reaction, using oxidation numbers.
State symbols are not required.

(1)



- (ii) Describe a test to confirm that ammonia, NH₃, has been produced.

(1)

(Total for Question 4 = 13 marks)



- *5 Compare and contrast the reactions of bromine with benzene and of bromine with cyclohexene.

Consider the types of reaction that occur, the products that form, and any special conditions that may be required.

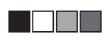
Justify your answers.

Detailed mechanisms for the reactions are **not** required.

(6)

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(Total for Question 5 = 6 marks)



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6 An ester **Q** has the molecular formula C₈H₁₆O₂.

- (a) When hydrolysed, 2.07 g of **Q** formed a carboxylic acid with a 78% yield.
Once separated, the carboxylic acid was neutralised using excess sodium hydrogencarbonate solution.



In this reaction, 269 cm³ of carbon dioxide gas was produced at room temperature and pressure.

Show that these data confirm that the relative molecular mass of **Q** is 144.

(3)

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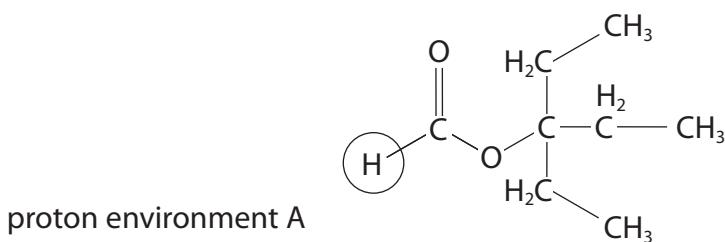
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- (b) The high resolution proton NMR spectrum of ester **Q** was obtained.
The structure of the ester **Q** is shown.



- (i) Explain the expected splitting pattern for the peak due to proton environment A circled in the structure.

(2)

- (ii) Label the structure to indicate the remaining equivalent proton environments in ester **Q**.

(1)

- (iii) Predict the chemical shifts, relative peak areas and splitting patterns in the high resolution proton NMR spectrum, due to the proton environments you have labelled.

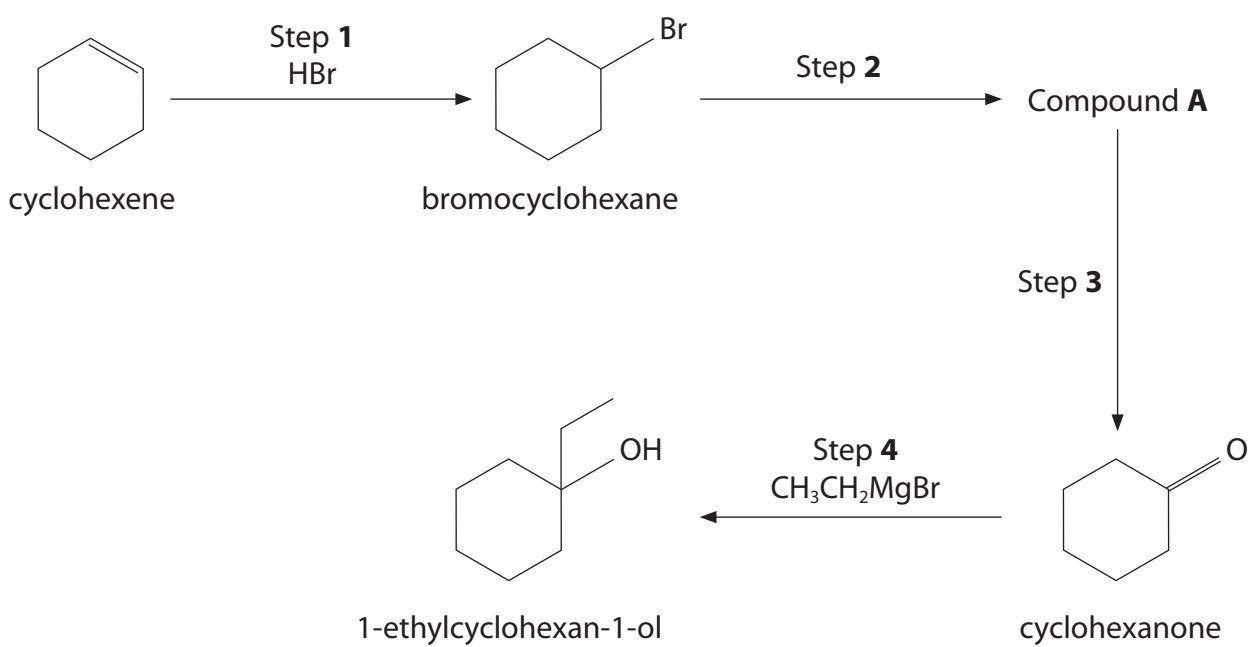
Do not consider proton environment A.

(3)

(Total for Question 6 = 9 marks)



- 7 The alcohol 1-ethylcyclohexan-1-ol can be synthesised from cyclohexene using the reaction scheme shown.



- (a) Draw the mechanism for Step 1.
Include curly arrows, and relevant lone pairs and dipoles.

(4)



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- (b) Devise a reaction scheme for Step **2** and Step **3** of the synthesis.
Include in your answer all reagents and conditions, the type of reaction in each step, and the structure of **A**.

(5)



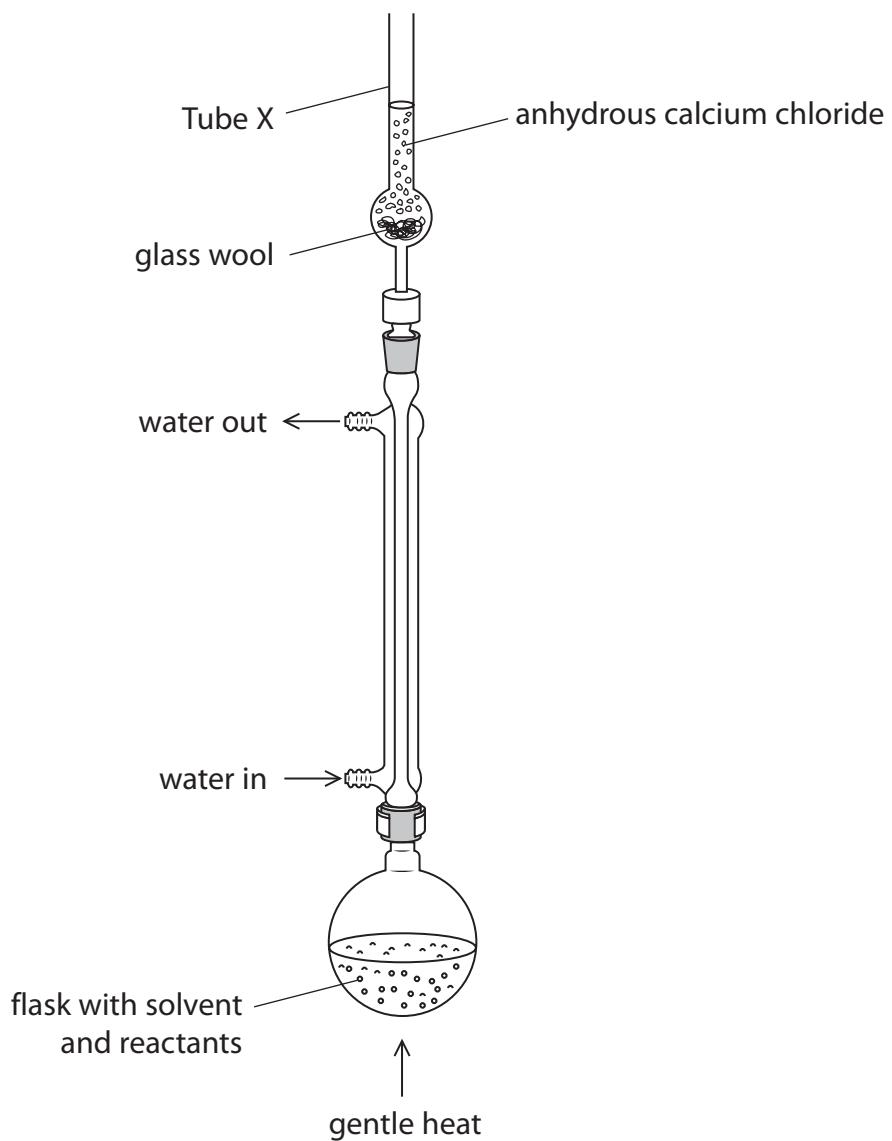
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- (c) The reaction to produce the $\text{CH}_3\text{CH}_2\text{MgBr}$ used in Step 4 requires bromoethane and magnesium to be warmed under reflux with a suitable solvent. The apparatus used for this reaction is shown.



- (i) Name the solvent used.

(1)



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(ii) Explain why the anhydrous calcium chloride in Tube X is necessary.

(2)

(iii) Give a reason why it is best practice for the water in the condenser to flow in the direction shown in the diagram.

(1)



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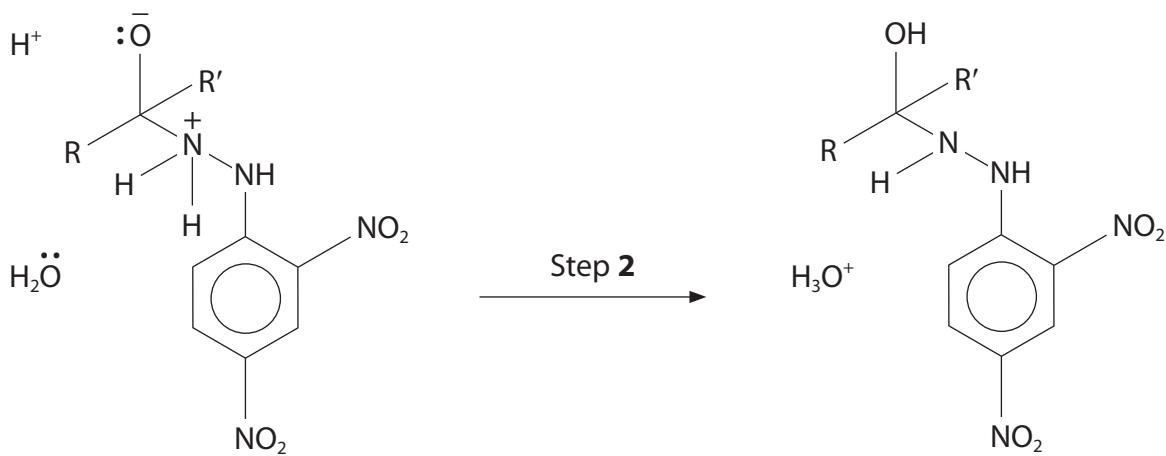
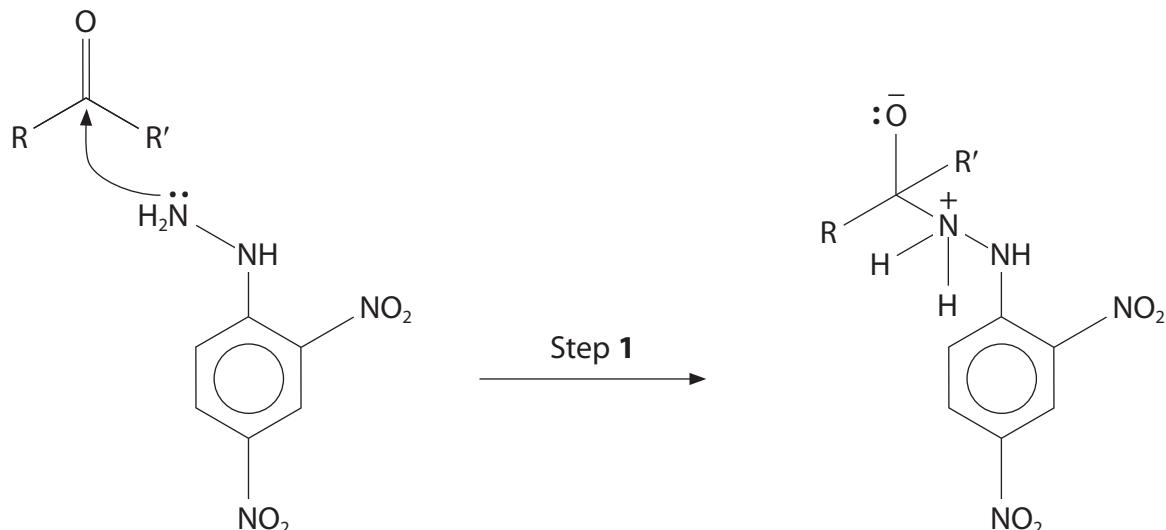
(d) Carbonyl compounds such as cyclohexanone can be identified by the reaction with 2,4-dinitrophenylhydrazine.

(i) Give the colour of the precipitate formed in this reaction.

(1)

(ii) Complete the simplified mechanism for the first two steps of this reaction by adding **four** curly arrows.

(4)



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(iii) Describe how you would obtain pure dry crystals from the precipitate.

(4)

(Total for Question 7 = 22 marks)



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8 This question is about acids and bases.

- (a) A student devised a method to determine the acid dissociation constant, K_a , of ethanoic acid, CH_3COOH , using a solution of the acid of unknown concentration.

The student's outline procedure is shown.

Step 1 Titrate 25.0 cm^3 of the ethanoic acid solution with a solution of ammonia of known concentration, using phenolphthalein to find the end-point.

Step 2 Add a further 25.0 cm^3 of the same ethanoic acid solution to the mixture from **Step 1**.

This gives a solution where the acid has been half-neutralised.

Step 3 Record the pH of the solution from **Step 2**.

- (i) Explain why this procedure would give invalid results.

(3)

- (ii) State how **Step 1** should be amended to give valid results.

(1)

- (iii) Show that when the solution has been half-neutralised, the acid dissociation constant is given by the expression

$$K_a = 10^{-\text{pH}}$$

(3)



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- (b) Calculate the mass of sodium ethanoate needed to be dissolved in 250 cm^3 of 0.520 mol dm^{-3} ethanoic acid to form a buffer solution with $\text{pH} = 4.48$

$[K_a \text{ for ethanoic acid} = 1.74 \times 10^{-5} \text{ mol dm}^{-3}]$

(5)



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- (c) A student mixes 50.0 cm^3 of $0.900 \text{ mol dm}^{-3}$ NaOH(aq) with 20.0 cm^3 of $0.400 \text{ mol dm}^{-3}$ $\text{H}_2\text{SO}_4(\text{aq})$.

Calculate the pH of the resultant solution.

[Ionic product of water, $K_w = 1.00 \times 10^{-14} \text{ mol}^2 \text{ dm}^{-6}$]

(5)

(Total for Question 8 = 17 marks)



- 9 Acidity in wine is mainly due to the dicarboxylic acid tartaric acid, $C_4H_6O_6$.

Sulfur dioxide is added to wine as an antioxidant, to prevent the colour and flavour of the wine from deteriorating. It also inhibits the growth of bacteria.

The concentration of tartaric acid and the concentration of sulfur dioxide present in wine can both be determined by titrations.

- (a) The procedure to find the total amount of tartaric acid in a white wine is shown.

Step 1 Pipette a 20.0 cm^3 sample of the wine into a conical flask.

Step 2 Add anti-bumping granules to the flask, and then place it on a preheated hot plate.

Step 3 Boil the sample to remove any dissolved gases and allow it to cool.

Step 4 Use 0.100 mol dm^{-3} sodium hydroxide solution, NaOH(aq) , in a burette to titrate the sample using a suitable indicator.

- (i) State a precaution you would take in carrying out Step 3.
Justify your answer.

(2)

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- (ii) Explain what effect the omission of Step 3 would have on the titre.

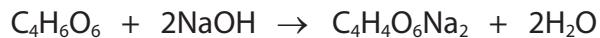
(2)

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(iii) A mean titre of 20.60 cm^3 was obtained.

Calculate the concentration, in g dm^{-3} , of tartaric acid in the white wine.



(3)

(iv) A student carrying out the experiment forgot to remove an air bubble from the tip of the burette **before** carrying out a titration. During the titration, the air bubble escaped.

Explain the effect this mistake would have on the value of this titre.

(2)



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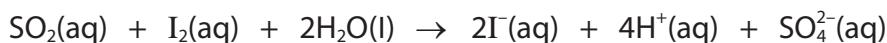
(b) A procedure to find the amount of sulfur dioxide, SO_2 , in a wine is shown.

Step 1 50.0 cm^3 of a white wine is added to a conical flask containing 5 cm^3 of dilute sulfuric acid and a few drops of starch indicator.

Step 2 A spatula measure of sodium hydrogencarbonate is then added to reduce the loss of sulfur dioxide from the flask.

Step 3 The contents of the flask are titrated rapidly using $0.0100 \text{ mol dm}^{-3}$ iodine solution, $\text{I}_2(\text{aq})$.

The equation for the reaction that takes place is shown.



- (i) A student carrying out the experiment obtained a titre of 11.80 cm^3 .
The maximum permitted level of sulfur dioxide in white wine in the UK is 200 mg dm^{-3} .

Show by calculation that the concentration of sulfur dioxide in the white wine is below the permitted maximum.

(4)



P 7 4 4 5 5 A 0 2 5 3 2

- (ii) Explain how the addition of sodium hydrogencarbonate in Step 2 reduces the loss of sulfur dioxide from the flask.

(2)

(Total for Question 9 = 15 marks)



- 10** The progress of the reaction between ethyl ethanoate and water with a hydrochloric acid catalyst can be followed in an experiment using a titrimetric method. An outline procedure is shown.

Step 1 Add 100 cm³ of 0.500 mol dm⁻³ HCl(aq) to a conical flask.

Step 2 Using a pipette, add 5.0 cm³ of ethyl ethanoate to the conical flask while mixing.

Step 3 Immediately remove a 5.0 cm³ sample of the reaction mixture from Step 2 and add it to a separate conical flask containing crushed ice.

Step 4 Titrate the removed sample using 0.200 mol dm⁻³ NaOH(aq).

Step 5 Repeat Steps 3 and 4, removing 5.0 cm³ samples every 10 minutes.

Step 6 After 60 minutes, heat the remaining reaction mixture under reflux for 15 minutes.

Step 7 Remove a 5 cm³ sample of the reaction mixture after completing Step 6, titrating this sample with 0.200 mol dm⁻³ NaOH(aq).

- (a) (i) Explain the purpose of the crushed ice in Step 3.

(2)

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- (ii) Explain why Step 6 is carried out before the final sample is removed and titrated in Step 7.

(2)

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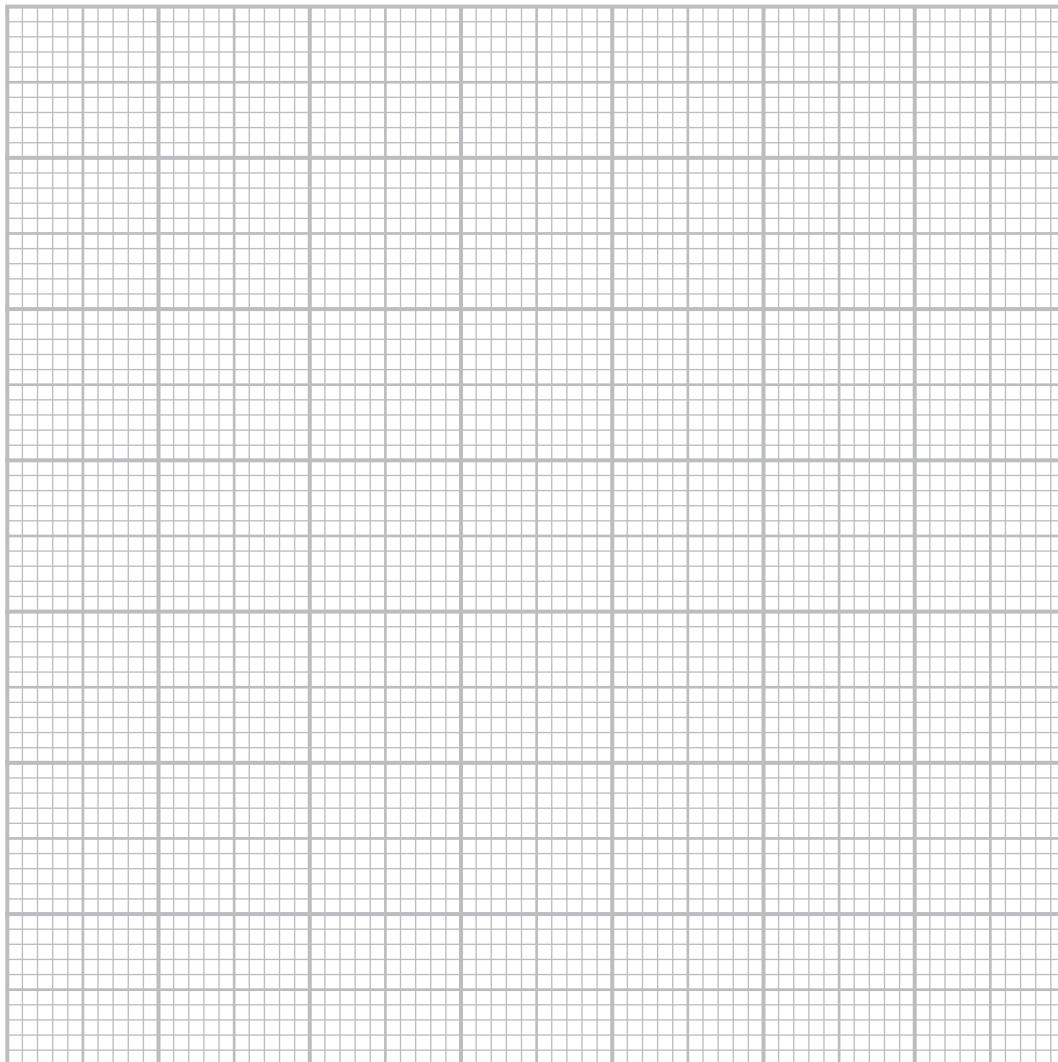
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- (b) Some data from the experiment are shown. V_{final} is the titre volume from Step 7, V_t is the titre volume at time t.

Time reaction sample is added to crushed ice / min	0	10	20	30	40	50	60
$V_{\text{final}} - V_t$ of sodium hydroxide / cm ³	11.60	6.50	3.80	1.80	0.90	0.80	0.80

- (i) Plot a graph of the $V_{\text{final}} - V_t$ of sodium hydroxide against the time the reaction sample is added to the crushed ice.

(3)



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- (ii) Explain why $V_{\text{final}} - V_t$ is plotted rather than the concentration of sodium hydroxide solution for each titre.

(2)

- (iii) Determine two half-lives for the reaction, showing your working on the graph.

(2)

- (iv) Deduce the order of the reaction, using your answer from (b)(iii) to justify your deduction.

(1)



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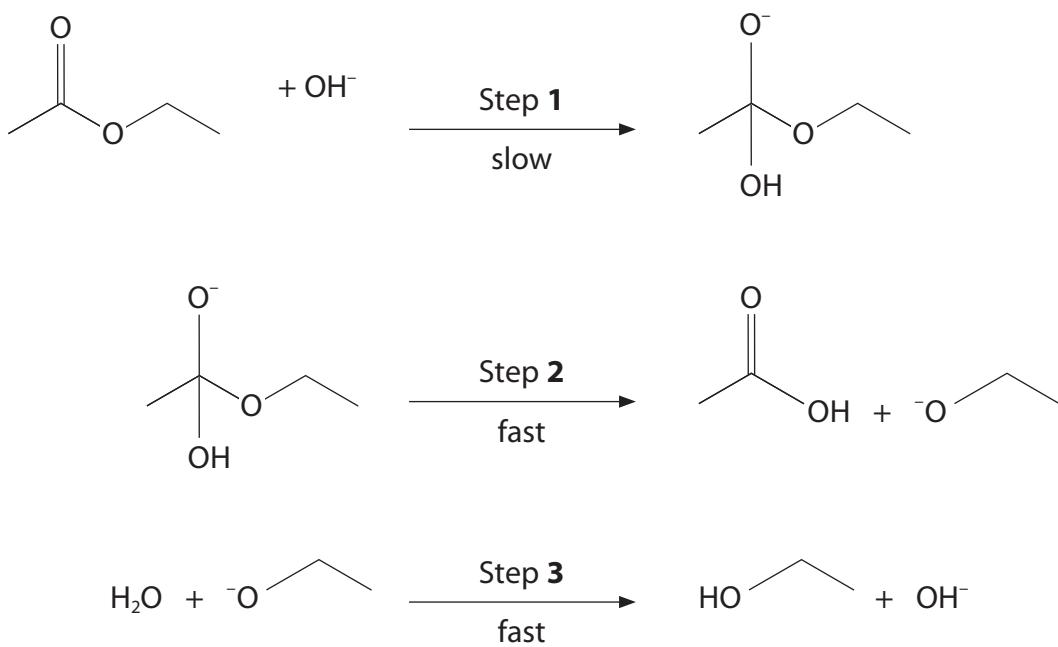
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(c) Ethyl ethanoate can also be hydrolysed under alkaline conditions.

A student researching the mechanism for the reaction found this example.



Deduce a rate equation that is consistent with this reaction mechanism.
Justify your answer.

(2)

(Total for Question 10 = 14 marks)

TOTAL FOR PAPER = 120 MARKS



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The Periodic Table of Elements

1 2

1.0	H	hydrogen
1		

Key

relative atomic mass
atomic symbol
name
atomic (proton) number

(1)	(2)	(3)	(4)	(5)	(6)	(7)	(8)	(9)	(10)	(11)	(12)	(13)	(14)	(15)	(16)	(17)	(18)
6.9 Li lithium 3	9.0 Be beryllium 4	12.0 B boron 5	19.0 C carbon 6	24.0 N nitrogen 7	31.0 O oxygen 8	32.1 F fluorine 9	35.5 Ne neon 10	40.0 He helium 2									
23.0 Na sodium 11	24.3 Mg magnesium 12	30.0 Ca calcium 20	45.0 Sc scandium 21	50.9 Ti titanium 22	52.0 V vanadium 23	54.9 Cr chromium 24	55.8 Mn manganese 25	58.9 Fe iron 26	63.5 Co cobalt 27	65.4 Cu copper 29	69.7 Ni nickel 28	72.6 Ga gallium 31	74.9 Ge germanium 32	79.0 As arsenic 33	83.8 Br bromine 35	Kr krypton 36	
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	58.7 Ni nickel 28	63.5 Co cobalt 27	65.4 Cu copper 29	69.7 Ga gallium 31	72.6 Ge germanium 32	79.0 Se selenium 34	83.8 Kr krypton 36	Xe xenon 36		
85.5 Rb rubidium 37	87.6 Sr strontium 38	91.2 Y yttrium 39	92.9 Zr zirconium 40	95.9 Nb niobium 41	95.9 Mo molybdenum 42	95.9 Tc technetium 43	95.9 Ru ruthenium 44	101.1 Rh rhodium 45	102.9 Pd palladium 46	106.4 Ag silver 47	112.4 Cd cadmium 48	114.8 In indium 49	118.7 Sn tin 50	121.8 Sb antimony 51	126.9 Te tellurium 52	131.3 I iodine 53	
132.9 Cs caesium 55	137.3 Ba barium 56	138.9 La* lanthanum 57	178.5 Hf hafnium 72	180.9 Ta tantalum 73	183.8 W tungsten 74	186.2 Re rhenium 75	190.2 Os osmium 76	192.2 Ir iridium 77	195.1 Pt platinum 78	197.0 Au gold 79	200.6 Hg mercury 80	204.4 Tl thallium 81	207.2 Pb lead 82	209.0 Bi bismuth 83	210.0 Po polonium 84	222.0 At astatine 85	
[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	[268] Hs hassium 108	[277] Mt meitnerium 109	[268] Ds darmstadtium 110	[271] Rg roentgenium 111							
140 Ce cerium 58	141 Pr praseodymium 59	144 Nd neodymium 60	144 Pm promethium 61	150 Sm samarium 62	152 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	[245] Cf californium 98	[253] Es einsteinium 99	[256] Fm fermium 100	[254] Md mendelevium 101	[254] No nobelium 102	[257] Lr lawrencium 103				

* Lanthanide series
* Actinide series

Elements with atomic numbers 112-116 have been reported but not fully authenticated

