Please check the examination details be	low before ente	ering your candidate information
Candidate surname		Other names
Centre Number Candidate N	umber	
Pearson Edexcel Inter	nation	al Advanced Level
Time 1 hour 20 minutes	Paper reference	WCH16/01
Chemistry		0
International Advanced Lo	evel	
UNIT 6: Practical Skills in	Chemist	try II
		,
You must have:		Total Marks
Scientific calculator, ruler		

## **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 50.
- The marks for **each** question are shown in brackets
  - use this as a guide as to how much time to spend on each question.
- You will be assessed on your ability to organise and present information, ideas, descriptions and arguments clearly and logically, including your use of grammar, punctuation and spelling.
- 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.
- Try to answer every question.
- Check your answers if you have time at the end.

Turn over ▶







# Answer ALL the questions. Write your answers in the spaces provided. Solution **A** and solution **B** are aqueous solutions. The compound in each solution contains one cation and one anion. The cations are different but the anions are the same. Both solutions are green. (a) Give the formulae of **three** cations which could be responsible for the green colour. (2) (b) A student added dilute aqueous sodium hydroxide, drop by drop, to samples of each solution. Initially a green precipitate formed in both solutions. More aqueous sodium hydroxide was added until the sodium hydroxide was present in excess. The precipitate produced from solution **A** was insoluble and turned brown on standing. The precipitate produced from solution **B** dissolved to give a dark green solution **C**. (i) Give the formula of the green precipitate from solution **A**. (1)

(c) The student added hydrogen peroxide solution to **C** and warmed the mixture, which turned yellow. The yellow solution was boiled to remove excess hydrogen peroxide, cooled and then pure ethanoic acid was added, drop by drop. The yellow solution gradually turned orange.

(ii) Give the formula of the species responsible for the dark green colour in

(i) State the type of reaction involved when the dark green solution turned yellow.

(ii) Identify, by name or formula, the ion responsible for the final orange colour of the solution.

(1)

(1)

(1)



solution C.

<ul> <li>(iii) Pure ethanoic acid is corrosive.</li> <li>Identify the appropriate control measure to reduce the risk associated w this hazard.</li> <li>Assume the student carried out the addition in a fume cupboard, wearin safety glasses and a lab coat.</li> </ul>	
(iv) Describe the corrosive hazard label present on a bottle containing pure ethanoic acid. You may use a diagram in your answer.	(2)



(d)	The student attempted to identify the anion present in the green solution <b>A</b> . The student added dilute nitric acid and a few drops of aqueous silver nitrate to about 2 cm³ of solution <b>A</b> . A pale precipitate formed, which the student thought might be white or cream coloured.	
	(i) Identify, by name or formula, <b>two</b> anions which might give this precipitate.	(1)
	(ii) The precipitate formed in (d)(i) was separated from the mixture and aqueous ammonia was added.	
	Describe how this test allows the student to distinguish between the two anions.	
	two amons.	(2)
	(iii) Explain why, with solution <b>A</b> , the precipitate must be separated before adding aqueous ammonia.	
		(2)
	(Total for Operation 1 – 14 may	rks)
	(Total for Question 1 = 14 mai	ik5)



**2** The reaction between the tertiary halogenoalkane 2-chloro-2-methylpropane and hydroxide ions to form 2-methylpropan-2-ol is shown.

$$(CH_3)_3CCI + OH^- \rightarrow (CH_3)_3COH + CI^-$$

The progress of the reaction can be followed by titrating portions of the reaction mixture with a solution of hydrochloric acid of known concentration.

An experiment to determine the order of the reaction was carried out.

#### **Procedure**

Step **1** A flask containing 250 cm<sup>3</sup> of an ethanolic solution of 2-chloro-2-methylpropane, with a concentration 0.100 mol dm<sup>-3</sup>, was placed in a water bath at 25 °C.

A similar flask, containing  $250\,\mathrm{cm^3}$  of aqueous sodium hydroxide, with a concentration  $0.100\,\mathrm{mol\,dm^{-3}}$ , was placed in the same water bath. The temperature of each solution was allowed to reach  $25\,^{\circ}\mathrm{C}$ .

- Step **2** Seven conical flasks were prepared, each containing about 50 cm<sup>3</sup> of propanone.
- Step 3 The reaction was started by mixing the two solutions from the water bath in a large flask. The flask was returned to the water bath.

  A timer was started as the solutions were mixed.
- Step **4** At intervals, 25.0 cm<sup>3</sup> samples of the reaction mixture were transferred to a conical flask containing propanone. The time was noted.
- Step **5** Each sample was immediately titrated with a solution of hydrochloric acid, of concentration 0.0500 mol dm<sup>-3</sup>, using methyl orange as the indicator.

(ii) Explain the effect on the volume of hydrochloric acid required if the titration

(a) (i) Give the expected colour **change** at the end-point.

was not carried out immediately after Step **4**. (2)



(1)

(iii) Explain how the use of aqueous ethanol in the reaction mixture allows the reaction in Step 3 to proceed at a relatively fast rate.

A description of the forces involved is not required.

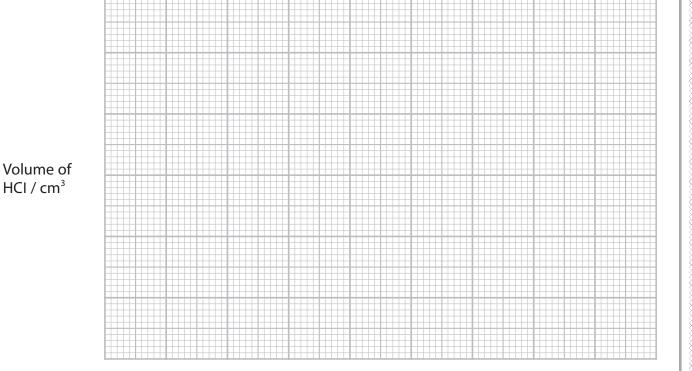
(2)

(b) An experiment was carried out using this procedure. The results are shown.

Time / s	50	50 320 750 1200 2050				3180	4020
Volume of HCI / cm <sup>3</sup>	24.5	22.0	18.5	15.5	11.0	7.0	5.0

(i) Plot a graph of the data using the axes given.

(2)



Time / s



(ii) Determine two successive half-lives for the reaction. You must show your working on the graph.	(2)
(iii) State the order of this reaction. Justify your answer.	(1)
(Total for Question 2 = 10	marks)

Moss in lawns can be treated with 'lawn sand', a mixture that contains sand and a double salt, ammonium iron(II) sulfate,  $(NH_4)_2Fe(SO_4)_2\cdot 6H_2O$ . The percentage by mass of  $Fe^{2+}$  in the mixture can be found by titration with a solution of potassium manganate(VII) of known concentration.

## **Procedure**

4.50 g of lawn sand was accurately weighed in a 250 cm<sup>3</sup> conical flask. The sample was shaken with 50 cm<sup>3</sup> of dilute sulfuric acid (an excess). The resulting mixture was titrated with potassium manganate(VII) solution of concentration 0.0200 mol dm<sup>-3</sup>.

The titration volume was 40.35 cm<sup>3</sup>.

(á	a) (I)	State what would happen in the titration if the mixture was <b>not</b> acidified.	(1)
	(ii)	Neither hydrochloric acid nor nitric acid can be used to acidify the titration mixture.	
		For each acid give a reason why it cannot be used.	(2)



(b) The ionic half-equations for the reactions of the iron(II) ions and the manganate(VII) ions are shown.

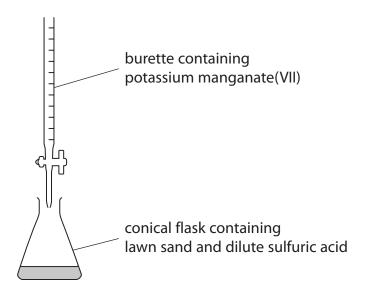
$$Fe^{2+} \ \to \ Fe^{3+} \ + \ e^{-}$$
 
$$MnO_{4}^{-} \ + \ 8H^{+} \ + \ 5e^{-} \ \to \ Mn^{2+} \ + \ 4H_{2}O$$

Calculate the percentage by mass of  $Fe^{2+}$  in the sample of lawn sand.

(4)

(3)

(c) The titration is carried out using the apparatus shown.



At the end-point, the solution changes to a permanent pale pink.

To repeat the experiment, the burette is refilled with potassium manganate(VII) and the procedure carried out again.

Give **three** ways in which a titration is carried out to give the most accurate possible burette reading.

Assume that the equipment is the same, the burette is clamped in a vertical position and that the titration volumes are read at eye level from the bottom of the meniscus.

(Total for Question 3 = 10 marks)



**BLANK PAGE** 



(2)

4 The compound 2-ethanoylaminobenzoic acid can be prepared by the reaction of 2-aminobenzoic acid and ethanoyl chloride.

## **Procedure**

- Step **1** Add 5.00 g of 2-aminobenzoic acid and a few anti-bumping granules to a dry 100 cm<sup>3</sup> pear-shaped flask and fit a reflux condenser.
- Step 2 Add 10 cm<sup>3</sup> of ethanoyl chloride (an excess) by pouring it slowly down the condenser.
- Step 3 Gradually bring the mixture to boil and heat under reflux for 15 minutes.
- Step **4** Allow the mixture to cool and slowly add 10 cm<sup>3</sup> of water down the condenser.
- Step 5 Heat the solution slowly until boiling.
- Step **6** Allow the solution to cool to room temperature.
- Step **7** Collect the crystals of 2-ethanoylaminobenzoic acid by filtration under reduced pressure.
- Step 8 Recrystallise the impure product from a mixture containing equal volumes of ethanoic acid and water.
- (a) Give **two** reasons why it is often necessary to heat a reaction under reflux as in Step **3**.

(b) Explain why, in Step <b>4</b> , water is added <b>slowly</b> to the cooled solution.	(2)
(c) Draw a labelled diagram of the apparatus used for filtration under reduced pressure in Step <b>7</b> .	(3)

- (d) The melting temperature may be used to confirm the identity of the crystals.
  - (i) Draw a labelled diagram of the apparatus you would use to measure the melting temperature.

(2)

(ii) State how this melting temperature determination would show that a pure sample of 2-ethanoylaminobenzoic acid had been prepared.

(2)

(e) Show by calculation that, in the preparation, 10 cm<sup>3</sup> of ethanoyl chloride is an excess compared to 5.00 g of 2-aminobenzoic acid.

## Data

 $Molar \ mass \ / \ g \ mol^{-1} \qquad CH_3COCI = 78.5 \qquad C_6H_4(NH_2)COOH = 137$ 

Density / g cm<sup>-3</sup>  $CH_3COCI = 1.1$ 

(2)

(f) A student carried out the preparation using the amounts of reagents given in (e) and obtained a yield of 56.7 % of 2-ethanoylaminobenzoic acid.

Calculate the mass of 2-ethanoylaminobenzoic acid obtained.

(3)

(Total for Question 4 = 16 marks)

**TOTAL FOR PAPER = 50 MARKS** 



Md No Lr mendetevium nobelium tawrencium

103

102

101

[257]

[254]

[256] 69

lutetium

ytterbium

Tm

167 Er erbium

20

68

3

ted		Radon 86	[222]	Xe xenon 54	131.3	krypton 36	8 2 8	39.9 Ar argon	10	20.2	He helium 2	0 (8)
been repor		At astatine 85	[210]	lodine 53	126.9	Br bromine 35	70.0	35.5 CI chlorine 17	fluorine 9	19.0	(17)	7
116 have t		Po polonium 84	[506]	Te tellurium 52	127.6	Se selenium 34	70.07	32.1 S sulfur 16	oxygen 8	16.0	(16)	•
tomic numbers 112-116 hav but not fully authenticated		Bi bismuth 83	209.0	Sb antimony 51	121.8	AS arsenic 33	74.0	31.0 Phosphorus	nitrogen 7	14.0	(15)	'n
atomic nur but not fo		Pb tead 82	207.2	<b>S</b> 0 th	118.7	Ge germanium 32	77.4	Si siltcon	carbon 6	12.0	(14)	4
Elements with atomic numbers 112-116 have been reported but not fully authenticated		TI thallium 81	204.4	Indium 49	114.8	Ga gallium 31	407	27.0 Al atuminium	boron 5	10.8	(13)	m
Elem		Hg mercury 80	200.6	Cd cadmium 48	112.4	Zn zinc 30	1 37	(2)				
Rg centgenium	[272]	Sold 79	197.0	Ag silver 47	107.9	Cu copper 29	3 67	(11)				
Mt Ds Rg mettorerium damstadtum roentgenium	[271]	Pt platinum 78	195.1	Pd palladium 46	106.4	<b>Ni</b> nicket 28	202	(01)				5
Mt	[268] Mt	iridium 77	192.2	Rh rhodium 45	102.9	Co cobalt 27	0 02	(6)				
-	265	Os osmium 76	190.2	Ru ruthenium 44	101.1	Fe iron 26	0 22	(8)			1.0 Hydrogen	
-	-	Re rhenium 75	186.2		[86]	Mn manganese 25	0 73	8				)
Sg seaborgium	[592]	tungsten 74	183.8	Mo Tc motybdenum technetium 42 43	6.26	E	200	9	ımber	mass		
- E	[292]	Ta tantalum 73	180.9	Nb niobium n	67.6	V vanadium 23	0 00	9	name atomic (proton) number	relative atomic mass	Key	
E	[261]	Hf hafnium 72	178.5	Zr zirconium 40	91.2	Ti titanium 22	47.0	3	atomic	relativ		
. E	[227]	La* lanthanum 57	138.9	Y yttrium 2	88.9	E	0 37	8				
	[526]	Ba barium k	137.3	Sr strontium 38	9.78	Ca calcium s	* 07	24.3 Mg magnesium	beryllium 4	0.6	(2)	7
Fr	[223]	CS caesium 55	132.9	Rb rubidium 37	85.5	K potassium 19	* 00	Na sodium	E	6.9	(1)	-

P 7 1 8 8 9 A 0 1 6 1 6

[253] Fm femium 100 Bk Cf Es berkelium catifornium einstenium fe 97 98 oo 165 Ho holmium 67 
 Pr
 Nd
 Pm
 Sm
 Eu
 Gd
 Tb
 Dy

 præecodymium neodymium promethium samarium
 samarium
 europium
 gadotinium
 terbium
 dysprosium
 99 65 £ # % [247] 64 Np Pu Am Pu Am Putonium plutonium americium 63 62 [147] 61 uranium 9 238 92 protactinium [231] Pa 29 6 thorium 6 5 5 cerium 232 8 28 · Lanthanide series Actinide series