



Cambridge International Examinations

Cambridge International Advanced Subsidiary and Advanced Level

CANDIDATE NAME										
CENTRE NUMBER						CANDIDATE NUMBER				_
CHEMISTRY									9701/33	3
Paper 3 Advance	ced Prac	tical Skill	ls 1			Od	tober/l	Noven	nber 2016	3
									2 hours	3
Candidates ans	wer on th	he Quest	ion Pap	oer.						
Additional Mater	rials:	As liste	d in the	Cor	nfidential Instructions					

READ THESE INSTRUCTIONS FIRST

Write your Centre number, candidate number and name on all the work you hand in.

Give details of the practical session and laboratory where appropriate, in the boxes provided.

Write in dark blue or black pen.

You may use an HB pencil for any diagrams or graphs.

Do not use staples, paper clips, glue or correction fluid.

DO **NOT** WRITE IN ANY BARCODES.

Answer all questions.

Electronic calculators may be used.

You may lose marks if you do not show your working or if you do not use appropriate units.

Use of a Data Booklet is unnecessary.

Qualitative Analysis Notes are printed on pages 10 and 11.

A copy of the Periodic Table is printed on page 12.

At the end of the examination, fasten all your work securely together. The number of marks is given in brackets [] at the end of each question or part question.

Session
Laboratory

For Examiner's Use		
1		
2		
3		
Total		

This document consists of 11 printed pages and 1 blank page.



1 Hydrogen peroxide, H_2O_2 , can be oxidised to give oxygen, O_2 . This reaction happens rapidly in the presence of acidified potassium manganate(VII), KMnO₄.

$$2MnO_4^{-}(aq) + 5H_2O_2(aq) + 6H^+(aq) \rightarrow 2Mn^{2+}(aq) + 5O_2(g) + 8H_2O(I)$$

You will determine the concentration of a solution of hydrogen peroxide. You will first dilute the solution and then carry out a titration using acidified potassium manganate(VII), KMnO₄.

FA 1 is aqueous hydrogen peroxide, H_2O_2 .

FA 2 is 0.0200 mol dm⁻³ potassium manganate(VII), KMnO₄.

FA 3 is 1.0 mol dm⁻³ sulfuric acid, H₂SO₄.

(a) Method

Dilution

- Pipette 25.0 cm³ of FA 1 into the 250 cm³ volumetric flask.
- Add distilled water to make 250 cm³ of solution and shake the flask thoroughly.
- Label this diluted solution of hydrogen peroxide FA 4.

Titration

- Fill the burette with FA 2.
- Rinse the pipette thoroughly with distilled water and then with a little **FA 4**.
- Pipette 25.0 cm³ of **FA 4** into a conical flask.
- Use the measuring cylinder to add 25 cm³ of **FA 3** to the conical flask.
- Perform a rough titration and record your burette readings in the space below.

The rough titre	is		cm ³
-----------------	----	--	-----------------

- Carry out as many accurate titrations as you think necessary to obtain consistent results.
- Make certain any recorded results show the precision of your practical work.
- Record in a suitable form below all of your burette readings and the volume of FA 2 added in each accurate titration.

Keep FA 3 and FA 4 for use in Question 2.

I	
II	
III	
IV	
V	
VI	
VII	
	II III IV V VI

[7]

(b)		m your accurate titration results, obtain a suitable value for the volume of FA 2 to be used our calculations. Show clearly how you obtained this value.
		25.0 cm ³ of FA 4 required cm ³ of FA 2 . [1]
(c)	Cal	culations
		ow your working and appropriate significant figures in the final answer to each step of your culations.
	(i)	Calculate the number of moles of potassium manganate(VII) present in the volume of $\textbf{FA 2}$ calculated in (b).
		moles of KMnO ₄ = mol
	(ii)	Use your answer to (i) and the equation at the top of page 2 to calculate the number of moles of hydrogen peroxide present in 25.0 cm³ of FA 4 .
		moles of H_2O_2 = mol
	(iii)	Calculate the concentration, in mol dm $^{-3}$, of H_2O_2 in FA 4 .
		concentration of H_2O_2 in FA 4 = mol dm ⁻³
	(iv)	Calculate the concentration, in mol dm $^{-3}$, of H_2O_2 in FA 1 .
		concentration of H_2O_2 in FA 1 = mol dm ⁻³ [4]
		[Total: 12]

2 In Question 1, hydrogen peroxide was oxidised to form oxygen. Hydrogen peroxide can also be reduced to form water. This occurs, for example, when hydrogen peroxide, H_2O_2 , reacts with iodide ions, $I^-(aq)$, to form iodine, $I_2(aq)$.

$$H_2O_2(aq) + 2H^+(aq) + 2I^-(aq) \rightarrow I_2(aq) + 2H_2O(I)$$

The rate of this reaction can be measured by adding acidified hydrogen peroxide, H_2O_2 , to a mixture of iodide ions, I^- , thiosulfate ions, $S_2O_3^{2-}$, and starch indicator. As the iodine is produced, it reacts immediately with the thiosulfate ions and is reduced back to iodide ions.

$$I_2(aq) + 2S_2O_3^2(aq) \rightarrow 2I^-(aq) + S_4O_6^2(aq)$$

When all the thiosulfate has reacted, the iodine then turns the starch indicator blue-black. The rate of reaction may be determined by timing how long it takes the reaction mixture to turn blue-black.

In this experiment you will measure two reaction times and use one of these to calculate the concentration of the thiosulfate ions in **FA 6**.

FA 3 is 1.0 mol dm⁻³ sulfuric acid, H₂SO₄.

FA 4 is the diluted hydrogen peroxide, H_2O_2 , you prepared in **1(a)**.

FA 5 is 1.0 mol dm⁻³ potassium iodide, KI.

FA 6 is aqueous sodium thiosulfate, $Na_2S_2O_3$.

starch indicator

Read through the instructions carefully before starting any practical work.

(a) Method

- Use the measuring cylinder to transfer 20 cm³ of **FA 3** into the 100 cm³ beaker.
- Use the measuring cylinder to add 10 cm³ of **FA 4** into the 100 cm³ beaker.
- Rinse the measuring cylinder with distilled water and drain.
- Use the measuring cylinder to transfer 20 cm³ of **FA 5** into the 250 cm³ beaker.
- Use the measuring cylinder to transfer 20 cm³ of **FA 6** into the 250 cm³ beaker.
- Use the measuring cylinder to transfer 10 cm³ of starch indicator into the 250 cm³ beaker.
- Add the contents of the 100 cm³ beaker to the 250 cm³ beaker and start timing immediately.
- Stir the mixture once and place the beaker on a white tile.
- Stop timing as soon as the solution turns blue-black.
- Record this reaction time to the nearest second.
- Rinse both beakers and drain.

Keep all FA solutions and starch indicator for use in (c).

reaction time = s [2]

(b) Calculations

Show your working and appropriate significant figures in the final answer to **each** step of your calculations.

(i) The expression for the rate of reaction is shown.

The concentration of iodine at the reaction time is the concentration of iodine that would have been present at the time the blue-black colour appeared if no thiosulfate ions had been added.

In this experiment, you should assume the rate = 2.61×10^{-5} mol dm⁻³ s⁻¹.

Use this value for the rate to calculate the concentration of iodine that would have been present at the reaction time if no thiosulfate ions had been added.

concentration of
$$I_2$$
 = $mol \, dm^{-3}$

(ii) Use your answer to (i) and the total reaction volume, to calculate the number of moles of iodine that would have been present at the reaction time if no thiosulfate ions had been added.

(If you were unable to calculate an answer to (i), you should use the value of 1.32×10^{-3} mol dm⁻³. This may not be the correct value.)

moles of
$$I_2$$
 = mol

(iii) Calculate the number of moles of thiosulfate ions that reacted with the moles of I_2 , calculated in (ii).

$$I_2(aq) + 2S_2O_3^{2-}(aq) \rightarrow 2I^{-}(aq) + S_4O_6^{2-}(aq)$$

moles of
$$S_2O_3^{2-}$$
 = mol

(iv) Use your answer to (iii) to calculate the concentration of thiosulfate ions in FA 6.

concentration of
$$S_2O_3^{2-}$$
 in **FA 6** = mol dm⁻³

(c)	Repeat the	experiment in	(a)	usina th	e followina	quantities	of (each reagent.
\ ⁻ /			\ · /				_	

add to the 100 cm³ beaker
 20 cm³ of FA 3

10 cm³ of **FA 4**

• add to the 250 cm³ beaker

20 cm³ of **FA 5** 10 cm³ of **FA 6**

10 cm³ of starch indicator

10 cm³ of distilled water

- Add the contents of the 100 cm³ beaker to the 250 cm³ beaker and start timing immediately.
- Stir the mixture once and place the beaker on a white tile.
- Stop timing as soon as the solution turns blue-black.
- Record this reaction time to the nearest second.

	reaction time = s [1]
(d) (i)	Explain the relationship between the value of the reaction time in (a) and the value of the reaction time in (c) .
(ii)	A student states that the error in the total volume of the reaction mixture in (a) is the same as the error in the total volume of the reaction mixture in (c) .
	State whether or not you agree with the student and explain your answer.

[Total: 11]

[4]

3 Qualitative Analysis

At each stage of any test you are to record details of the following.

- colour changes seen
- the formation of any precipitate
- the solubility of such precipitates in an excess of the reagent added

Where gases are released they should be identified by a test, **described in the appropriate place in your observations**.

You should indicate clearly at what stage in a test a change occurs.

No additional tests for ions present should be attempted.

If any solution is warmed, a boiling tube MUST be used.

Rinse and reuse test-tubes and boiling tubes where possible.

Where reagents are selected for use in a test, the name or correct formula of the element or compound must be given.

(a) FA 7, FA 8, FA 9 and FA 10 are solutions that each contain a metal cation from those listed in the Qualitative Analysis Notes on page 10.

Carry out tests using reagents that will allow you to identify which metal ions are present. Draw a single table to record your observations.

Complete the table below with your conclusions.

solution	FA 7	FA 8	FA 9	FA 10
metal ion present				

(b) Carry out the following test and record your observations.

test	observations
To a 1 cm depth of FA 10 in a boiling tube add a 0.5 cm depth of 20 'vol' hydrogen peroxide, then add one drop of aqueous sodium hydroxide.	

[2]

(c) FA 11 is a solid that contains one anion from those listed in the Qualitative Analysis Notes on page 11.

Place all of FA 11 into a boiling tube and add a 3 cm depth of distilled water. Stir until all the solid has dissolved.

(i) Carry out the following test and record your observations.

(ii) Use your observations from (i) to identify the anion in FA 11.

test	observations
To a 1 cm depth of the solution of FA 11 in a test-tube add dilute sulfuric acid.	

(iii)	Use the remaining solution of FA 11 in the boiling tube to carry out a further test to support
	vour identification of the anion.

Record details of this test.

The anion is

[4]

[Total: 17]

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Qualitative Analysis Notes

Key: [ppt. = precipitate]

1 Reactions of aqueous cations

ion	reaction with									
ion	NaOH(aq)	NH ₃ (aq)								
aluminium, Al³+(aq)	white ppt. soluble in excess	white ppt. insoluble in excess								
ammonium, NH₄⁺(aq)	no ppt. ammonia produced on heating	_								
barium, Ba²+(aq)	no ppt. (if reagents are pure)	no ppt.								
calcium, Ca ²⁺ (aq)	white ppt. with high [Ca ²⁺ (aq)]	no ppt.								
chromium(III), Cr³+(aq)	grey-green ppt. soluble in excess	grey-green ppt. insoluble in excess								
copper(II), Cu²+(aq)	pale blue ppt. insoluble in excess	blue ppt. soluble in excess giving dark blue solution								
iron(II), Fe ²⁺ (aq)	green ppt. turning brown on contact with air insoluble in excess	green ppt. turning brown on contact with air insoluble in excess								
iron(III), Fe ³⁺ (aq)	red-brown ppt. insoluble in excess	red-brown ppt. insoluble in excess								
magnesium, Mg²+(aq)	white ppt. insoluble in excess	white ppt. insoluble in excess								
manganese(II), Mn²+(aq)	off-white ppt. rapidly turning brown on contact with air insoluble in excess	off-white ppt. rapidly turning brown on contact with air insoluble in excess								
zinc, Zn²+(aq)	white ppt. soluble in excess	white ppt. soluble in excess								

2 Reactions of anions

ion	reaction
carbonate, CO ₃ ²⁻	CO ₂ liberated by dilute acids
chloride, C <i>l</i> ⁻ (aq)	gives white ppt. with Ag ⁺ (aq) (soluble in NH ₃ (aq))
bromide, Br ⁻ (aq)	gives cream ppt. with Ag ⁺ (aq) (partially soluble in NH ₃ (aq))
iodide, I^(aq)	gives yellow ppt. with Ag ⁺ (aq) (insoluble in NH ₃ (aq))
nitrate, NO ₃ ⁻ (aq)	NH ₃ liberated on heating with OH ⁻ (aq) and A <i>l</i> foil
nitrite, NO ₂ ⁻ (aq)	NH_3 liberated on heating with $OH^-(aq)$ and Al foil; NO liberated by dilute acids (colourless $NO \rightarrow$ (pale) brown NO_2 in air)
sulfate, SO ₄ ²⁻ (aq)	gives white ppt. with Ba ²⁺ (aq) (insoluble in excess dilute strong acids)
sulfite, SO ₃ ²⁻ (aq)	gives white ppt. with Ba ²⁺ (aq) (soluble in excess dilute strong acids)

3 Tests for gases

gas	test and test result				
ammonia, NH ₃	turns damp red litmus paper blue				
carbon dioxide, CO ₂	gives a white ppt. with limewater (ppt. dissolves with excess CO ₂)				
chlorine, Cl ₂	bleaches damp litmus paper				
hydrogen, H ₂	"pops" with a lighted splint				
oxygen, O ₂	relights a glowing splint				

The Periodic Table of Elements

			_		Π								- ~		_			_	
F 2	helium 4.0	10	Ne	neon 20.2	18	Ā	argon 39.9	36	궃	kryptor 83.8	42	Xe	xenon 131.3	98	R	radon			
		6	ш	fluorine 19.0	17	Cl	chlorine 35.5	35	B	bromine 79.9	53	Н	iodine 126.9	82	¥	astatine -			
		80	0	oxygen 16.0	16	S	sulfur 32.1	34	Se	selenium 79.0	52	<u>e</u>	tellurium 127.6	84	Ъ	polonium –	116	^	livermorium –
		7	z	nitrogen 14.0	15	۵	phosphorus 31.0	33	As	arsenic 74.9	51	Sb	antimony 121.8	83	Ξ	bismuth 209.0			
		9	ပ	carbon 12.0	14	S	silicon 28.1	32	Ge	germanium 72.6	20	Sn	tin 118.7	82	Pb	lead 207.2	114	Εl	flerovium -
		2	В	boron 10.8	13	Ν	aluminium 27.0	31	Ga	gallium 69.7	49	In	indium 114.8	18	11	thallium 204.4			
							12	30	Zu	zinc 65.4	48	g	cadmium 112.4	80	Нg	mercury 200.6	112	ပ်	copernicium
							7	29	Cn	copper 63.5	47	Ag	silver 107.9	79	Au	gold 197.0	111	Rg	roentgenium -
							10	28	Ż	nickel 58.7	46	Pd	palladium 106.4	78	చ	platinum 195.1	110	Ds	darmstadtium -
							6	27	රි	cobalt 58.9	45	뫈	rhodium 102.9	77	٦	iridium 192.2	109	₹	meitnerium -
- エ	hydrogen 1.0						80	26	Ъе	iron 55.8	44	Ru	ruthenium 101.1	9/	SO	osmium 190.2	108	Hs	hassium
		J					7	25	Mn	manganese 54.9	43	ည	technetium -	75	Re	rhenium 186.2	107	Bh	bohrium –
			loc	SS			9	24	ပ်	chromium 52.0	42	Mo	molybdenum 95.9	74	≥	tungsten 183.8	106	Sg	seaborgium -
	Key	stomic number	mic sym	name tive atomic ma			2	23	>	vanadium 50.9	41	g	niobium 92.9	73	д	tantalum 180.9	105	9	dubnium –
			ato	rela			4	22	F	titanium 47.9	40	Zr	zirconium 91.2	72	Ξ	hafnium 178.5	104	꿆	rutherfordium -
					_		က	21	Sc	scandium 45.0	39	>	yttrium 88.9	57-71	lanthanoids		89–103	actinoids	
		4	Be	beryllium 9.0	12	Mg	magnesium 24.3	20	Ca	calcium 40.1	38	တ်	strontium 87.6	56	Ba	barium 137.3	88	Ra	radium
		3	:=	ithium 6.9	1	Na	odium 23.0	19	¥	stassium 39.1	37	Rb	ubidium 85.5	55	Cs	aesium 132.9	87	ь Г	ancium –
		T hydrogen 1.0	Key hydrogen 1.0 5 6 7 8 9	Key hydrogen 1.0 5 6 7 8 9 Be atomic symbol B C N O F	Key 1.0 H Prodrogen atomic number 1.0 F F B T 8 9 F Be atomic symbol B C N O F beryllum relative atomic mass 9.0 relative atomic mass 10.8 12.0 14.0 16.0	Key hydrogen 1,0 F F F F B F B C N O F Bequired atomic name benching mass name atomic mass boron carbon ritrogen carbon ritrogen oxygen fluorine atomic mass number of thorine properties atomic mass number of thorine properties atomic mass 12.0 14.0 16.0 17.0 12 13 14 15 16 17	Key hydrogen 1.0 F	Key Invalvagen 1 modification 1 modification	Key hydrogen 1.0 Telegy monitor number 1.0 Fe production of the produc	H	The control of the	The color of the	The color The	A	H	Harmonidar Har	H	The control of the	The control of the

71	P	lutetium 175.0	103	۲	lawrencium	ı
20	ΥÞ	ytterbium 173.1	102	Š	nobelium	ı
69	E	thulium 168.9	101	Md	mendelevium	ı
89	ш	erbium 167.3	100	Fn	ferminm	1
29	웃	holmium 164.9	66	Es	einsteinium	-
99	٥	dysprosium 162.5	86	ర	californium	ı
65	Д	terbium 158.9	26	Ř	berkelium	ı
29	gg	gadolinium 157.3	96	CH	curium	ı
63	En	europium 152.0	92	Am	americium	ı
62	Sm	samarium 150.4	94	Pn	plutonium	I
61	Pm	promethium	93	ď	neptunium	ı
09	PZ	neodymium 144.4	92	\supset	uranium	238.0
59	፵	praseodymium 140.9	91	Ра	protactinium	231.0
58	Ce	cerium 140.1	06	드	thorium	232.0
57	Га	lanthanum 138.9	89	Ac	actinium	ı

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lanthanoids

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