

Surname	Centre Number	Candidate Number
First name(s)		2



GCE A LEVEL

A410U30-1



FRIDAY, 21 JUNE 2024 – MORNING

CHEMISTRY – A level component 3

Chemistry in Practice

1 hour 15 minutes

For Examiner's use only		
Question	Maximum Mark	Mark Awarded
1.	8	
2.	13	
3.	8	
4.	19	
5.	12	
Total	60	

ADDITIONAL MATERIALS

- A calculator, pencil and ruler
- **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.

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.

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 60.

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

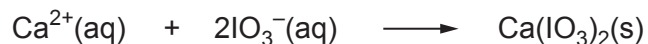
The assessment of the quality of extended response (QER) will take place in **Q2(a)(i)**.



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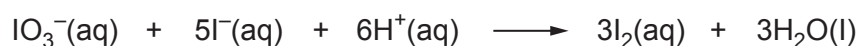
Answer **all** questions.

1. Calcium iodate(V) is a sparingly soluble salt and can be prepared by the reaction between solutions of potassium iodate(V) and a suitable soluble calcium salt, such as calcium chloride.



It is usually obtained as a hydrate, $\text{Ca}(\text{IO}_3)_2 \cdot x\text{H}_2\text{O}$, when precipitated from aqueous solution.

In acid solution, iodate(V) ions react with excess iodide ions to liberate iodine.



The iodine formed can be determined by titration against standard sodium thiosulfate solution.



Using this technique, the relative formula mass of the salt, $\text{Ca}(\text{IO}_3)_2 \cdot x\text{H}_2\text{O}$, and the value of x were determined as follows.



Step	Method
1	An accurately weighed sample of solid $\text{Ca}(\text{IO}_3)_2 \cdot x\text{H}_2\text{O}$ was dissolved in hot water in a conical flask.
2	After cooling, excess acidified potassium iodide solution was added to the solution in the flask.
3	The solution containing the iodine (I_2) formed was titrated against $0.540 \text{ mol dm}^{-3}$ sodium thiosulfate solution until the dark brown colour changed to pale yellow. Starch indicator was added and the titration continued until the end-point. The volume of sodium thiosulfate solution used was recorded.
4	The method was repeated with a further two samples of the solid $\text{Ca}(\text{IO}_3)_2 \cdot x\text{H}_2\text{O}$.

Results

Titration number	1	2	3
Mass of $\text{Ca}(\text{IO}_3)_2 \cdot x\text{H}_2\text{O}$ / g	0.481	0.493	0.474
Volume of $\text{S}_2\text{O}_3^{2-}(\text{aq})$ / cm^3	21.45	22.00	21.15

- (a) For step 3, give the colour change of the starch indicator at the end-point. [1]

.....

- (b) Identify the titration that has the smallest percentage error in the **mass** of $\text{Ca}(\text{IO}_3)_2 \cdot x\text{H}_2\text{O}$ used. Give your reasoning. [1]

Titration number

.....

.....



(c) The five steps of the calculation are shown below.

- (i) Number these in the correct order. The first and last steps have been done for you. [1]

	Correct order
Convert moles of IO_3^- to moles of $\text{Ca}(\text{IO}_3)_2 \cdot x\text{H}_2\text{O}$	
Use the balanced equation(s) to calculate the number of moles of IO_3^- ions present in the solid sample	
Calculate the value of x in $\text{Ca}(\text{IO}_3)_2 \cdot x\text{H}_2\text{O}$	5
Convert moles of $\text{Ca}(\text{IO}_3)_2 \cdot x\text{H}_2\text{O}$ to M_r of $\text{Ca}(\text{IO}_3)_2 \cdot x\text{H}_2\text{O}$	
Calculate the number of moles of sodium thiosulfate used in the titration	1

- (ii) Carry out the calculation to determine the relative formula mass of the salt, $\text{Ca}(\text{IO}_3)_2 \cdot x\text{H}_2\text{O}$, and hence the value of x . Use the results from **titration 2** only.

You **must** show your working. [5]

$x =$



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2. (a) Peroxydisulfate ions react with iodide ions to form iodine.



The rate equation for the reaction is as follows.

$$\text{rate} = k [\text{S}_2\text{O}_8^{2-}] [\text{I}^-]$$

One way of measuring the rate is to time how long it takes to produce a certain amount of iodine using the iodine clock reaction.

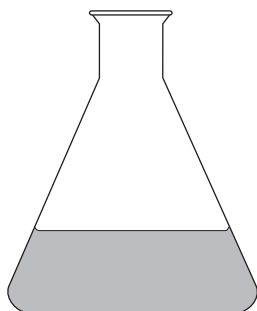
A student was asked to carry out an experiment to confirm that the reaction is first order with respect to peroxydisulfate ions.

The student was provided with the following aqueous solutions and deionised water.

0.1 mol dm⁻³ potassium peroxydisulfate
 0.3 mol dm⁻³ potassium iodide
 0.02 mol dm⁻³ sodium thiosulfate
 1% starch solution

Assume that all solutions are at the same temperature.

The student started the experiment by accurately measuring the solutions into a conical flask.



25.0 cm³ of potassium iodide
 10.0 cm³ of sodium thiosulfate
 1.0 cm³ of starch solution
 4.0 cm³ of deionised water

They then added 10.0 cm³ of 0.1 mol dm⁻³ potassium peroxydisulfate solution.

They recorded the time taken for the expected colour change to be seen.

The student needed to carry out one more run to complete the experiment.



Give the practical details, including reference to the apparatus used. State how the data would confirm that the reaction is first order with respect to peroxydisulfate ions.



- (ii) The peroxydisulfate-iodide reaction can be catalysed by some *d*-block metal ions.

Outline how the method described in part (i) could be **modified** to investigate the catalytic effect of $\text{Fe}^{2+}(\text{aq})$ on the reaction. State how the results would show that $\text{Fe}^{2+}(\text{aq})$ acts as a catalyst. [2]

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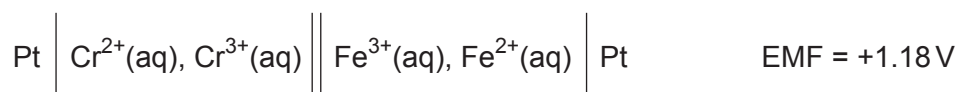
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- (b) Consider the cell diagram below where the negative electrode is shown on the left-hand side.



- (i) Give the half-equation for the reaction occurring at the negative electrode. [1]

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- (ii) Give the observation at the positive electrode after the cell has been connected for a period of time. [1]

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- (iii) Give the overall equation for the cell reaction. Identify the oxidising agent and the reducing agent. [2]

Equation

.....

Oxidising agent

Reducing agent

- (iv) The concentrations of the $\text{Cr}^{2+}/\text{Cr}^{3+}$ and $\text{Fe}^{3+}/\text{Fe}^{2+}$ ions are all 1.0 mol dm^{-3} .

State and explain how the value on a high-resistance voltmeter would change if the concentration of the Cr^{3+} ions was decreased whilst the concentrations of all the other ions were left unchanged. [1]

Change in EMF

Explanation

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3. Organic acids derived from fats are described as weak acids.

(a) Explain the terms **weak** and **dilute** as applied to acid solutions. [2]

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(b) A student is given samples of nitric acid and ethanoic acid, both of concentration 1.0 mol dm^{-3} .

Describe **one chemical** test, apart from the use of an indicator, that could be used to distinguish between both acids.

Your answer should include an explanation of the different observations made. [2]

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- (c) A laboratory technician prepared a buffer solution of pH 5.4 by mixing 1.0 mol dm^{-3} aqueous ethanoic acid and 1.0 mol dm^{-3} aqueous sodium ethanoate.

- (i) Calculate the ratio of the concentration of sodium ethanoate to ethanoic acid in the buffer mixture. [3]

(K_a for ethanoic acid = $1.78 \times 10^{-5} \text{ mol dm}^{-3}$ at 298 K)

Ratio of concentration of sodium ethanoate to ethanoic acid :

- (ii) Calculate the volumes of sodium ethanoate and ethanoic acid used to prepare 1 dm^3 of the buffer solution. [1]

Volume of ethanoic acid = cm^3

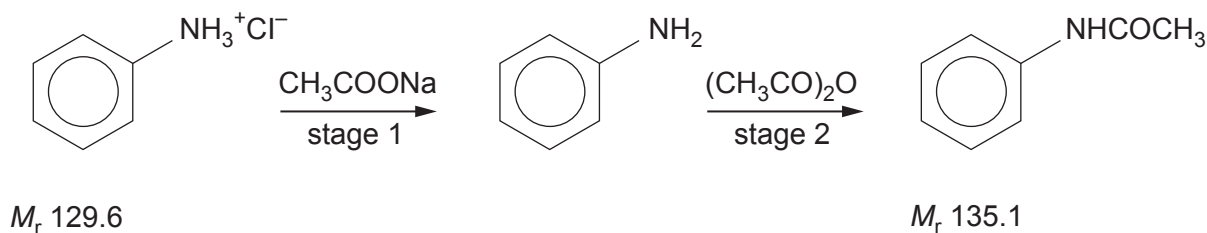
Volume of sodium ethanoate = cm^3



4. Acylation is the term used to describe the substitution of a hydrogen atom in a phenol, alcohol or amine by the acyl group, RCO— .

Ethanoic anhydride, $(\text{CH}_3\text{CO})_2\text{O}$, can be used as an acylating agent.

N-phenylethanamide, $\text{C}_6\text{H}_5\text{NHCOCH}_3$, can be prepared from the salt phenylammonium chloride, $\text{C}_6\text{H}_5\text{NH}_3^+\text{Cl}^-$. The reaction takes place in two stages.



The method below was used to prepare a pure sample of N-phenylethanamide.

- 4.78 g of phenylammonium chloride was dissolved in 150 cm^3 of deionised water in a conical flask.
- 10.0 cm^3 of ethanoic anhydride was added to the solution in the flask. The solution was stirred to ensure that all of the ethanoic anhydride had dissolved.
- 30.0 g of hydrated sodium ethanoate was dissolved in 100 cm^3 of deionised water and added to the flask. The mixture was stirred for 5 minutes.

The sodium ethanoate reacted with the phenylammonium chloride in stage 1, forming phenylamine, ethanoic acid and sodium chloride. The reaction in stage 2 happened readily with N-phenylethanamide forming as a white solid. Ethanoic acid was also formed.

The N-phenylethanamide was recrystallised to give 3.59 g of the pure product.

- (a) Write the equation for the reaction occurring in stage 1.

[1]



- (b) N-phenylethanamide is much more soluble in hot water than it is in cold water.

Use this fact to outline how you would purify N-phenylethanamide to obtain a pure, dry product. [3]

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- (c) 4.78 g of phenylammonium chloride gave 3.59 g of pure N-phenylethanamide.

- (i) Show that an excess of ethanoic anhydride was present in the reaction mixture. [2]

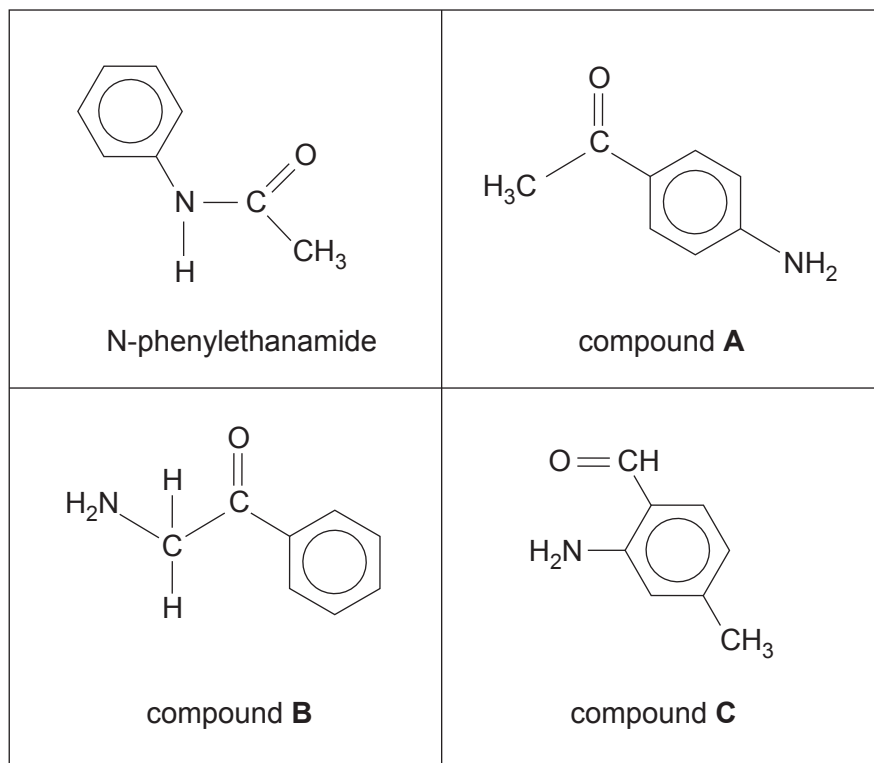
density of ethanoic anhydride = 1.08 g cm^{-3}

- (ii) Calculate the overall percentage yield of this two-stage reaction. [2]

Percentage yield = %



- (d) N-phenylethanamide, C_8H_9NO , and compounds **A**, **B** and **C** are structural isomers. They were investigated using test tube reactions and NMR spectroscopy.



- (i) Describe how the low resolution 1H NMR spectrum of compound **C** would differ from those of the other three compounds.

Assume that all aromatic protons are equivalent. You do not need to refer to the chemical shift values or peak areas. [2]

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- (ii) State a chemical test that will give a positive result for compound **C** but not for compound **A**. Give the reagent(s) and observation. [1]

Reagent(s)

Observation



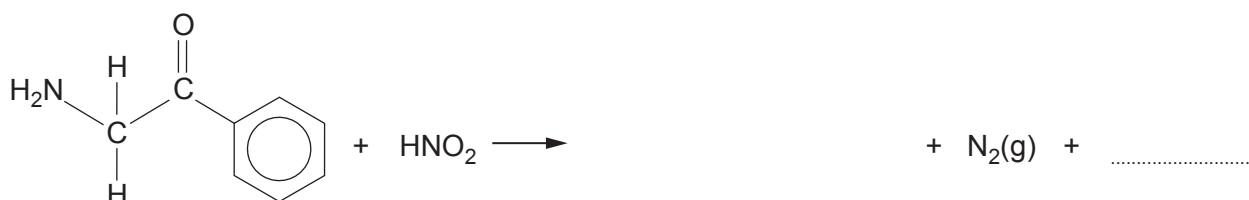
- (iii) State a chemical test that will give a positive result for N-phenylethanamide and compound **A**, but not for compounds **B** and **C**. Give the reagent(s) and observation. [1]

Reagent(s)

Observation

- (iv) Nitric(III) acid (nitrous acid), HNO_2 , is an unstable compound and is made when required by the reaction of a dilute acid, for example HCl , with sodium nitrate(III), NaNO_2 .

- I. Complete the equation for the reaction of compound **B** with nitric(III) acid. Give the structure of the organic compound formed. [1]



- II. In an experiment, 3.59 g of compound **B** (M_r 135.1) was dissolved in water and the solution made up to 500 cm^3 . A 25.0 cm^3 sample of this solution was reacted with an excess of nitric(III) acid.

Calculate the volume, in cm^3 , of nitrogen gas formed at 20°C and 1 atm. [4]

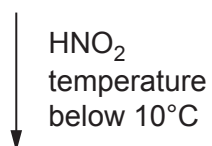
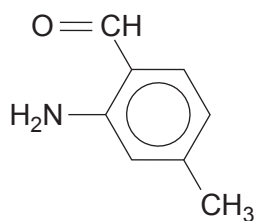
Volume = cm^3



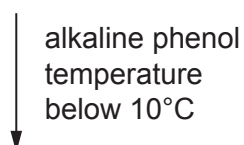
- (v) Compound **D** can be prepared from compound **C** in a two-stage process.

Give the structures of the intermediate and compound **D**.

[2]

compound **C**

intermediate

compound **D**

- The following tests were carried out on a Tutton salt where A^{a+} is a Group 1 cation and B^{b+} is a transition metal cation. A solution of the salt was prepared for use in tests 2 and 3.

[12]

Test	Result	Conclusion
Test 2A		
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.....		Cr^{3+} not present so $\text{B}^{\text{b}+}$ is Fe^{2+}
.....		
.....		

Test 2B		
Leave the precipitate that forms exposed to the air for 24 hours	Precipitate turns reddish-brown	



Test	Result	Conclusion
Test 3A		
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.....		
.....		
.....		
Equation		
.....		
Test 3B		
Add excess reagent	50.0 cm ³ sample of a 0.736 mol dm ⁻³ solution of the salt produces a precipitate with a dry mass of 17.16 g	
		y =



12



[illegible]