Write your name here	Other na	mes
Samuric	o the rid	mes
Pearson Edexcel Level 3 GCE	Centre Number	Candidate Number
Chemistry Advanced Paper 3: General and		les in Chemistry
Sample Assessment Materials for first Time: 2 hours 30 minutes		Paper Reference 9CH0/03
You must have: Data Booklet Scientific calculator, ruler		Total Marks

Instructions

- Use **black** ink or ball-point pen.
- **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.
- You may use a scientific calculator.
- For questions marked with an *, 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.
- Try to answer every question.
- Check your answers if you have time at the end.
- Show all your working in calculations and include units where appropriate.

Turn over ▶

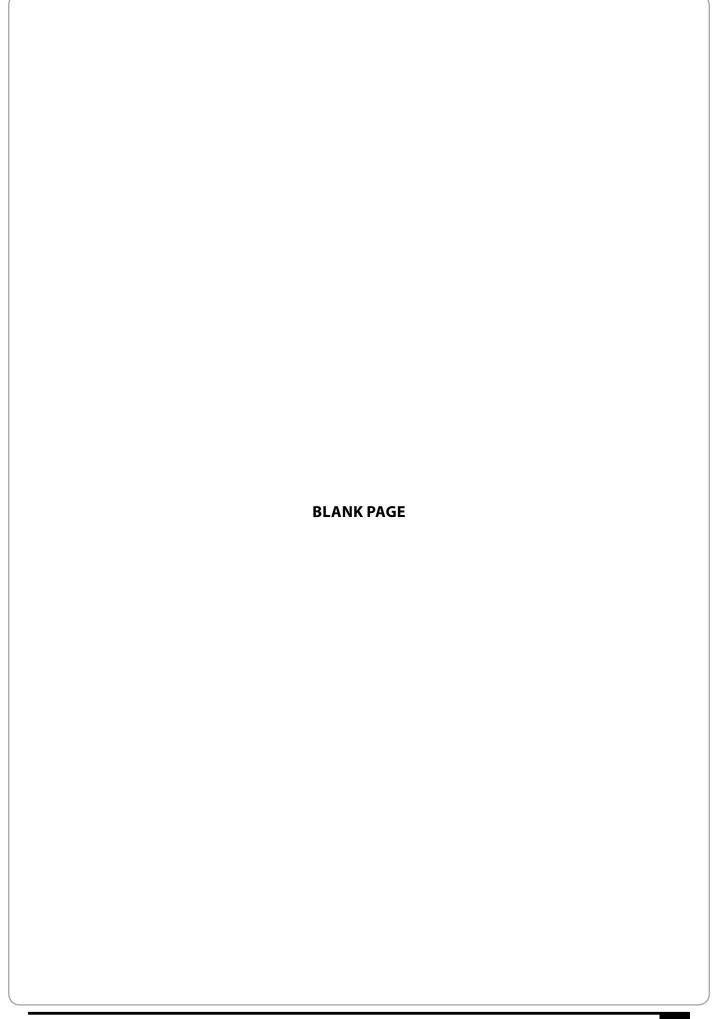
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Answer ALL questions.

Write your answers in the spaces provided.	
Solid benøic acid is usually purified by recrystallisation from hot water. The method used is:	
1 The impure solid is dissolved in the minimum volume of hot water.	
2 The solution is quickly filtered.	
3 The solution is allowed to cool and crystallise.	
4 The crystals are filtered from the remaining solution.	
5 The crystals are washed with a little cold water.	
6 The crystals are left to dry.	
(a) State how steps 2 , 4 and 5 remove impurities from the benøic acid.	β)
b) Describe how impurities affect the melting temperature of an impure solid.	
	2)
 Tatalfan Osa tian 1 Fu	
▼ otal for Q es tion 1 = 5 r	narks



2	Compound ${\bf A}$ is a green solid containing one cation and one anion. ${\bf t}$ dissolves in water to form a green solution.	
	(a) © e the formulae of two cations which could be responsible for the green colour in the solid.	
		2)
•••••	b) A student adds dilute sodium hydroide solution, drop by drop, to an aqueous	
	solution of A . A green precipitate forms, which dissolves in exe ss sodium hydroide solution to form a dark green solution.	
	() & e the formula of the cation in A .	1
		R*
	(i) 🕼 e a formula for the green precipitate.	1
		,
	(ii) 🛱 e a formula for the species present in the dark green solution.)
		J

()	The student adds a few drops of acidified potassium manganately solution to another sample of a solution of A in a test tube.	
	State the colour change that occurs.	(1)
(d)	The student acidifies 2 cm ³ of a solution of A with dilute nitric acid in a test tube	
	and then adds a few drops of aqueous silver nitrate. A white precipitate is formed. () () () () () () () () () ()	(1)
	(i) This test is usually followed by the addition of ammonia solution to test the solubility of the precipitate.	
	Exp lain why this procedure in not suitable to confirm the identity of the anion in ${\bf A}$.	(2)
	(Total for Question 2 = 9 ma	rks)

3 The equation for the reaction between iodine and propanone is:

$$\mathsf{CH}_{3}\mathsf{COCH}_{3}\mathsf{\acute{e}q}\;)\;+\;\mathsf{I}_{2}\mathsf{\acute{e}q}\;)\;\rightarrow\;\mathsf{CH}_{3}\mathsf{COCH}_{2}\mathsf{\acute{e}q}\;)\;+\;\mathsf{H}^{+}\!\!\left(\mathsf{\acute{e}q}\;\right)\;+\;\mathsf{I}^{-}\!\!\left(\mathsf{\acute{e}q}\;\right)$$

The kinetics of this reaction were studied.

(a) The order of the reaction with respect to iodine was determined by preparing a mixture of solutions of iodine and sulfuric acid in a conical flask. A solution of propanone was then added and a timer started. After one minute, a 10.0 cm³ sample of the reaction mixture was removed. Further 10.0 cm³ samples of the reaction mixture were removed at regular time intervals during the experiment.

After removal, each sample was immediately added to sodium hydrogencarbonate solution and then titrated with sodium thiosulfate solution to determine the concentration of iodine.

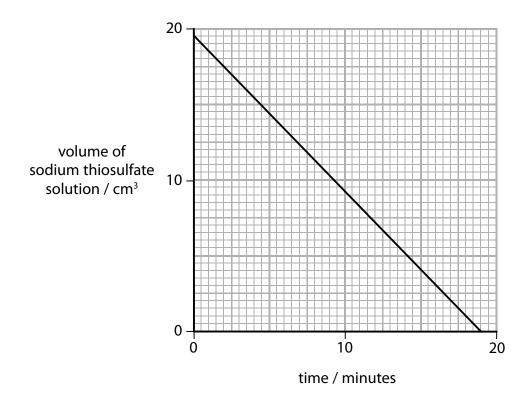
The table shows the volumes and initial concentrations of the substances in one experiment.

&b stance	Volume / cm³	Concentration / mol dm ⁻³
iodine	50	0.020
sulfuric acid	20	2.5
propanone	25	2.0

() Deduce, by calculation of the amounts used, whether propanone or iodine was in exe ss.

(I)	Name the piece of apparatus which is most suitable for measuring the volume of sulfuric acid.	(1)
(ii)	Name the piece of apparatus which is most suitable for removing the sample	
(11)	from the reaction mixure .	(1)

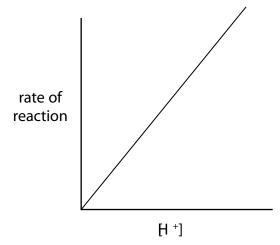
 $\ensuremath{\mathfrak{b}}$) The graph shows the results of titrating the samples of the reaction mixture $% {\ensuremath{\mathfrak{b}}}$.



Ep lain how you can deduce the order of reaction with respect to iodine from this graph.

() These graphs were obtained in other exp eriments to determine the order of reaction with respect to propanone and to sulfuric acid.

rate of reaction [CH₃COCH₃]

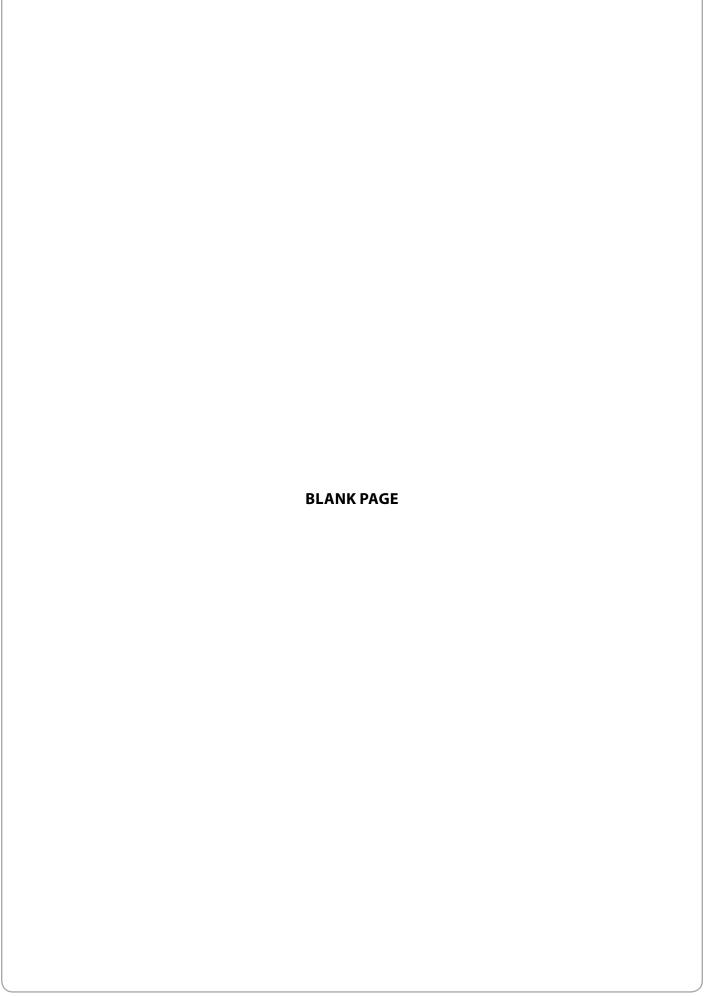


Use these graphs and your answer from part ${\mathfrak b}$) to comment on whether this proposed mechanism for the reaction is correct.

(3)

(Total for Question 3 = 9 marks)

slow



- 4 This question is about the effect of changes in temperature on reactions.
 - (a) An exp eriment to determine the activation energy for the reaction between magnesium and hydrochloric acid was carried out. The time taken for 0.100 g of magnesium to react completely when added to 20.0 cm³ of 1.00 mol dm⁻³ hydrochloric acid was recorded at different temperatures. A summary of the processed data is shown.

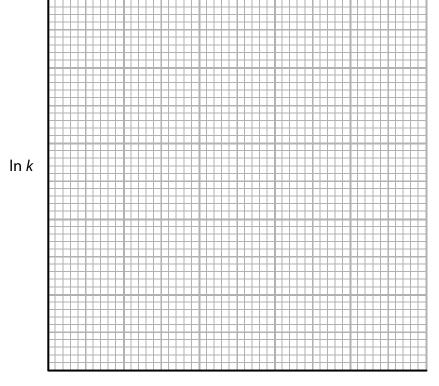
Temperature / K	1/K ⁻¹	ln <i>k</i>
283	3.53×10^{-3}	-4.80
299	3.34×10^{-3}	-3.56
311	3.22×10^{-3}	-3.00
322	3.11×10^{-3}	-2.25
329	3.04×10^{-3}	-1.79

The activation energy, E_{a} , of the reaction can be found using the equation:

$$\ln k = -\frac{E_a}{RT} + c$$

Use this data to plot a graph of $\ln k$ against $\frac{1}{T}$ and hence determine the activation energy in kJ mol⁻¹.

(4)



$$\frac{1}{T} / K^{-1}$$

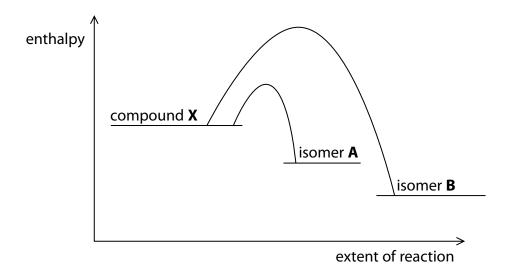
Activation energy =kJ mol $^{-1}$

† b) h the study of a different reaction, a compound **X** reacted reversibly with concentrated sulfuric acid to form two isomers, **A** and **B**.

compound
$$\mathbf{X} + \mathbf{H}_2 SO_4 \mathbf{\hat{c}}$$
 onc.) \rightleftharpoons isomer \mathbf{A} + isomer \mathbf{B}

At 40 °C, approximate ly 95% of the product was isomer **A**. At 160 °C, the product contained approximate ly 85% of isomer **B**.

The diagram shows the reaction profiles for the formation of the two isomers.



Use the information to comment on the different yields of isomer **A** and isomer **B** at different temperatures.

(Total for Question 4 = 0 marks)

(6)

5 Ammonia is produced industrially by reacting nitrogen and hydrogen.

$$N_2 g$$
) + $3H_2 g$) \Rightarrow 2NH₃g) $\Delta H^{\oplus} = -92.0 \text{ kJ mol}^{-1}$

A temperature in the range of 673 to 773 K is used.

The standard entropies, S^{\ominus} , of $N_2 G$) $H_2 G$) and $NH_3 G$) at 298 K are given in the table.

substance	N ₂ g)	H ₂ g)	NH ₃ g)
S [⊕] / J K ⁻¹ mol ⁻¹	192	131	193

(a) Show that this reaction is feasible at 298 K by calculating ΔG^{\ominus} in kJ mol⁻¹. Go e your answer to an appropriate number of significant figures.

(5)

b)	Εpο	lain, in terms	of entropy,	why this	reaction is	not feasible	e at very	high
	ten	nperatures.						

(c) Nitrogen and hydrogen were reacted together at 673 K and 200 atm pressure in a closed vessel.

$$N_2 g) + 3 H_2 g) \rightleftharpoons 2 N H_3 g)$$

The reaction mixture was allowed to reach equilibrium. The number of moles of each gas at equilibrium were found to be:

$$N_2 g$$
) 2.88 $H_2 g$) 8.64 $NH_3 g$) 6.48

Calculate the value of $K_{\rm p}$ at 673 K, giving appropriate units.

(5)

(Total for Question 5 = 2 marks)

6	Thi	is question is about the chemistry of the alcohol functional group.	
	(a)	Ethanol and water mix together readily because of the formation of hydrogen bonds between the molecules.	
		Devise an experiment to estimate the strength of this interaction including how you would process the data collected.	
			(6)

b) The painkiller aspirin can be synthesised by the reaction between 2-hydrox benzoic acid, which contains a hydrox group, and ethanoic anhydride, using concentrated phosphoric acid as a catalyst. The reagents are heated under reflux then the exe ss ethanoic anhydride is removed by reacting it with water.

$$CO_2H$$
 CO_2H CO_2H CO_2C CO_2C CO_3C CO_3

1) The percentage yield for this synthesis is 65%.

Calculate the mass of aspirin you would obtain using 2.0 g of 2-hydroxybenzoic acid.

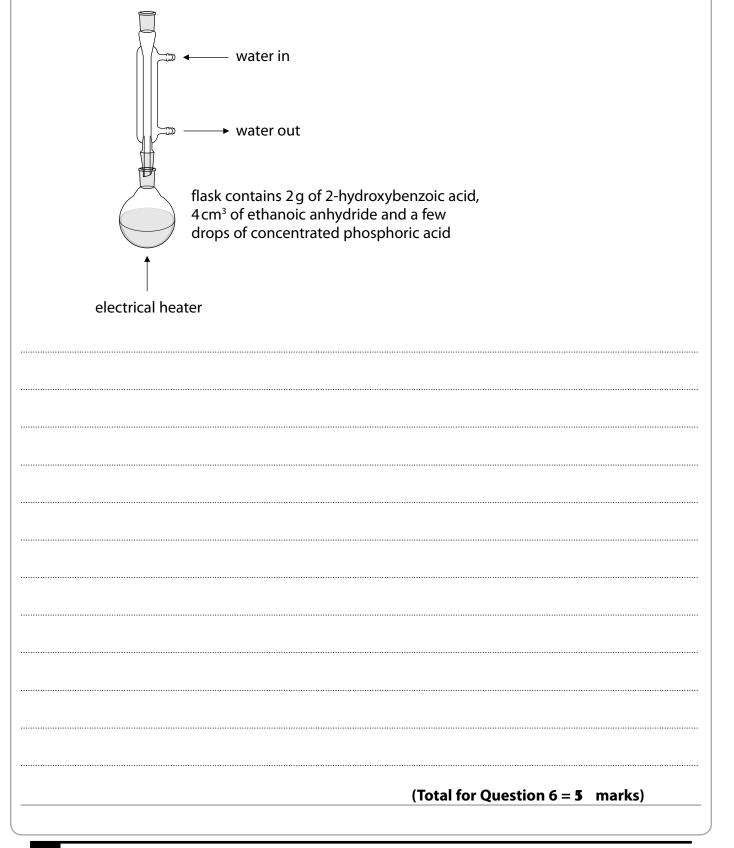
(3)

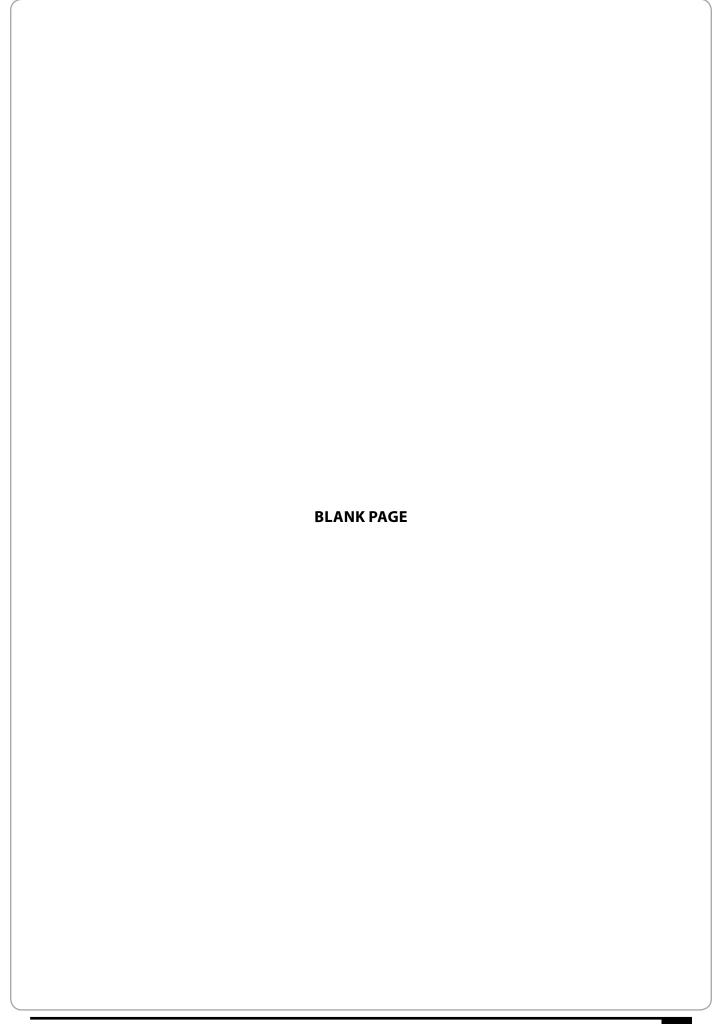
(i) The diagram shows a proposed set-up of apparatus used for the stage of the synthesis that requires heating under reflux

de ntify **three** improvements that should be made to this set-up. **©** e a reason for each improvement made.

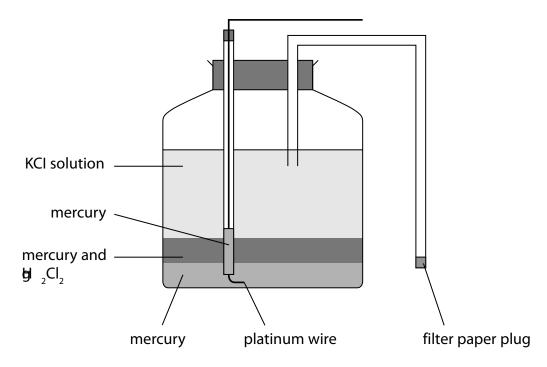
You may assume suitable clamps are used.

(6)





7 The diagram shows a calomel electrode, which can be used for measuring standard electrode potentials.



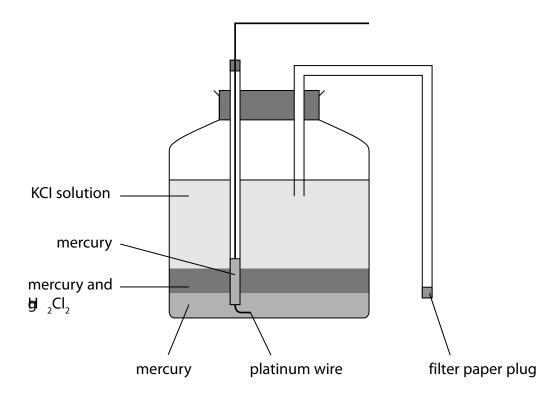
A)	n	State a solution that could be used in the side arm to act as the salt bridge.
ų,	v	State a solution that could be used in the side and to det as the sait bridge.

(1)

(i)	Fın	lain the	purpose	of the	salt	hridae
U)	ᅜ	ומווו נווכ	puipose	OI LITE	sait	Diluye.



 $\mathfrak b$) A calomel electrode can be used to measure the standard electromotive force $\mathfrak e$ mf) $E_{\rm cell}^{\ominus}$ of the cell:



() Complete the diagram above by drawing a suitable silver half-cell including the appropriate electric circuit.

(3)

(i) State the concentration of the solution in the silver half-cell and a suitable experimental condition.

the discussion of the batta bookies to calculate the standard enil of the central by	Use the Da	calculate the standard emf of the cell in bi)	given
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(i) The emf of the cell shown below is:

Pt
2
 6 1 1 1 1 2 1 2 2 1 2

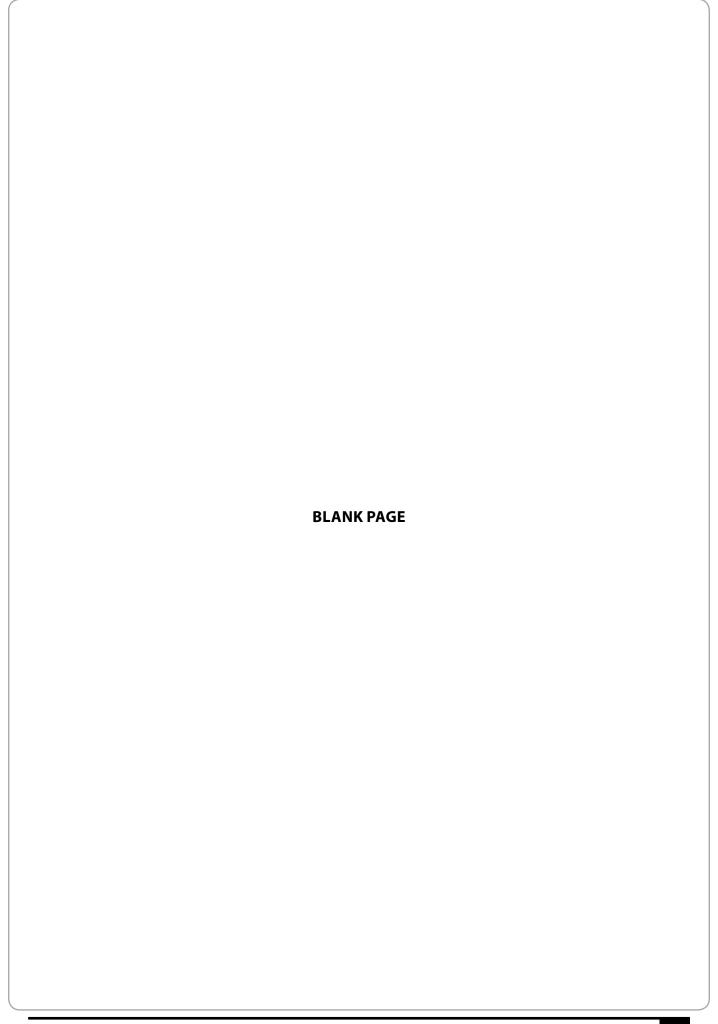
Use this value to calculate the standard emf of the following cell:

PtFe
$$^{2+}$$
aq) Fe $^{3+}$ aq] :: Ag+aq Ag \$) (1)

(ii) Write the overall equation for the cell reaction when current is being drawn.

(v) Calculate the numerical value of the equilibrium constant for this reaction at 298 K using $\Delta G^{\oplus} = -2892$ J mol⁻¹ and the relationship $\Delta G^{\oplus} = -RT \ln K$ (2)

(Total for Question 7 = 4 marks)



8 Cinnamon is a spice that is extracted from the bark of the cinnamon tree.

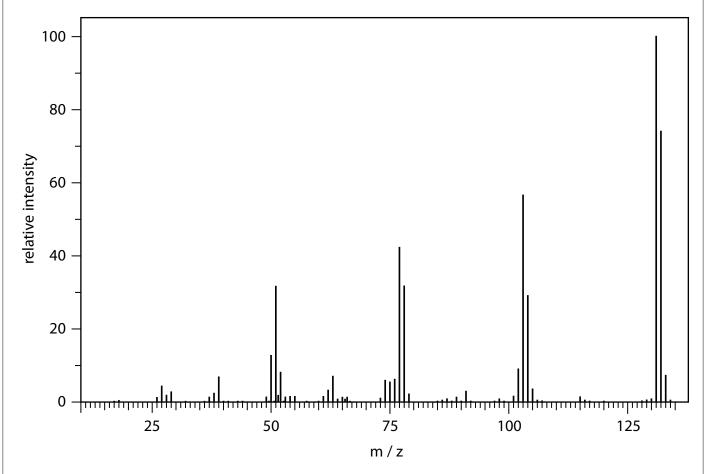
It is extracted by crushing the soft bark with water, and then steam distilling the mixture to produce cinnamon oil.

- (a) The steam distillate is washed with saturated sodium chloride solution and the oil separated and dried.
 - () Draw a diagram of the apparatus you would use to separate the oil, labelling the oil.

D ensities: cinnamon oil 1.050 g cm⁻³; sodium chloride solution 1.122 g cm⁻³]

(ii) State the change in appearance of the oil when it is being dried. (1) (1) (1) (1) (1) (1) (1) (1	(1) b) Compound Q is the main component of cinnamon oil. Some chemical tests are carried out to try to find the structure of Q . The results of each test are given in parts () to (ii) Deduce from the results of each test what functional group may be present in Q . () Q decolourises both bromine water and acidified potassium manganate(V (1) (i) Q gives a yellow precipitate with 2,4-dinitrophenylhydraine. (1)	(1) b) Compound Q is the main component of cinnamon oil. Some chemical tests are carried out to try to find the structure of Q . The results of each test are given in parts () to (ii) Deduce from the results of each test what functional group may be present in Q . () Q decolourises both bromine water and acidified potassium manganate(V) (1) (i) Q gives a yellow precipitate with 2,4-dinitrophenylhydraine. (1)	(i) (i) (i) e the name of a chemical that could be used to dry the oil.	(1)
carried out to try to find the structure of Q . The results of each test are given in parts () to (ii) Deduce from the results of each test what functional group may be present in Q . () Q decolourises both bromine water and acidified potassium manganate(V) (i) Q gives a yellow precipitate with 2,4-dinitrophenylhydraine . (1) (ii) Q forms a red precipitate when boiled with either Benedict's or Fehling's solution.	carried out to try to find the structure of Q . The results of each test are given in parts () to (ii) Deduce from the results of each test what functional group may be present in Q . () Q decolourises both bromine water and acidified potassium manganate(V) (i) Q gives a yellow precipitate with 2,4-dinitrophenylhydraine . (1) (ii) Q forms a red precipitate when boiled with either Benedict's or Fehling's solution.	carried out to try to find the structure of Q . The results of each test are given in parts () to (ii) Deduce from the results of each test what functional group may be present in Q . () Q decolourises both bromine water and acidified potassium manganate(V) (1) (i) Q gives a yellow precipitate with 2,4-dinitrophenylhydraine . (1)	(ii) State the change in appearance of the oil when it is being dried.	(1)
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(i) Q decolourises both bromine water and acidified potassium manganate(V (1) (1) (1) (1) (1) (1) (1) (1) (1) (1)	(i) Q decolourises both bromine water and acidified potassium manganate(V (1) (1) (1) (2) gives a yellow precipitate with 2,4-dinitrophenylhydraine . (1) (1)	(i) Q decolourises both bromine water and acidified potassium manganate(y (i) Q gives a yellow precipitate with 2,4-dinitrophenylhydraine . (ii) Q forms a red precipitate when boiled with either Benedict's or Fehling's solution.		
(i) Q gives a yellow precipitate with 2,4-dinitrophenylhydraine . (1) (ii) Q forms a red precipitate when boiled with either Benedict's or Fehling's solution.	(i) Q gives a yellow precipitate with 2,4-dinitrophenylhydraine . (1) (ii) Q forms a red precipitate when boiled with either Benedict's or Fehling's solution.	(i) Q gives a yellow precipitate with 2,4-dinitrophenylhydraine . (1) (ii) Q forms a red precipitate when boiled with either Benedict's or Fehling's solution.		! .
(ii) Q forms a red precipitate when boiled with either Benedict's or Fehling's solution.	(ii) Q forms a red precipitate when boiled with either Benedict's or Fehling's solution.	(ii) Q forms a red precipitate when boiled with either Benedict's or Fehling's solution.	() Q decolourises both bromine water and acidilled potassium manganate()	(1)
				(1)

 ξ) **Q** has the empirical formula C_9H_8O and its mass spectrum is shown.



Predict the identity of the fragment ions causing the peaks at 77 and 103.(2)

m/z	formula of fragment ion
77	
103	

(i) Deduce **two** possible displayed formulae for **Q**.

	 (ii) The peak at m/z = 132 in the mass spectrum of Q is the molecular ion peak. The mass spectrum also shows a peak at m/z = 133. G e a reason why the peak at m/z = 133 occurs. 	(1)
(b)	Compound Q can be converted into cinnamic acid which contains a carboxylic acid functional group and is a monobasic acid. 1.78 g of cinnamic acid is reacted with 250 cm³ of 0.500 mol dm⁻³ NaOH 25.0 cm³ of the resulting solution was titrated with 0.400 mol dm⁻³ B I. 28.25 cm³ was needed for complete neutralisation. Calculate the <i>M</i> _r of cinnamic acid, giving your answer to one decimal place.	(5)
	(Total for Question 8 = 7 m	arks)

9	Brass i	s an alloy of copper and inc .	
	(a) ()	Epp lain why copper is classified as a transition element but inc is not.	(2)
			(2)
	(i)	Brass has a structure similar to that of metallic copper, but with inc ions replacing some copper ions in the lattice.	
		Ep lain why brass is malleable whereas a crystal of sodium chloride is not.	(0)
			(3)

(b)	A sample of brass was weighed on a balar	ce, reading to	two deci	mal places.	The mass
	of the sample was recorded as 5.00 g.				

This sample of brass was reacted with excess concentrated nitric acid and the resulting solution was made up to 250 cm³ in a volumetric flask using distilled water.

25.0 cm³ portions were taken from this solution using a pipette. Each portion was neutralised by adding sodium carbonate solution; and excess potassium iodide solution was then added. The liberated iodine was titrated with 0.250 mol dm⁻³ sodium thiosulfate solution, using a freshly prepared solution of starch as indicator.

The mean titre was 22.70 cm³.

() Use the equation for the reaction to give two observations when nitric acid reacts with copper.

Cu\$) +
$$2NO_3^-$$
(aq) + $4H^+$ (aq) $\rightarrow Cu^{2+}$ (aq) + $2NO_2$ (g) + $2H_2O$ (1)

(2)

(i)	Complete the equation for the reaction between iodine and thiosulfate ions
	hclude state symbols.

(1)

$$l_2 (q) + 2S_2O_3^{2} (q) \rightarrow$$

(ii) Copper(ions react with iodide ions to form iodine.

$$2Cu^{2+}\!\!\left(\!\!\!\left(q\right)\right) \,+\, 4l^{-}\!\!\left(\!\!\!\left(q\right)\right) \,\rightarrow\, 2Cu\xi) \quad +\, l_{_{2}}\!\!\left(\!\!\!\left(q\right)\right)$$

(5)

	ent wants to identify the pie rement uncertainties in this ed.			
	ercentage measurement unc on the volumetric flask as ±0.0		the pipette as $\pm 0.24\%$),
By usir most s	ng appropriate calculations f significant source of measure	or the other apparatusement uncertainty in t	s used, deduce the his procedure.	(2)
		(Total fo	r Question 9 = 5 ma	nrks)

10

Benzene

Benzene was first discovered by Michael Faraday in London in 1825. El identified it by distilling a liquid obtained from condensing a gas produced by heating whale oil.

In 1834, Mitscherlich discovered the same liquid could be obtained by heating benzoic acid with lime.

Benzene is now produced from petroleum. Petroleum is fractionally distilled and benzene is obtained from the heane in the naphtha fraction.

El ane is heated to about 770 K. It then passes to a reactor where it reacts to form cycloheane and hydrogen. The cycloheane is then dehydrogenated to form benzene. Other aromatic products like methylbenzene and dimethylbenzenes are also produced. The aromatic products are separated by further distillation.

Benzene is the starting compound for a large number of useful chemicals and materials. For eam ple it is used to make the polymer, poly() henylethene)

(a) Write an equation for the reaction between benzoic acid and lime, calcium oide . State symbols are not required.

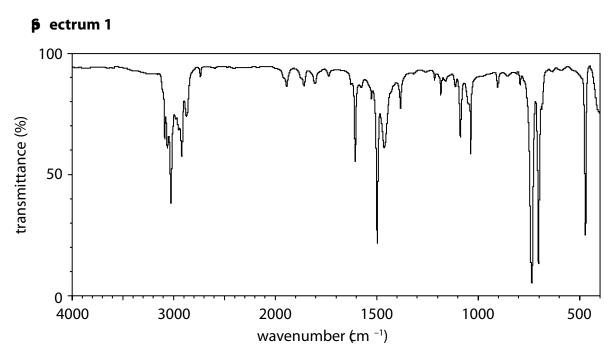
(1)

(b)	Deduce, with references to oida	tion numbers, what has happened to the carbo)n
	in the conversion of cyclohexne	to benzene.	

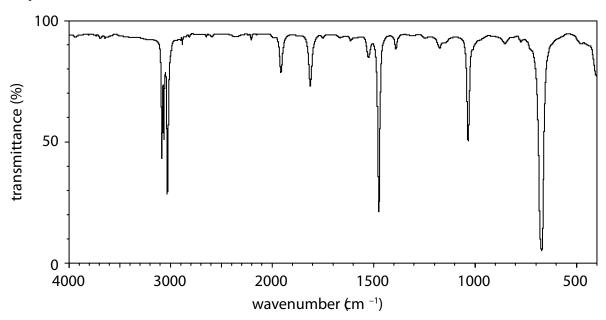
(c) The infrared spectra of benzene and methylbenzene are shown.

ble ntify which infrared spectrum is that of methylben ne. ne.

lustif y your answer by identifying **one** distinguishing feature in your chosen spectrum.





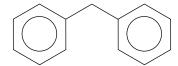


(1) Phenlyethene is used to make the polymer poly(2) henylethene)

Write an equation for the polymerization of phenylethene.

(2)

(e) Diphenylmethane is used to provide the geranium fragrance in soap.



It can be synthesised in five steps from benzene. The first two steps in the process are

() Name the catalyst in Step 1.

(1)

(i)	Write a three step synthesis to convert phenylmagnesium bromide to diphenylmethane, including reagents used in each step.	(6)
	(Total for Question 0 = 4 ma	rks)
	TOTAL FOR PAPER = 120 MA	RKS

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0 (8)	(18) 4.0 He helium 2	20.2 Ne neon	39.9 Ar argon 18	83.8	Kr krypton 36	131.3	Xe	xenon 54	[222]	Rn radon	86	ted						
7	(77)	19.0 F fluorine 9	35.5 Cl chlorine 17	79.9	Br bromine 35	126.9	-	odine 53	[210]	At astatine	85	oeen repor	175	Lu	lutetium 71	[257]	ב	lawrencium 103
9	(16)	16.0 O oxygen 8	32.1 S sulfur 16	79.0	Selenium 34	127.6	<u>Te</u>	tellurium 52	[506]	Po polonium	84	·116 have t	173	ΥÞ	ytterbium 70	[254]	2	nobelium 102
5	(15)	14.0 N nitrogen 7	31.0 P	74.9	As arsenic 33	121.8	Sp	antimony 51	209.0	Bi bismuth	83	tomic numbers 112-116 hav but not fully authenticated	169	Tm	thulium 69	[526]	ÞΨ	mendelevium 101
4	(14)	12.0 C carbon 6	28.1 Si silicon 14	72.6	Ge germanium 32	118.7	Sn	20 ti	207.2	Pb lead	82	atomic nu but not f	167	Ъ	erbium 68	[253]		termium 100
æ	(13)	10.8 B boron 5	27.0 Al aluminium 13	69.7	Ga gallium 31	114.8	드	ndium 49	204.4	T thallium	81	Elements with atomic numbers 112-116 have been reported but not fully authenticated	165		holmium 67	[254]	S :	catifornium einsteinium 98 99
			(12)	65.4	Zn zinc 30	112.4	8	cadmium 48	200.6	Hg mercury	80		163	٥	dysprosium 66	[251]	ָל	98
			(11)	63.5	Cu copper 29	107.9	Ag	silver 47	197.0	Au	79	Rg roentgenium 111	159	ተ	terbium 65	[245]	8	Derketium 97
			(10)	58.7	nickel 28	106.4	Pd	palladium 46	195.1	Pt platinum	78	Ds damstadtium 110	157	В	gadolinium 64			96
			(6)	58.9	Co cobalt 27	102.9	- 3	rhodium 45	192.2	– iridium	77	[268] Mt meitnerium 109	152		europium 63	[243]	Am	95
	1.0 hydrogen	(8)			Fe iron 26	101.1	Ru.	technetium ruthenium 43 44	190.2	Os osmium	92	[277] Hs hassium 108	150		samarium 62	[242]	-	94
			0	54.9	Cr Mn chromium manganese 24 25	[86]			186.2	Re rhenium	75	[264] Bh bohrium 107	[147]	Pa	promethium 61	[237]	δ.	n neptunium plutonium amencium 93 95
		mass bol number	(9)	52.0	Cr chromium 24	95.9	Wo	molybdenum 42	183.8	W tungsten	74	Sg seaborgium 106	144	P	praseodymium neodymium 59 60		>	uranıur 92
	Key	relative atomic mass atomic symbol name atomic (proton) number	(5)	50.9	V vanadium 23	92.9	£	niobium 41	180.9	Ta tantalum	73	[262] Db dubnium 105	141	ዋ	praseodymium 59	[231]	Pa	protactinium 91
		relat ato atomic	(4)	47.9	Ti titanium 22	91.2	Zr	zirconium 40	178.5	Hf hafnium	72	[261] Rf rutherfordium 104	140	o e	cerium 58	232	ַב בּ	thorium 90
(3)				45.0	Sc scandium 21	88.9	>	yttrium 39	138.9	La* lanthanum	22	[227] Ac* actinium 89		Se				
7	(2)	9.0 Be beryllium	24.3 Mg magnesium 12	40.1	Ca calcium 20	87.6	S	strontium 38	137.3	Ba barium	56	[226] Ra radium 88		* Lanthanide series	* Actinide series			
-	(1)	6.9 Li lithium 3	23.0 Na sodium 11	39.1	K potassium 19	85.5	2	rubidium 37	132.9	CS caesium	55	[223] Fr francium 87		* Lanth	* Actini			
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