## ZigZag Practice Exam Papers





Write-on

# Chemistry A Unit H432

### **Practice Paper 1A**

Name	
Itallic	

#### Time allowed

2 hours 15 minutes

#### **Information**

- The total marks available for this paper is 100. The number of marks available for each question is shown in brackets.
- Answer all questions and show all working

#### You will need:

An OCR A Chemistry data sheet

#### You may use:

- A scientific or graphical calculator
- A pencil for graphs and drawings
- A ruler

Question	Mark
MCQs	
16	
17	
18	
19	
20	
21	
Total	

#### **SECTION A**

You should aim to finish this section within 20 minutes.

1 Calculate the relative atomic mass of sulfur based on the following isotopic distribution.

Number of neutrons in isotope	Isotopic abundance (%)
16	94.90
17	0.80
18	4.30

- **A** 32.09
- **B** 32.10
- C 16.09
- **D** 16.10

Your answer

- 2 Identify which of the following can act as a reducing agent for  $Cu^{2+}_{(aq)}$ .
  - A OH-
  - B NH<sub>3</sub>
  - C H<sub>2</sub>O<sub>2</sub>
  - D I-

Your answer

- Which of the following gives the correct ionic equation for the reaction of nitric acid with sodium carbonate?
  - $\textbf{A} \quad Na^{+}{}_{(aq)} + CO_{3}{}^{2-}{}_{(aq)} + 2H^{+}{}_{(aq)} + NO_{3}{}^{-}{}_{(aq)} \Rightarrow Na^{+}{}_{(aq)} + CO_{2(g)} + H_{2}O_{(l)} + NO_{3}{}^{-}{}_{(aq)}$
  - $B \qquad Na^{+}{}_{(aq)} + CO_{3}{}^{2-}{}_{(aq)} + 2H^{+}{}_{(aq)} + NO_{3}{}^{-}{}_{(aq)} \rightarrow Na^{+}{}_{(aq)} + CO_{2(g)} + H_{2(g)} + NO_{3}{}^{-}{}_{(aq)}$
  - $C \quad CO_{3^{2^{+}}\!(aq)} + 2H^{+}\!_{(aq)} \boldsymbol{\rightarrow} NaNO_{3(aq)} + CO_{2(g)} + H_{2}O_{(1)}$
  - $D \quad CO_{3^{2-}(aq)} + 2H^{+}_{(aq)} \rightarrow CO_{2(g)} + H_2O_{(l)}$

Your answer

If 5.0 g of magnesium chloride is obtained from the reaction of 5.0 g of magnesium with hydrochloric acid, what is the percentage yield?

- 25 %
- 50 %
- C 20 %
- 41 %

Your answer	
-------------	--

Define 'enthalpy of formation'.

Enthalpy change when a compound is formed from one mole of its elements at 273 K and 100 kPa, with all reactants and products in their standard states

Enthalpy change when one mole of a compound is formed from one mole of its elements at 273 K and 100 kPa, with all reactants and products in their standard states

Enthalpy change when one mole of a compound is formed from its elements at 298 K and 100 kPa, with all reactants and products in their standard states

Enthalpy change when one mole of a compound is formed from one mole of its elements at 298 K and 100 kPa, with all reactants and products in their standard states

		_
Your ans	swer	

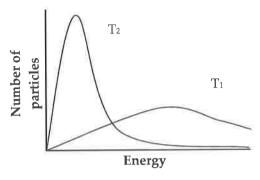
Which of the complexes below cannot exhibit stereoisomerism?

Your answer

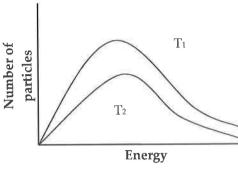
7	Wh Ide	en green copper carbonate is heated using a Bunsen burner, it decomposes to form black CuO. ntify which of the following <b>cannot</b> be used to determine an initial rate of reaction.
	A	Conduct the reaction on a balance. Time taken for the mass of starting material to decrease by a certain amount.
	В	Attach the reaction to a gas syringe. Time taken for a certain volume of gas to be produced (allowing for expansion of air).
	С	Observe the reaction carefully, holding a timer. Time taken for the green copper carbonate to go completely black.
	D	Connect the reaction flask to a U-tube filled with lime water. Time taken for lime water to go milky.
	You	ur answer
8	Ide	ntify which of the following has the most negative value:
	A	The second electron affinity of oxygen
	В	The first electron affinity of oxygen
	C	The first ionisation energy of oxygen
	D	The fifth ionisation energy of oxygen
	Yo	ur answer
9	Th	e indicator bromocresol green exists in two forms:
		$C_{21}H_{12}Br_4O_5S^- \rightleftharpoons C_{21}H_{12}Br_4O_5S^{2-} + H^+$
		Yellow Blue
	WI	nich of the following statements explains the effect of adding acid?
	A	Adding acid increases the concentration of H <sup>+</sup> , causing the position of equilibrium to shift right and making the solution more yellow.
	В	Adding acid increases the concentration of H <sup>+</sup> , causing the position of equilibrium to shift left and making the solution more yellow.
	C	Adding acid decreases the concentration of H <sup>+</sup> , causing the position of equilibrium to shift righ and making the solution more blue.
	D	Adding acid increases the concentration of H <sup>+</sup> , causing the position of equilibrium to shift right and making the solution more blue.
	Yo	our answer

10 Which of the following shows the Maxwell–Boltzmann distribution of molecular energies for a given sample of gas at two temperatures where  $T_1 > T_2$ ?

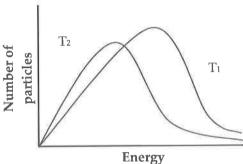
A



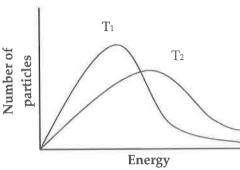
В



C



D



Your answer

11 Which of the following statements is true of a fuel cell?

- A Oxygen is always required, hydrogen or alternatives may be used.
- B Hydrogen is always required, oxygen or alternatives may be used.
- C Oxygen and hydrogen are both always used in a fuel cell.
- D Hydrogen and oxygen are common in fuel cells, but both are optional.

Your answer



12 Iodine atoms combine to form iodine molecules in the gas phase in a second-order reaction. If  $1.00 \times 10^{-5}$  atoms of iodine gives a rate of production of iodine molecules of 0.78 mol dm<sup>-3</sup> s<sup>-1</sup>, calculate the value of the rate constant.

- **A**  $7.8 \times 10^4$
- **B** 7.8 × 10<sup>-4</sup>
- C  $7.8 \times 10^9$
- **D**  $7.8 \times 10^{-9}$

Your answer

13		monium nitrate is a common fertiliser that dissolves in water. Which of the following is true of solution?
	1.	If it is warmed gently, it will turn damp litmus blue.
	2.	It can conduct electricity.
	3.	It will form a white precipitate if aqueous silver ions are added to it.
	A	1 and 2 only
	В	1 and 3 only
	C	2 and 3 only
	D	1, 2 and 3
	Υοι	ar answer
14	Wh	nich of the following statements about ionisation energy are true?
	1.	Oxygen has a higher first ionisation energy than nitrogen because it has an extra proton.
	2.	Boron has a higher first ionisation energy than beryllium because it has a higher nuclear charge.
	A	Statement 1 only
	В	Statement 2 only
	C	Statements 1 and 2
	D	Neither statement 1 nor statement 2
	You	ur answer
15	Wł	nich of the following show a conjugate acid-base pair?
	1.	Acid: NH <sub>3</sub> Base: NH <sub>2</sub> -
	2.	Base: PO <sub>4</sub> <sup>3-</sup> Acid: HPO <sub>4</sub> <sup>2-</sup>
	3.	Acid: ArH <sup>+</sup> Base: Ar
	A	Only 1
	В	Only 1 and 2
	C	Only 2 and 3
	D	1, 2 and 3
	Yo	ur answer

#### **SECTION B**

16	A st	tudent carried out a seri	es of experiments using	g the following electrochemical data	as a basis:
		•	$Ag^{+}_{(aq)} + e^{-} \rightleftharpoons Ag_{(s)}$	E∘=+0.80 V	
		2	$Al^{3+}_{(aq)} + 3e^- \rightleftharpoons Al_{(s)}$	$E_{e} = -1.66 \text{ V}$	
		3	$Fe^{3+}{}_{(aq)}+e^-{\ \rightleftharpoons\ }Fe^{2+}{}_{(aq)}$	$E^{e} = +0.77 \text{ V}$	
	a)	From the list above, ide	entify the best reducing	g agent, explaining your answer.	
					(2)
	b)	Calculate the standard	cell potential if equation	ons ① and ② were combined.	
	ŕ		_		
			<b>30 (50 (4) (50 (5) (5) (5) (5) (5) (5) (5) (5) (5) (5)</b>		
					(1)
	c)	Draw a labelled diagra	m of a circuit that can	be used to experimentally determine	e the emf
	۷,	generated when half co	ells ② and ③ are comb	ined, identifying any key conditions.	
				× ′	
					(4)

d)	Explain why the combination of half cells ${\mathbb O}$ and ${\mathbb O}$ is unlikely to give a reaction even though it may be feasible according to electrochemical values.	
		•
	(1	[)
e)*	Determination of the concentration of $Fe^{2+}$ in a sample may be achieved by titration of the sample with acidified potassium permanganate solution.	
	Outline how this experiment can be used for determination of the concentration of $Fe^{2+}$ in a 25.00 cm <sup>3</sup> sample of factory waste water.	
	You should include: <ul> <li>equations for all reactions taking place</li> <li>equipment needed</li> <li>details of how to carry out the titration</li> </ul>	
	You do not need to include details of calculations or repeats.	
		i.e.
		••
		**
		**
		**
		**
		×
		.11
	(	 6)

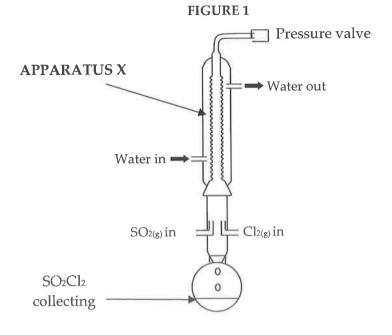
**QUESTION TOTAL: 14 MARKS** 

<b>17</b>			nydroxide, Ba(OH) <sub>2</sub> , is an alkali formed from the dissolution of barium metal in water. forms calcium hydroxide in the same way.	
	a)	Wri	te a balanced equation for the reaction of barium with water.	
			[2	 2]
	b)	A ch	nemist wanted to accurately compare the pH of a solution formed from the same number of es of barium and calcium.	•
		i)	Describe, experimentally, how the chemist would go about making this comparison.	
				•
				***
				3000
				***
				3]
		ii)	Predict, qualitatively, the outcome of the comparison. Justify your prediction.	
				### *****
				[1]
	c)	i)	A solution of Ba(OH)2 was found to have a pH of 10.94 at 298 K. Using the value of $K_w = 1.00 \times 10^{-14}$ , calculate the [OH-] that this contains.	
				•••
				 [2]
		ii)	$50.00 \text{ cm}^3$ of this solution was neutralised by $25.00 \text{ cm}^3$ of nitrous acid, HNO <sub>2</sub> , a monobasic weak acid with a $K_a$ of $4.51 \times 10^{-4}$ . Calculate the concentration of the nitrous acid.	
				••••
				****
				****
			QUESTION TOTAL: 12 MAR	[4] <b>K</b> S

	$SO_{2(g)} + Cl_{2(g)} \rightleftharpoons S$	6O2Cl2(g)	
a)	Explain why, in terms of atom economy, this we	ould be considered a sustainable process.	****
b)	Draw and name the shape of the molecule SO <sub>2</sub> C	Cl2 and predict the bond angle.	
c)	i) Use the following bond energy data to pre		
c)			
c)	i) Use the following bond energy data to pre	dict the enthalpy change for this reaction.  Average bond enthalpy (kJ mol <sup>-1</sup> )  523	
c)	i) Use the following bond energy data to pre  Bond  S=O  S-Cl	Average bond enthalpy (kJ mol <sup>-1</sup> )  523  253	
c)	i) Use the following bond energy data to pre  Bond  S=O	dict the enthalpy change for this reaction.  Average bond enthalpy (kJ mol <sup>-1</sup> )  523	
c)	i) Use the following bond energy data to pre  Bond  S=O  S-Cl	Average bond enthalpy (kJ mol <sup>-1</sup> )  523  253	
c)	i) Use the following bond energy data to pre  Bond  S=O  S-Cl	Average bond enthalpy (kJ mol <sup>-1</sup> )  523  253	
c)	i) Use the following bond energy data to pre  Bond  S=O  S-Cl	Average bond enthalpy (kJ mol <sup>-1</sup> )  523  253	

	ii)	Suggest why this value does not fit with values calculated from the experiment.
		[1]
d)	i)	0.400 moles of $SO_{2(g)}$ were added to an equal number of moles of $Cl_{2(g)}$ at a pressure of 2.00 kPa and temperature of 303 K in a closed system. At equilibrium 0.0240 moles of $SO_{2(g)}$ remained. Calculate the value and unit of $K_P$ .
		[5]
	ii)	The equation can be written to form $SO_2Cl_2$ as a liquid. How does this affect the equation for $K_p$ ?
		[1]

e) The reaction takes place in a vessel, as shown in Figure 1.



#### STAGE 1

- SO<sub>2</sub> and Cl<sub>2</sub> gases are passed into apparatus X
- They react to form SO<sub>2</sub>Cl<sub>2</sub>
- SO<sub>2</sub>Cl<sub>2</sub> collects at the bottom of the flask

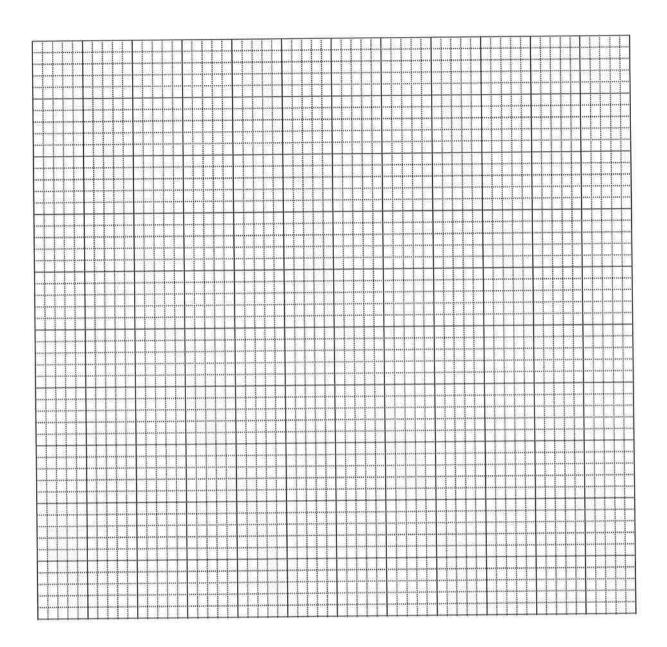
#### **STAGE 2**

- After a period of time, the flask of SO<sub>2</sub>Cl<sub>2</sub> is removed, then connected to distillation apparatus.
- The SO<sub>2</sub>Cl<sub>2</sub> is heated to its boiling point and is distilled into a different flask.

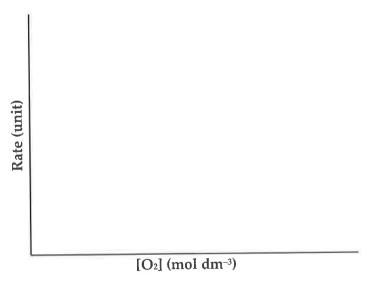
i)	Explain why this experiment would be carried out in a fume cupboard.
	[1]
ii)	Suggest the name and purpose of Apparatus X.
	[2]
iii)	Explain the purpose of Stage 2.

**QUESTION TOTAL: 20 MARKS** 

1)	i)	Explain ho	ow a catalyst works.		
		***************************************			
					***************************************
					***************************************
		EXPLANATION OF ELECTRIC			
	ii)	Identify tv	wo potential disadvantage	es of using a catalyst in an industrial pro	ocess.
					a a wata at aiw
b)	Αl	Iternatively,	instead of studying cataly	ysis, colorimetry can be used to study the	ne rate of air
b)	Al ox	ternatively, idation of Fe	instead of studying cataly e <sup>2+</sup> to Fe <sup>3+</sup> , which is impor	ysis, colorimetry can be used to study the tant for understanding rusting.	ne rate of air
b)	OX	idation of Fe	e <sup>2+</sup> to Fe <sup>3+</sup> , which is impor	tant for understanding rusting.	
b)	ox Us	idation of Fe	e <sup>2+</sup> to Fe <sup>3+</sup> , which is impor	ysis, colorimetry can be used to study the tant for understanding rusting.  of Fe <sup>3+</sup> (aq) formed can be estimated and a	
b)	OX Us	idation of Fosing a colorion	e <sup>2+</sup> to Fe <sup>3+</sup> , which is impor meter, the concentration of against time drawn.	tant for understanding rusting. of Fe³+(aq) formed can be estimated and a	plot of
b)	OX Us co:	idation of Fosing a coloring a coloring a coloring incentration of the follow	e <sup>2+</sup> to Fe <sup>3+</sup> , which is impor meter, the concentration of against time drawn.	tant for understanding rusting.	plot of
0)	OX Us co:	idation of Fosing a coloring a coloring a coloring incentration of the follow	e <sup>2+</sup> to Fe <sup>3+</sup> , which is impor meter, the concentration of against time drawn. ving data on the graph pa	tant for understanding rusting. of Fe³+(aq) formed can be estimated and a	plot of
6)	OX Us co:	idation of Fosing a coloring a coloring a coloring incentration of the follow	e <sup>2+</sup> to Fe <sup>3+</sup> , which is impormeter, the concentration of against time drawn.  ving data on the graph pagiving the unit.	tant for understanding rusting.  of Fe <sup>3+</sup> (aq) formed can be estimated and a  per overleaf, and determine the rate of	plot of
b)	OX Us co:	idation of Fosing a coloring a coloring a coloring incentration of the follow	e <sup>2+</sup> to Fe <sup>3+</sup> , which is impormeter, the concentration of against time drawn.  ving data on the graph pagiving the unit.  Time (s)	tant for understanding rusting.  of $Fe^{3+}_{(aq)}$ formed can be estimated and a per overleaf, and determine the rate of $[Fe(H_2O)_6]^{3+}_{(aq)}$ (mol dm <sup>-3</sup> )	plot of
b)	OX Us co:	idation of Fosing a coloring a coloring a coloring incentration of the follow	e <sup>2+</sup> to Fe <sup>3+</sup> , which is impormeter, the concentration of against time drawn.  ving data on the graph participations the unit.  Time (s)	tant for understanding rusting.  of Fe <sup>3+</sup> (aq) formed can be estimated and a sper overleaf, and determine the rate of  [Fe(H <sub>2</sub> O) <sub>6</sub> ] <sup>3+</sup> (aq) (mol dm <sup>-3</sup> )  0.00	plot of
(b)	OX Us co:	idation of Fosing a coloring a coloring a coloring incentration of the follow	e <sup>2+</sup> to Fe <sup>3+</sup> , which is impormeter, the concentration of against time drawn.  ving data on the graph participating the unit.  Time (s)  0 120	tant for understanding rusting.  of $Fe^{3+}_{(aq)}$ formed can be estimated and a sper overleaf, and determine the rate of	plot of
b)	OX Us co:	idation of Fosing a coloring a coloring a coloring incentration of the follow	e <sup>2+</sup> to Fe <sup>3+</sup> , which is impormeter, the concentration of against time drawn.  ving data on the graph participation of the unit.  Time (s)  0 120 240	tant for understanding rusting.  of Fe <sup>3+</sup> (aq) formed can be estimated and a sper overleaf, and determine the rate of	plot of
b)	OX Us co:	idation of Fosing a coloring a coloring a coloring incentration of the follow	e <sup>2+</sup> to Fe <sup>3+</sup> , which is impormeter, the concentration of against time drawn.  ving data on the graph participation of the unit.  Time (s)  0  120  240  360	tant for understanding rusting.  of Fe <sup>3+</sup> (aq) formed can be estimated and a sper overleaf, and determine the rate of	plot of
b)	OX Us co:	idation of Fosing a coloring a coloring a coloring incentration of the follow	e <sup>2+</sup> to Fe <sup>3+</sup> , which is impormeter, the concentration of against time drawn.  ving data on the graph participation of the unit.  Time (s)  0  120  240  360  480	tant for understanding rusting.  of Fe <sup>3+</sup> (aq) formed can be estimated and a sper overleaf, and determine the rate of	plot of
	OX Us co: Pla 30	sing a coloring a colo	e <sup>2+</sup> to Fe <sup>3+</sup> , which is impormeter, the concentration of against time drawn.  ving data on the graph participation of the unit.  Time (s)  0 120 240 360 480 600	tant for understanding rusting.  of Fe <sup>3+</sup> (aq) formed can be estimated and a oper overleaf, and determine the rate of	plot of reaction at
	OX Us co:	tidation of Festing a colorist incentration of the follows 00 seconds, g	e <sup>2+</sup> to Fe <sup>3+</sup> , which is impormeter, the concentration of against time drawn.  ving data on the graph participation of the unit.  Time (s)  0  120  240  360  480  600	tant for understanding rusting.  of Fe <sup>3+</sup> (aq) formed can be estimated and a sper overleaf, and determine the rate of	plot of reaction at
	OX Us co: Pla 30	sing a coloring a colo	e <sup>2+</sup> to Fe <sup>3+</sup> , which is impormeter, the concentration of against time drawn.  ving data on the graph participation of the unit.  Time (s)  0  120  240  360  480  600	tant for understanding rusting.  of Fe <sup>3+</sup> (aq) formed can be estimated and a oper overleaf, and determine the rate of	plot of reaction at
b) c)	OX Us co: Pla 30	tidation of Festing a colorist incentration of the follows 00 seconds, g	e <sup>2+</sup> to Fe <sup>3+</sup> , which is impormeter, the concentration of against time drawn.  ving data on the graph participation of the unit.  Time (s)  0  120  240  360  480  600	tant for understanding rusting.  of Fe <sup>3+</sup> (aq) formed can be estimated and a oper overleaf, and determine the rate of	plot of reaction at



ii) The reaction is zeroth order with respect to  $[O_2]$ . Sketch the shape of a rate against  $[O_2]$  graph on the axes below.



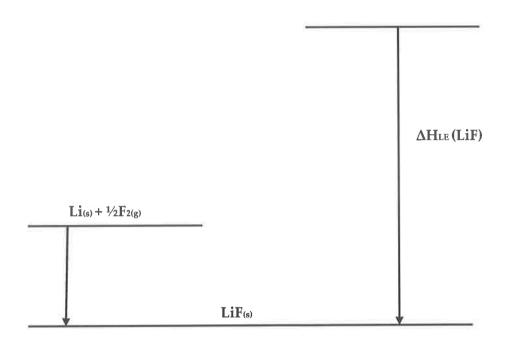
	d)	Explain why it is necessary to calibrate a colorimeter before using it, and explain how the calibration would be done.
		[3]
		QUESTION TOTAL: 16 MARKS
20 <b>*</b>	The	chemistry of transition metal solutions and complexes is varied, and includes: ligand substitution reactions redox reactions
	•	precipitation reactions
	solt	scribe what each of these reactions involves using example equations for reactions of $Cu^{2+}$ or $Fe^{2+}$ ations or complexes, and give an example of the importance of each type of reaction that involves her $Cu^{2+}$ or $Fe^{2+}$ ions.
	*****	
	*****	
	******	
	*****	
	300000	
		Tr'
		QUESTION TOTAL 6 MARKS

21 Lithium fluoride is a crystalline solid that is used as in X-rays and LEDs. It has the second most exothermic lattice enthalpy of any compound for a given mass of starting material. The following data can be used to calculate the lattice energy.

	Enthalpy Change (kJ mol-1)
1st Ionisation Energy (Li)	+520
1st Electron Affinity (F)	-326
Enthalpy of Formation (LiF(s))	-616
Enthalpy of Atomisation (F(g))	+79
Enthalpy of Atomisation (Li(s))	+158

a)	Describe the bonding in lithium fluoride,
	(2)
b)	Define the term 'lattice enthalpy',
	[2]

c) Complete the Born–Haber cycle for lithium fluoride below, labelling the enthalpy changes, and use the data above to calculate the lattice enthalpy of lithium fluoride.



d) Beryllium oxide, BeO, has the most exothermic lattice enthalpy for a given mass of substance.

Explain why BeO has a more exothermic lattice enthalpy than LiF for a given mass of reactant.

[2]

[6]

A student suggested that $LiF_{(g)}$ would be more soluble in water than $NaCl_{(g)}$ because the enthalpy of hydration of $Li^{+}_{(g)}$ and $F^{-}_{(g)}$ would be more exothermic than the enthalpies of hydration of $Na^{+}_{(g)}$ and $Cl^{-}_{(g)}$ . In fact, the enthalpy of solutions of lithium fluoride and sodium chloride are as follows:	ı
Enthalpy of solution (LiF) = 4.73 kJ mol <sup>-1</sup>	
Enthalpy of solution (NaCl) = 3.88 kJ mol <sup>-1</sup>	
Suggest, by referring to enthalpy terms only, why the student's prediction was incorrect.	*****
	[2]
Between the temperatures of 273 K and 373 K , the dissolution of lithium fluoride is feasible. considering the linear nature of the equation $\Delta G = \Delta H - T\Delta S$ , sketch a graph of $\Delta G$ against T over this temperature range, and explain your answer.	Ву
QUESTION TOTAL: 17 MA	[3] RKS
	enthalpy of hydration of Lit and Fr@ would be more exothermic than the enthalpies of hydration of Na' and Clr and Infact, the enthalpy of solutions of lithium fluoride and sodium chloride are as follows:  Enthalpy of solution (LiF) = 4.73 kJ mol <sup>-1</sup> Enthalpy of solution (NaCl) = 3.88 kJ mol <sup>-1</sup> Suggest, by referring to enthalpy terms only, why the student's prediction was incorrect.  Between the temperatures of 273 K and 373 K, the dissolution of lithium fluoride is feasible. considering the linear nature of the equation ΔG = ΔH – TΔS, sketch a graph of ΔG against T over this temperature range, and explain your answer.