Mark schemes

Q1.

(a)	Heat (energy) change at constant pressure Ignore conditions even if wrong Ignore energy change	1
(b)	M2 Ca ²⁺ (g) + 2 e ⁻ + Cl ₂ (g) Alternative M2 Ca ⁺ (g) + e ⁻ + 2 Cl(g)	1
	M3 Ca ²⁺ (g) + 2 Cl ⁻ (g)	1
	M1 Ca(s) + Cl ₂ (g)	1
(c)	M1 –795 + LE = 193 + 590 +1150 + (2 x 121) + (2 x –364) Numbers and factors used correctly from cycle	1
	M2 LE = (+) 2242 (kJ mol ⁻¹) Rearrangement to calculate LE If one or both factors of 2 missing award 1 mark for (+) 2485, (+)2121 or (+)2606 (kJ mol ⁻¹) Allow 1 mark for – 2242 (kJ mol ⁻¹)	1
(d)	$\begin{split} MgCl_2(s) &\to Mg^{2+}(aq) + 2 \ Cl^-(aq) \\ & Allow \ MgCl_2(s) \rightleftharpoons Mg^{2+}(aq) + 2 \ Cl^-(aq) \\ & Allow \ MgCl_2(s) + aq \rightleftharpoons Mg^{2+}(aq) + 2 \ Cl^-(aq) \end{split}$	1
(e)	M1 ΔH soln MgCl ₂ = ΔH latt diss+ ΔH hyd Mg ²⁺ + 2 ΔH hyd Cl- OR 2493 –1920 + (2 x –364) <i>M1 for expression with or without numbers</i>	1
	M2 = – 155 (kJ mol ⁻¹) M2 for answer If factor of 2 missing for ΔH hyd Cl ^{-,} allow 1 mark for 209	1
(f)	M1 Ca ²⁺ (ion) bigger/lower charge to size ratio (than Mg ²⁺) Allow converse answers M1 Do not accept Ca ²⁺ is a bigger atom/molecule M1 Allow Ca ²⁺ has more shells/ more distance of outer e to nucleus Ignore more shielding	

	1
M2 weaker attraction/bond to (O^{δ} in) water	1 [11]

Q2.				
(a)	CO ₂	/ gas is more disordered (than solid)		
		Allow answers based on carbon		
		Ignore CO_2 is a gas and C is a solid	1	
			1	
(b)	0 K			
		Units essential		
		Allow absolute zero OR –273 °C	1	
(c)	M1	$\Delta H = (3^{\times} - 394) - (-1669 \times 2)$		
(0)		$M1 = (3^{-} - 3^{-} 3^{-}) - (-1003 \times 2)$ M1 correct expression		
			1	
	M2	= 2156 (kJ mol ⁻¹)		
		M2 if –2156 seen allow 1 mark out M1 and M2		
			1	
	М3	$\Delta S = (28 \times 4 + 214 \times 3) - (51 \times 2 + 6 \times 3)$		
		M3 correct expression		
			1	
	M4	= 634 (J K ⁻¹ mol ⁻¹)		
		M4 if – 634 allow 1 mark from M4 and M4		
			1	
	M5	$\Delta G = \Delta H - T \Delta S \text{ or } \Delta H = T \Delta S \text{ or } T = \Delta H \div \Delta S$		
		M5 expression or rearranged expression or with		
		numbers	1	
	Me	A = 0.624 k K - 1 m o l - 1		
	M6	$\Delta S = 0.634 \text{ kJ K}^{-1} \text{mol}^{-1}$ M6 $\Delta S = M4 \div 1000$		
			1	
		2156		
	M7	T = 0.634 = 3400 to 3401 (K)		
		$M7 = M2 \div M6$ but must be a positive answer		
			1	[9]
				[3]

Q3.

(a) **M1** $\Delta_{f}H = \Delta_{a}H(Sr) + 2\Delta_{a}H(CI) + \Delta_{1st IE}H(Sr) + \Delta_{2nd IE}H(Sr) + 2\Delta_{EA}H(CI) + \Delta_{LE}H(Sr)$ (Sr)

1

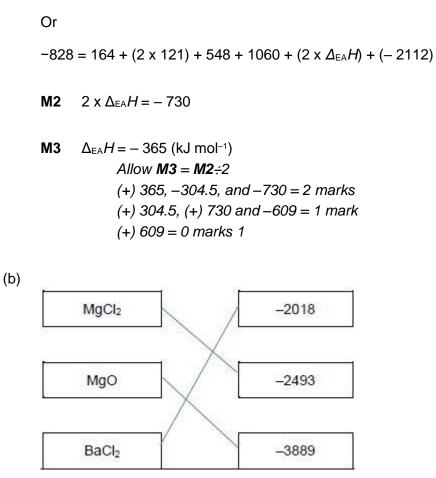
1

1

1

1

1



All three lines must be shown

(c) (Has) covalent character or partial covalent bonding (as well as ionic bonding)

Allow chloride **ion** has been polarised or chloride **ion** distorted Ignore not perfectly ionic Ignore ions are not spheres Do not allow references to molecules or ions with covalent character Do not allow it is covalently bonded alone

(d) **M1** (From Li⁺ to K⁺) size (of ion) increases OR charge density (of ion) decreases

M1 Allow K+ has more shells or larger distance between nucleus and outer electrons or larger ionic radius

Do not allow atomic radius or molecules

M2 (Electrostatic) attraction between metal ion and O^δ- of water decreases or attraction between lone pair on O and + ion decreases

M2 Not dependent on M1

		Allow converse arguments	1
(e)	M1 /		
	or M1	$-110 = 2176 + (-1650) + 2x \Delta_{hyd} H (Br)$	1
	M2	$(2x \Delta_{hyd} H (Br)) = -636$	1
	МЗ	$\Delta_{hyd}H(Br^{-}) = -318 (kJ mol^{-1})$ Allow M3 = M2 ÷ 2 (+)1858, (+)318 and -636 = 2 marks +3716, -1858 and (+)636 = 1 mark -3716 = 0 marks	1 [10]
Q4. (a)	M1	$\Delta H = (-201 + -242) - (-394)$	1
	M2	$\Delta H = -49 \text{ (kJ mol}^{-1}\text{)}$ Allow consequential marking M2 1 mark for $\Delta H = +49 \text{ (kJ mol}^{-1}\text{)}$	1
	М3	$\Delta S = -180$ (J K ⁻¹ mol ⁻¹)	1
	M4	$\Delta G = \Delta H - T\Delta S$ M4 Recall this equation. If M4 incorrect cannot score M6	1
	М5	$\Delta G = -49 - \begin{bmatrix} 890 \text{ x} - \underline{180} \\ 1000 \end{bmatrix} \text{ or } \mathbf{M3} \div 1000$ M5 Conversion of ΔS into kJ mol ⁻¹ $\underline{(890 \times M3)}$ $\Delta G = \mathbf{M2} - \underline{1000}$	
	M6	$\Delta G = 111 \text{ (kJ mol}^{-1}\text{)}$ If ΔS not converted to kJ in M5 , answer is +160151 kJ mol}^{-1} = 5 marks	1
(b)	M1	ΔH = intercept of y axis = 145 (kJ mol ⁻¹) M1 Value between 144 and 146 kJ mol ⁻¹	1

	M2 Gradient = $-\Delta S$ or a negative value			
	М3	Gradient x 1000	1	
	М4	$\Delta S = +167 \text{ to } +173 \text{ (J K}^{-1} \text{ mol}^{-1})$ $M4 + 0.167 \text{ to } +0.173 \text{ scores } 2 \text{ for } \Delta S$ $-167 \text{ to } -173 \text{ scores } 2 \text{ for } \Delta S$ $-0.167 \text{ to } -0.173 \text{ scores } 1 \text{ for } \Delta S$	1	
(C)	<u>Above</u> 845 K reaction becomes (thermodynamically) feasible OR <u>Below</u> 845 K reaction is not (thermodynamically) feasible <i>Allow 845 to 860</i>			[11]
Q5. (a)	Top li		1	
	Lowe	er line $Cs(s) + \frac{1}{2}I_2(s)$	1	
(b)	79 + × + 376 − 314 = −337 +585			
	So er	nthalpy change = 107 (kJ mol ⁻¹) Allow I mark for -107 (kJ mol ⁻¹) Allow answer to 2sf or more	1	
(c)	(Almost/Mostly) purely/ perfectly ionic If ionic not mentioned, allow no/little covalent bonding/character Penalise references to atoms/molecules			
		Ignore electronegativity	1	
(d)	M 1 ∆	S = $[(82.8 + \frac{1}{2} \times 117) - 130] = \frac{11.3}{(J \text{ K}^{-1} \text{ mol}^{-1})}$ M1 Correct entropy change value	1	
	M2 $\Delta G = \Delta H - T\Delta S$ M2 equation or equation with numbers			
	M3 ∆	G = 337 – 298 × 11.3 × 10 ⁻³ OR 337000 – 298 × 11.3 M3 for converting units: ΔS into kJK ⁻¹ mol ⁻¹ or ΔH into J mol ⁻¹		

1

$$M4 \ \Delta G = (+)334 \ kJ \ mol^{-1} \text{ or } 334000 \ J \ mol^{-1}$$

 $M4 \ answer \ with \ correct \ units$
Any negative answer loses $M4$
1
[9]
Q6.
 $Mg^{2+}(g) + 2e^{-} + O(g)$

(a)
$$Mg^{2*}(g) + e^{-} + O^{-}(g)$$
$$Mg^{2*}(g) + e^{-} + O^{-}(g)$$

(b)
$$\Delta_{\rm f} H = \Delta_{\rm a} H (Mg) + \frac{1}{2} \Delta_{\rm BD} H (O_2) + \Delta_{\rm 1st \ IE} H (Mg) + \Delta_{\rm 2nd \ IE} H (Mg) +$$

$$\Delta_{1st EA}H(O) + \Delta_{2nd EA}H(O) + \Delta_{LE}H(MgO)$$

- 602 = 150 + (½ × 496) + 736 +1450 - 142 + 844 + $\Delta_{LE}H(MgO)$

[6]

1 1 1

1

1

1

Q7.

Α

Q9.

(a) <u>Enthalpy change</u> or heat energy change when <u>1 mol</u> of <u>solid ionic</u>

1

	compound/substance or <u>1 mol</u> of <u>ionic lattice</u>	1		
	is formed from its gaseous ions.			
	Allow: <u>enthalpy change</u> for:	1		
	$M^{+}(g) + X^{-}(g) \rightarrow MX(s) \text{ or } Ag^{+}(g) + I^{-}(g) \rightarrow AgI$ (s)			
	$CE=0/2$ if describing wrong process (e.g. ΔH of lattice dissociation or ΔH of formation / or heat energy required)			
	Ignore heat energy released			
(b)	lattice dissociation energy= (112 + 464 + 293) = + 869 (kJmol ⁻ 1)	1		
	lattice formation energy = − 869 (kJ mol ⁻ 1)			
	(+)869 = 1 mark	1		
	A st contains covalent character			
(c)	Agl contains <u>covalent</u> character CE=0/2 if atoms/molecules			
	For M1, allow the following:			
	not completely ionic / ions not spherical / ions distorted / some covalent bonding			
	Forces / bonds (holding the lattice together) are stronger Ignore covalent bonds stronger (than ionic bonds)	1		
	Ignore electronegativity			
	Ignore references to energy			
		1		
(d)	AgNO₃			
	<u>yellow</u> ppt Ignore ammonia/acidified/nitric acid/sulfuric acid			
		1		
	or			
	Cl ₂ or Br ₂			
	brown solution/black ppt			
	M2 dependent on correct M1 but mark on from Ag⁺ or Tollens			
		1		
			[8]	
0.10				
Q10.				
	= Σ_S products – ΣS reactants or + (2 × 198) – (2 × 223 + 2 × 5.7 + 50.2) (= 649 – 507.6)			
200	$T_{2} \times T_{30} = (2 \times 223 + 2 \times 3.1 + 30.2) (= 0+3 = 301.0)$			

This expression could also score M1

∆S = 141(.4) (J K⁻¹mol⁻¹)

			This scores M1 and M2 Allow ecf for M3, M4 and M5 from incorrect M2		
				1	
	∆G∶	= ΔH - TΔS		1	
	$\Delta G = -60 - (1262 \times 141(.4) \times 10^{-3})$				
			This expression also scores M3.		
			For M4, allow $\Delta G = -60 - (1262 \times \text{their } M2 \times 10^{-3})$	1	
				1	
	= -2	38 (kJ mol⁻¹) to 3 sig figs		
			If calculated in joules		
			M4: Allow $\Delta G = -60 \times 10^3 - (1262 \times 141(.4))$		
			M5: Allow - <u>238 000</u> J mol ⁻¹ providing units shown	1	
	feas	Ible since Δ	G is negative/less than zero		
			Allow consequential M6 from their ΔG	1	
					[6]
Q1	11.				
	(a)	Not possib	ble to prevent some dissolving		
			ALLOW It is soluble / dissolves / other hydrates		
			may form / suggestions related to difficulty of measuring T (change) of a solid		
				1	
	(b)	(/hurlH-)-	-155 – (–39)		
	(6)	(2111yu 11 —)	OR labelled cycle		
			Minimum needed for 'labelled cycle'		
			ΔΗ ΔΗ		
			-155 -39 or -155 (+)39		
				1	
		–116 (kJ n	nol ⁻¹)		
		·	1/2 for (+)116 or for –29 or for seeing –116 that has		
			then be processed further	1	
				1	
	(c)		ion is marked using levels of response. Refer to the Mark		
		question	nstructions for examiners for guidance on how to mark this		

Level 3 (5 – 6 marks)

All stages are covered and the explanation of each stage is correct and

virtually complete.

Stage 2 must include use of a graphical method for Level 3 (i.e. 'highest T reached' method is max Level 2)

Answer communicates the whole explanation, including reference to enthalpy, coherently and shows a logical progression through all three stages.Answer is full and detailed and is supported by an appropriate range of relevant points such as those given below:

For the answer to be coherent there must be some indication of **how** the graph is used to find ΔT

Level 2 (3 – 4 marks)

All stages are covered (NB 'covered' means min 2 from each of stage 1 and 3) but the explanation of each stage may be incomplete or may contain inaccuracies

OR two stages covered and the explanations are generally correct and virtually complete

Answer is coherent and shows some progression through all three stages. Some steps in each stage may be out of order and incomplete

Level 1 (1 – 2 marks)

Two stages are covered but the explanation of each stage may be incomplete or may contain inaccuracies **OR** only one stage is covered but the explanation is generally correct and virtually complete

Answer shows some progression between two stages

Level 0 (0 marks)

Insufficient correct Chemistry to warrant a mark

Indicative Chemistry Content

Stage 1 Method

(1a) Measures water with named appropriate apparatus

- (1b) Suitable volume/mass / volume/mass in range 10 200 cm³/g
- (1c) Into insulated container / polystyrene cup (NOT just 'lid')
- (1d) Add known mass of MgCl₂(s)

(1e) Use of 'before and after' weighing method. NOT 'added with washings'

Stage 2 Measurements (could mark from diagram)

(2a) Record initial temperature (min 2 measurements)

(2b) Record T at regular timed intervals for 5+ mins / until trend seen

(2c) Plot T vs time

Stage 3 Use of Results (3a and 3b could come from diagram)

(3a) Extrapolate lines to when solid added (to find initial and final T)

(3b) $T_{\text{final}} - T_{\text{initial}} = \Delta T / \text{ idea of finding } \Delta T \text{ from graph at point of addition}$

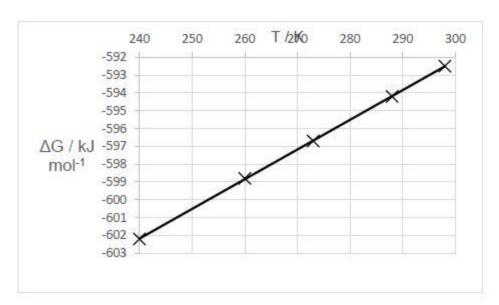
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(3c) $q = mc \Delta T$ (3d) amount = mass/Mr (0.80/95.3 = 8.39 × 10⁻³ mol) (3e) $\Delta H_{soln} = -q/8.39 \times 10^{-3}$ or in words

This could all be described in words without showing actual calculations but describing stages

If method based on 'combustion' Max Level 1





M1 = 5 points correctly plotted
M2 = line drawn correctly (NOT if curved, doubled
or kinked)
(Check line of best fit -if through 250, -600.5 and 280, -595.5 +/- one small
square then award M2, if all crosses on line award
M1 as well)

Gradient = $\Delta (\Delta G) / \Delta T = 0.167$ (kJ K⁻¹ mol⁻¹)

 $(\Delta G = \Delta H - T \Delta S$ so gradient = $-\Delta S$)

 $\Delta S = -167$ (J K⁻¹ mol⁻¹) **M4** = unit conversion i.e. **M3** × 1000; M5 = -sign (process marks) Correct answer with sign gets **M3**, **M4** and M5 ALLOW - 163 to -171

1+1

2

1

[14]

[1]