

A-Level Chemistry

Gibb's Free Energy

Question Paper

Time available: 69 minutes Marks available: 61 marks

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This question is about thermodynamics. Consider the reaction shown.

$$2 \text{ Al}_2\text{O}_3(s) + 3 \text{ C(s)} \rightarrow 4 \text{ Al(s)} + 3 \text{ CO}_2(g)$$

The table below shows some thermodynamic data.

Substance	Al ₂ O ₃ (s)	Al(s)	C(s)	CO ₂ (g)
$\Delta_{\rm f} H^{\Theta}$ / kJ mol $^{-1}$	-1669	0	0	-394
S ^e / J K ⁻¹ mol ⁻¹	51	28	6	214

(a)	Explain why the standard entropy value for carbon dioxide is greater than that for carb	on.
		(1
(b)	State the temperature at which the standard entropy of aluminium is 0 J K ⁻¹ mol ⁻¹	
		(1

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K, at which this reaction becomes feas	sible.	
	Minimum temperature	К
		(7)
		(Total 9 marks)

(c) Use the equation and the data in the table above to calculate the minimum temperature, in

$$CO_2(g) + 3 H_2(g) \rightleftharpoons CH_3OH(g) + H_2O(g)$$

The table contains enthalpy of formation and entropy data for these substances.

	CO ₂ (g)	H ₂ (g)	CH₃OH(g)	H ₂ O(g)
$\Delta_{\rm f}H/{\rm kJ~mol^{-1}}$	-394	0	-201	-242
S / J K ⁻¹ mol ⁻¹	214	131	238	189

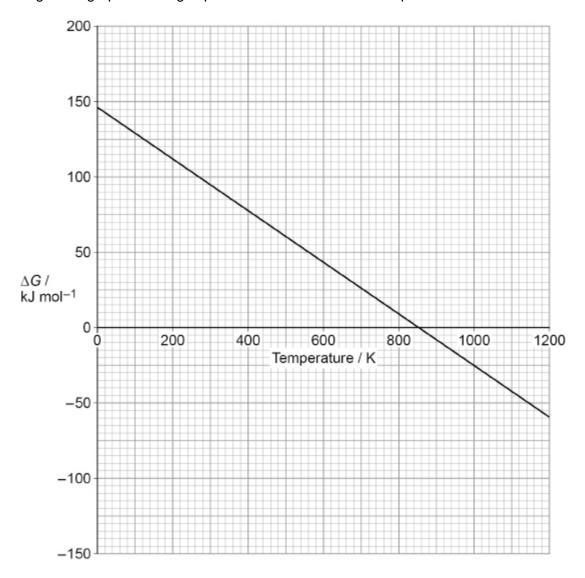
(a) Use the equation and the data in the table above to calculate the Gibbs free-energy change (ΔG) , in kJ mol⁻¹, for this reaction at 890 K

ΔG	kJ	mol^{-1}

(6)

The graph below shows how the Gibbs free-energy change varies with temperature in a different gas phase reaction.

The straight line graph for this gas phase reaction has been extrapolated to zero Kelvin.



٦.	ol ⁻¹ , for this reaction.	m the graph to cal ge (Δ <i>S</i>), in J K ⁻¹ n			
	kJ mol ⁻¹	ΔH			
	J K ⁻¹ mol ⁻¹	Λς			
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The diagram shows an incomplete Born–Haber cycle for the formation of caesium iodide. The diagram is not to scale.

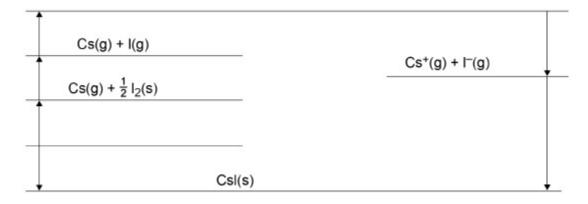


Table 1 gives values of some standard enthalpy changes.

Table 1

Name of enthalpy change	∆ <i>H</i> [⊕] / kJ mol ⁻¹
Enthalpy of atomisation of caesium	+79
First ionisation energy of caesium	+376
Electron affinity of iodine	-314
Enthalpy of lattice formation of caesium iodide	-585
Enthalpy of formation of caesium iodide	-337

(a) Complete the diagram above by writing the formulas, including state symbols, of the appropriate species on each of the two blank lines.

(b) Use the diagram above and the data in **Table 1** to calculate the standard enthalpy of atomisation of iodine.

Standard enthalpy of atomisation of iodine _____ kJ mol⁻¹

(2)

(2)

(c) The enthalpy of lattice formation for caesium iodide in **Table 1** is a value obtained by experiment.

The value obtained by calculation using the perfect ionic model is −582 kJ mol⁻¹

Deduce what these values indicate about the bonding in caesium iodide.

(1)

(d) Use data from Table 2 to show that this reaction is not feasible at 298 K

$$CsI(s) \to Cs(s) + \frac{1}{2}I_2(s)$$
 $\Delta H^{e} = +337 \text{ kJ mol}^{-1}$

Table 2

	CsI(s)	Cs(s)	I ₂ (s)
S [⊕] / J K ⁻¹ mol ⁻¹	130	82.8	117

(4) (Total 9 marks)

4.

The equation for the reaction between ammonia and oxygen is shown.

$$4NH_3(g) + 5O_2(g) = 4NO(g) + 6H_2O(g)$$
 $\Delta H = -905 \text{ kJ mol}^{-1}$

Some standard entropies are given in the table.

Gas	Sº / J K ⁻¹ mol ⁻¹
NH ₃ (g)	193
O ₂ (g)	205
NO(g)	211
H ₂ O(g)	189

)	Calculate the entropy change for the reaction between ammonia and oxygen.
	Entropy change J K ⁻¹ mol ⁻¹
	Calculate a value for the Gibbs free-energy change (ΔG), in kJ mol ⁻¹ , for the reaction between ammonia and oxygen at 600 °C
	(If you were unable to obtain an answer to part (a), you should assume that the entropy change is 211 J K^{-1} mol ⁻¹ . This is not the correct answer.)
	ΔG kJ mol $^{-1}$
	The reaction between ammonia and oxygen was carried out at a higher temperature.
	Explain how this change affects the value of ΔG for the reaction.

	_
	Deduce the change in oxidation state of nitrogen, when NH ₃ is oxidised to NO
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	A// :
	When ammonia reacts with oxygen, nitrous oxide (N_2O) can be produced instead of NO
	Give an equation for this reaction.

5.

Titanium(IV) chloride can be made from titanium(IV) oxide as shown in the equation.

$$TiO_2(s) + 2C(s) + 2Cl_2(g) \rightarrow 2CO(g) + TiCl_4(l)$$

$$\Delta H^{\Theta} = -60.0 \text{ kJ mol}^{-1}$$

Some entropy data are shown in the table.

Substance	TiO ₂ (s)	C(s)	Cl ₂ (g)	CO(g)	TiCl ₄ (I)
S ^e / J K ⁻¹ mol ⁻¹	50.2	5.70	223	198	253

Use the equation and the data in the table to calculate the Gibbs free-energy change for this reaction at $989\,^{\circ}\text{C}$

Give your answer to the appropriate number of significant figures.

Use your answer to explain whether this reaction is feasible.

	Gibbs free-energy change	kJ mol ⁻¹
Explanation		
		(Total 6 marks)

6.

Hydrogen can be manufactured from the reaction of steam with methane.

$$CH_4(g) + H_2O(g) \longrightarrow CO(g) + 3H_2(g)$$

(a) The table contains some enthalpy of formation and entropy data.

Substance	ΔH [÷] _f / kJ mol ⁻¹	S [⊕] / J K ⁻¹ mol ⁻¹
CH ₄ (g)	-75	186
H ₂ O(g)	-242	189
CO(g)	-111	198
H ₂ (g)	0	131
CO ₂ (g)	-394	214

Use data from the table to calculate the enthalpy change, ΔH , for the reaction of steam with methane.	
Use data from the table to calculate the entropy change, ΔS , for the reaction of sto	nam
with methane.	zaiii

(2)

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The te	emperature used for this manufacture of hydrogen is usually about 1300 K.	
	est one reason, other than changing the position of equilibrium, why this tempe	rature
is use	ed rather than the value that you calculated in part (b).	
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		-
Hydro	ogen can also be obtained by reaction of carbon monoxide with steam.	
CO(g) + $H_2O(g) \longrightarrow CO_2(g) + H_2(g)$ $\Delta H = -41 \text{ kJ mol}^{-1}$, $\Delta S = -42 \text{ J K}^{-1} \text{ mol}^{-1}$	
(i)	Explain, using a calculation, why this reaction should not occur at 1300 K.	

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