M1.B

[1]

M2.C

[1]

M3.B

[1]

M4. (a)
$$\Delta H = \Sigma \Delta H_t$$
(products) – $\Sigma \Delta H_t$ (reactants)

1

$$= -201 - 242 - (-394)$$

1

= -49 kJ mol-1

+49 kJ mol⁻¹ = 1 mark

units not required, wrong units lose 1 mark

1

(b)
$$\Delta S = \Sigma S(\text{products}) - \Sigma S(\text{reactants})$$

1

$$= 238 + 189 - (214 + 3 \times 131)$$

1

+180 = 1 mark

units not required, wrong units lose 1 mark

1

(c)
$$\Delta G = \Delta H - T \Delta S$$

If use G not Δ G penalise M1 but not M2 and M3

1

(ΔS is negative so) at high temp $-T\Delta S$ (is positive and) greater than ΔH /large

		Do not award M2 or M3 if positive ΔS value used	1
	So Δ <i>G</i> > 0	Independent mark unless positive ΔS value used	1
	(Limiting condition $\Delta G = 0$ so) $T = \Delta H/\Delta S$		1
	= 272 K	Allow 297-298 if used given values. Do not award M5 if T –ve or if M4 should give T –ve	1
	Reaction is	s too slow at this temperature/to speed up the reaction	1
(d)	CH₃OH +	3/2O₂ → CO₂ + 2H₂O Allow multiples. Ignore state symbols. Do not allow equation for wrong compound but mark on provided number of moles increases or stays the same. If no equation or equation that gives a decrease in the number of moles, CE = 0	1
	2.5 mol giv	ve 3 mol (gases) Allow statement 'increase in number of moles/molecules' If numerical values given, they must match the equation in M1 Ignore the effect of incorrect state symbols on the number of moles of particles unless used correctly	1
	Therefore	ΔS is positive/entropy increases If correct deduction from wrong equation is $\Delta S = 0$ or ΔS very small must say H –ve	1
	•	on exothermic so ΔH –ve so ΔH – $T\Delta S$) and hence snegative (less than zero) Allow G instead of ΔG Can score 3 out of 4 marks if equation wrong but leads to increase or no change in number of moles $M4$ dependent on $M3$ Note, if equation wrong AND there is an incorrect deduction about the change in number of moles, $CE = 0$	1

(e) CO₂/CO/CH₄ may be produced during H₂ manufacture/building the plant/transport/operating the plant

[17]

1

M5. (a) Standard enthalpy change, ΔH^{Θ} : $\Delta H_{R} = \Sigma \Delta H_{foroducts} - \Sigma \Delta H_{freactants}$ (1) or cycle

$$\Delta H_R = (0 + [2 \times -242]) - (4 \times -92)$$
 (1)
= -484 + 368
= -116 (kJ mol⁻¹)
Allow max 1 for +116

Standard entropy change, ΔS^{\bullet} : $\Delta S = \Sigma \Delta H_{f \text{ products}} - \Sigma \Delta H_{f \text{ reactants}}$

$$\Delta S = ([2 \times 223] + [2 \times 189]) - (205 + [4 \times 187])$$
 (1)
= 824 - 953
= -129 (J K⁻¹ mol⁻¹)
allow max one for +129

6

- (b) (i) Effect: Equilibrium displaced to right / to products (1)

 Explanation: Reaction is endothermic (1)

 Constraint reduced (1)

 mark separately
 - (ii) Feasible when $\Delta G \leq 0$ (1)

$$\Delta G = \Delta H - T\Delta S$$
 (1)
 $T = \Delta H/\Delta S = 208 \times 1000$ (1) / 253
= 822 K (1)

[13]

1

1

1

(ii) The negative S⁻ ion

repels the added electron

(iii) Step B is the atomisation enthalpy of sulphur

Step D is the second ionisation enthalpy of calcium

(iv) Electrons nearer to the nucleus

Electrons removed from a positive species or more strongly attracted

(b) The model used assumes the ions are spherical and in a lattice

The calculated value is smaller than the cycle value or stronger attraction

Indicating some covalent character or ions are polarised

(c)	(i)	For a reaction to occur $\Delta G < 0$	1	
		ΔS is positive and large as a gas is evolved	1	
		$T\Delta S$ is larger than ΔH and ΔG is negative	1	
	(ii)	ΔS is negative	1	
		Four moles gaseous reactant forming or more moles of gaseous product	1	
		At high temperature T Δ S is larger than Δ H and Δ G is positive	1	[18]
M8. C				[1]
M9. C				
IVIJ.				[1]