

- M1.** (a) M1 $K_p = (p_Y)^3 \cdot (p_Z)^2 / (p_W)^2 \cdot (p_X)$ NB [] wrong 1
- M2 temperature 1
- M3 increase 1
- M4 particles have more energy or greater velocity/speed 1
- M5 more collisions with $E > E_a$ or more successful collisions 1
- M6 Reaction exothermic or converse 1
- M7 Equilibrium moves in the left 1

Marks for other answers

Increase in pressure or concentration	allow M1, M5, M6	Max 3
Addition of a catalyst;	allow M1, M5, M6	Max 3
Decrease in temperature;	allow M1, M2, M6	Max 3
Two or more changes made;	allow M1, M6	Max 2

- (b) (i) Advantage; reaction goes to completion, not reversible or faster 1

Disadvantage; reaction vigorous/dangerous
(*exothermic must be qualified*)

or HCl(g) evolved/toxic
or CH₃COCl expensive

NB Allow converse answers

Do not allow reactions with other reagents e.g. water
or ease of separation

1

- (ii) $\Delta S = \Sigma S \text{ products} - \Sigma S \text{ reactants}$ 1

□□□□□□□□ $\Delta S = (259 + 187) - (201 + 161)$ 1

□□□□□□□□ $\Delta S = 84 \text{ (JK}^{-1} \text{ mol}^{-1})$ (*Ignore units*)

Allow – 84 to score (1) mark

1

□□□□□□□□ $\Delta G = \Delta H - T\Delta S$

1

$= -21.6 - 298 \times 84/1000$
 $= -46.6 \text{ kJ mol}^{-1}$ or $-46\,600 \text{ J mol}^{-1}$

1

Allow (2) for – 46.6 without units
(Mark ΔG consequentially to incorrect ΔS)
(e.g. $\Delta S = -84$ gives $\Delta G = +3.4 \text{ kJ mol}^{-1}$)

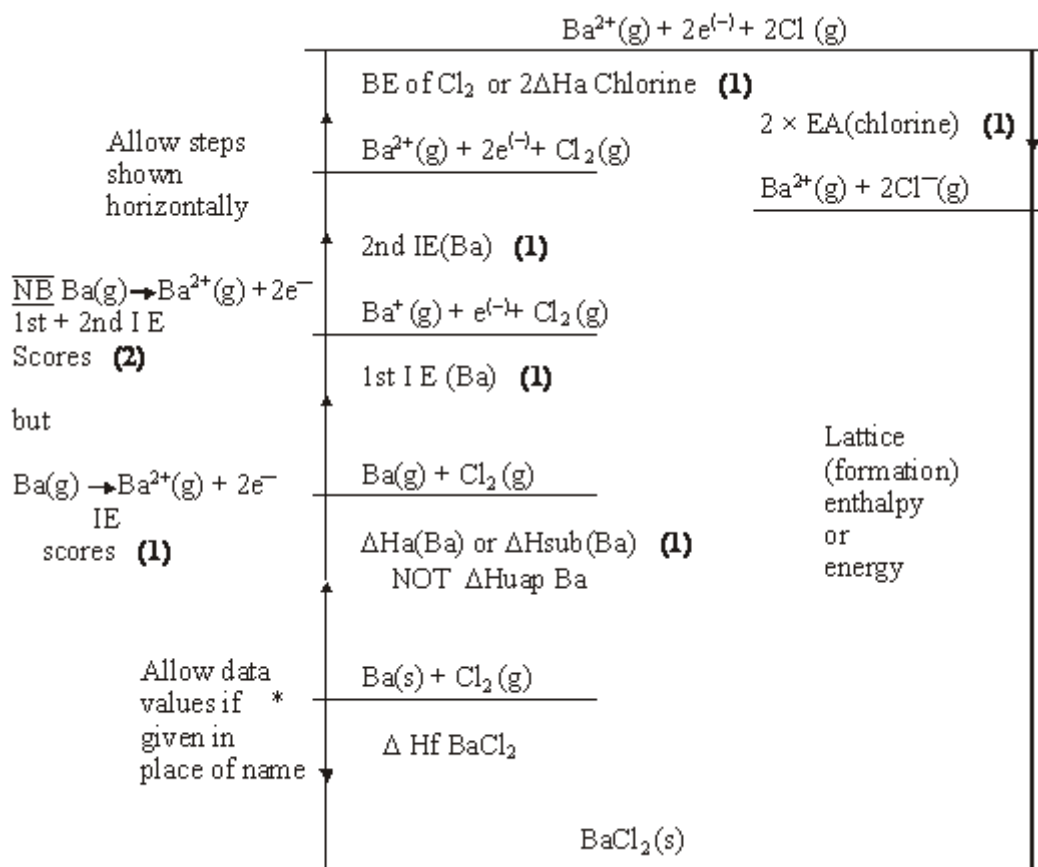
1

[15]

M2.B

[1]

M3. (a) (i)



Allow steps shown horizontally

$\overline{NE} Ba(g) \rightarrow Ba^{2+}(g) + 2e^{-}$
1st + 2nd IE
Scores **(2)**

but

$Ba(g) \rightarrow Ba^{2+}(g) + 2e^{-}$
IE
scores **(1)**

Allow data values if * given in place of name

Lattice (formation) enthalpy or energy

- (ii) Cycling clockwise about (*)
CE if step missing

$$\Delta H_f Ba + 1^{st} IE Ba + 2^{nd} IE Ba + 2\Delta H_a Cl + 2EA Cl + LE - \Delta H_f BaCl_2 = 0 \text{ (1)}$$

$$+180 + 503 + 965 + 2 \times 122 + 2EA - 2056 + 859 = 0 \text{ (1)}$$

$$EA = -695/2 = - (347 \text{ to } 348) \text{ (1)}$$

Ignore units

Calculation -1 for each error

Mark conseq.

Notes: -695 scores (2)

+(347 to 348) scores (2)

-(286 to 287) scores (2)

+(286 to 287) scores (1)

-573 scores (1)

+573 scores (0)

9

- (b) $\Delta S = \Sigma S \text{ products} - \Sigma S \text{ reactants}$
 $= (63 + 223) - 124 = 162 \text{ (1)}$

$$\Delta G = \Delta H - T\Delta S \quad \text{or} \quad \Delta H = T\Delta S \quad \text{or} \quad T = \Delta H/\Delta S \quad (1)$$

or used correctly

$$\Delta H = 859 \times 10^3 \quad (1) = T \times 162$$

$$T = (5\,300 \text{ to } 5304) \text{ K} \quad (1)$$

Penalise if units °C

-1 for each error + mark conseq.

4

[13]

M4.A

[1]

- M5.** (a) (i) *Standard enthalpy change:*
 $\Delta H_r = \Sigma\Delta H_f(\text{products}) - \Sigma\Delta H_f(\text{reactants}) \quad (1)$
or cycle

$$\Delta H_r = ([2 \times 0] + [3 \times -393.5]) - (-824.2 + [3 \times -110.5]) \quad (1)$$

$$= -24.8 \text{ (kJ mol}^{-1}\text{)} \quad (1)$$

Allow + 24.8 max one

Standard entropy change: $\Delta S = \Sigma S(\text{products}) - \Sigma S(\text{reactants}) \quad (1)$

$$\Delta S = ([2 \times 27.3] + [3 \times 213.6]) - (87.4 + [3 \times 197.6]) \quad (1)$$

$$= (54.6 + 640.8) - (87.4 + 592.8) \quad (1)$$

$$= 15.2 \text{ (JK}^{-1}\text{mol}^{-1}\text{)} \quad (1)$$

Allow -15.2 max one

- (ii) $\Delta G = \Delta H - T\Delta S \quad (1)$
 ΔH negative and $-T\Delta S$ is negative (**or ΔS positive or
or correct calc**) (1)
Hence ΔG is always negative
(or feasible when $\Delta G \leq 0$) (1)

9

- (b) $\Delta G = 0 = \Delta H - T\Delta S$ Hence $\Delta H = T\Delta S \quad (1)$

$$T = \Delta H / \Delta S = 492.7 \times 1000 / 542.6 \text{ (1)}$$
$$= 908 \text{ K (1)}$$

penalise missing 1000 by one mark

3

(c) $\Delta G (b) = \Delta G (a)$

$$(492.7 \times 10^3 - T \times 542.6) = (-24.8 \times 10^3 - T \times 15.2) \text{ (1)}$$
$$517.5 \times 10^3 = 527.4 T \text{ (1)}$$
$$T = 981.2 \text{ K(1)}$$

Allow 980 - 982

Penalise missing 1000 by one mark

3

[15]

M6.C

[1]