

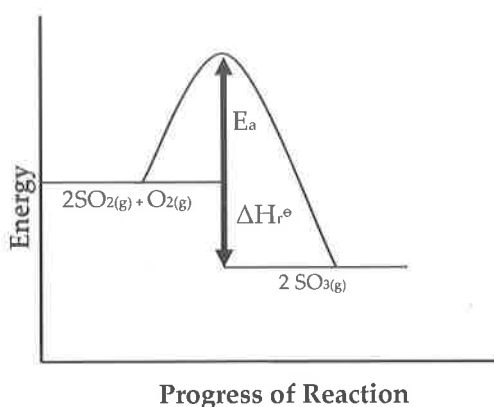
Practice Paper 1B

Section A

- | | | |
|-----|------|------|
| 1 C | 6 B | 11 D |
| 2 B | 7 C | 12 B |
| 3 D | 8 C | 13 B |
| 4 A | 9 A | 14 B |
| 5 D | 10 D | 15 D |

Section B

- 16 a) Sulfur trioxide is symmetrical, but sulfur dioxide is not ✓
Dipoles cancel out in sulfur trioxide ✓
- b) i) $\Delta H_r^\ominus = 2 \times \Delta H_f^\ominus(\text{SO}_3) - 2 \times \Delta H_f^\ominus(\text{SO}_2)$ ✓ (Award for symbols or numbers)
 $\Delta H_r^\ominus = (2 \times -395.7) - (2 \times -296.8)$
 $\Delta H_r^\ominus = -197.8 \text{ (kJ mol}^{-1}\text{)}$ ✓ (Sign must be correct)
- ii) Product line below reactant line and ΔH_r^\ominus line correctly drawn ✓
 E_a correctly labelled (ALLOW double-headed arrow) ✓



- c) i) $K_p = \frac{p(\text{SO}_3)^2}{p(\text{O}_2) \times p(\text{SO}_2)^2}$ ✓
Moles (SO_3) at equilibrium = $\frac{\text{Mass}}{M_r} = \frac{3.443}{80.1} = 0.0430$ ✓

	O ₂	SO ₂	SO ₃
Moles (initial)	0.160	0.160	0.00
Moles (eqm)	0.117	0.0740	0.0430

✓ (Correct eqm moles)

Total moles = 0.234

	O ₂	SO ₂	SO ₃
Partial Pressure	$\frac{0.117 \times 0.5}{0.234}$	$\frac{0.074 \times 0.5}{0.234}$	$\frac{0.043 \times 0.5}{0.234}$
Partial Pressure	0.2500	0.1581	0.0919

✓ (ALLOW ECF from incorrect moles)

$$K_p = \frac{0.0919^2}{0.25 \times 0.1581^2}$$

$K_p = 1.35$ ✓ ALLOW ECF from incorrect partial pressures

$$K_p = \frac{\text{kPa}^2}{\text{kPa} \times \text{kPa}^2} = \text{kPa}^{-1} = \checkmark \text{ (ALLOW Pa}^{-1} \text{ if Pascals used in calculation)}$$

- ii) (Pressure is doubled, so) the partial pressures of each of the reactants and products are initially doubled ✓
 Value of $\frac{p(\text{SO}_3)^2}{p(\text{O}_2) \times p(\text{SO}_2)^2}$ decreases OR denominator increases by more than the numerator (ALLOW argument made using actual figures from (i) ✓
 Equilibrium shifts to the right to restore the value of K_p OR equilibrium shifts to the right to increase the numerator and decrease the denominator ✓

d) i) V_2O_5 ✓

ii) No change ✓

Catalysts speed up both the forward and reverse reactions equally ✓ (IGNORE catalysts do not affect K_p – this is credited by the first mark)

iii) At lower temperatures the reaction is slower ✓

At higher temperatures there is a lower yield ✓

Higher temperatures are more expensive OR require more fuel OR result in more pollution from emissions ✓

17 a) Side product is HCl ✓



b) Reacts to form chlorinated hydrocarbons (which have been linked to ill health effects)

OR Chlorine is toxic ✓

c) To calibrate the pH meter ✓

So readings using the pH meter can be adjusted to the correct values (using a calibration curve) ✓

d) i) Even scale that uses half the graph paper AND axes labelled AND units for volume included ✓

Points plotted correctly ✓

Smooth curve with clear vertical section ✓

ii) Bromocresol purple AND range entirely within the vertical section of the graph ✓

e) i) $K_a = 10^{-pK_a} = 10^{-1.94} = 1.148 \times 10^{-2}$ ✓

$$[\text{H}^+] = \sqrt{K_a \times [\text{HA}]} = \sqrt{1.148 \times 10^{-2} \times 0.01} = 1.072 \times 10^{-2} \text{ mol dm}^{-3} \quad \checkmark$$

$$pH = -\log_{10}[\text{H}^+] = -\log_{10} 0.01072 = 1.97 \quad \checkmark$$

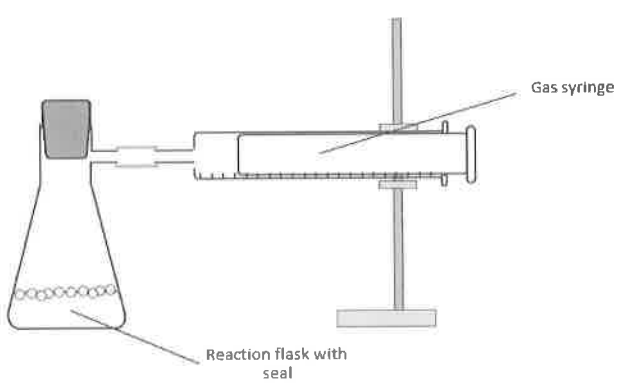
ii) $[\text{OH}^-] = \frac{K_w}{[\text{H}^+]} = \frac{1.00 \times 10^{-14}}{1.072 \times 10^{-2}} = 9.33 \times 10^{-13} \text{ (mol dm}^{-3}) \quad \checkmark$

iii) [HA] at equilibrium cannot be approximated to be the same as the initial [HA] if greater than 5 % dissociation ✓

This gives a $[\text{H}^+]$ that is too small ✓

18 a) (Greater concentration means) there are more particles in a given volume ✓
Therefore, there are more **frequent** (successful) collisions ✓

b)

LEVEL OF RESPONSE QUESTION	
Level 3: (5–6 marks)	The practical procedure is described in a comprehensive and logical fashion. The experimental set-up is completely viable.
Level 2: (3–4 marks)	The practical procedure is described, but omits some details or is not described in a completely logical order. The experimental set-up is mostly correct, with minor errors.
Level 1: (1–2 marks)	Some suggestions for an procedure are given, but not in a logical order. Some appropriate equipment is suggested.
0 marks	No creditworthy response.
Indicative Content	
	
Experimental Set-up	
<ul style="list-style-type: none"> • Experimental diagram with a set-up that allows gas collection (e.g. gas syringe or collection over water). Must be closed to prevent loss of gas to atmosphere. • Labelled piece of equipment in diagram to measure gas production 	
Experimental Description	
<ul style="list-style-type: none"> • Measure out given quantities of Fe (III) solution and H₂O₂ using a measuring cylinder / other appropriate measuring apparatus • Add to the reaction flask and replace bung/seal • Measure the time taken to produce a certain quantity of gas/oxygen • Repeat for different concentrations of Fe (III) • Identification of at least one control variable (e.g. volume of H₂O₂; concentration of H₂O₂; temperature) 	

c) When concentration of H₂O₂ is divided by 3, rate is divided by 3. Rate is first order with respect to H₂O₂. ✓

$$\text{Rate} = k [\text{H}_2\text{O}_2] \text{ so } k = \frac{\text{Rate}}{[\text{H}_2\text{O}_2]} \quad \checkmark$$

$$\text{From reaction 1 (ALLOW use of reaction 2): } k = \frac{3.65 \times 10^{-5}}{5.00 \times 10^{-3}} = 7.3(0) \times 10^{-3} \quad \checkmark$$

$$\text{Units: } k = \frac{\text{mol dm}^{-3} \text{ s}^{-1}}{\text{mol dm}^{-3}} = \text{s}^{-1} \quad \checkmark$$

$$\text{d) } k = \frac{\ln 2}{t_{1/2}} \text{ so } t_{1/2} = \frac{\ln 2}{k} \quad \checkmark$$

$$t_{1/2} = \frac{\ln 2}{7.3 \times 10^{-3}} = 95.0 \text{ (accept 94.95) OR } 6931(.47) \text{ from given value } \quad \checkmark$$

e) Overall order is 2 ✓

Rate is first order with respect to Fe (III) and H₂O₂ so sum of orders of individual reaction is 2. ✓

- 19 a) 4 ✓
- b) A central metal ion surrounded by / coordinately bonded to ligands ✓
- c) Correct groups bound to Pt ✓
Cis square planar complex ✓
Trans complex would have the same ligands on opposite side of the complex ✓



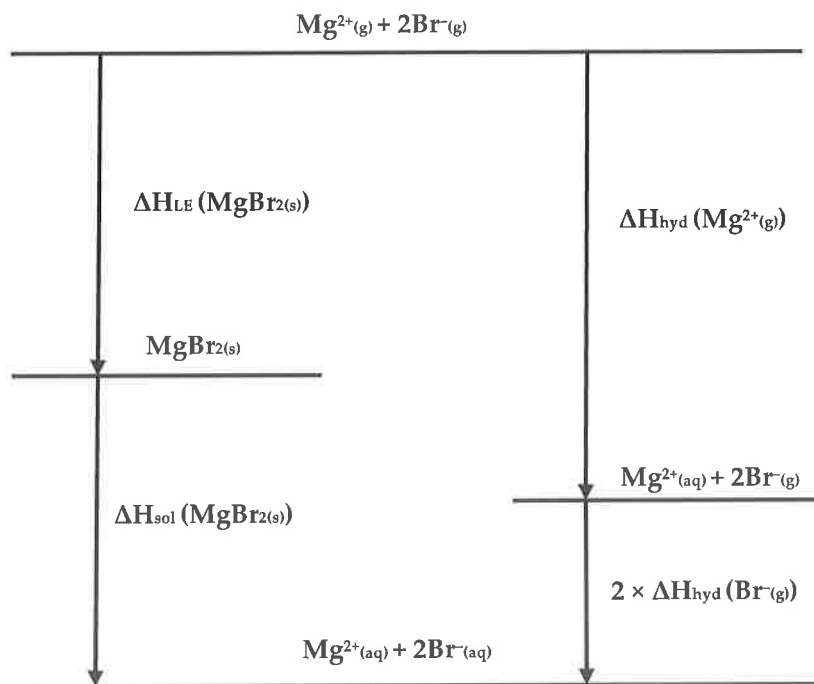
- d) $[\text{PtI}_4]^{2-}(\text{aq}) + 2 \text{NH}_3(\text{aq}) \rightleftharpoons \text{Pt}(\text{NH}_3)_2\text{I}_2(\text{s}) + 2 \text{I}^-(\text{aq})$ Correct, balanced species ✓ State symbols ✓
- e) Binds to the DNA ✓
Stops replication ✓
- f) Adverse side effects (ALLOW valid alternatives) ✓

20 a)

LEVEL OF RESPONSE QUESTION	
Level 3: (5–6 marks)	Answer is structured in an entirely ordered manner. All elements are included in discussion, with characteristics linked to explanations.
Level 2: (3–4 marks)	Answer is mostly well structured. The majority of the elements are included in the discussion, with correct characterisation of melting points in most cases, and some explanation.
Level 1: (1–2 marks)	Answer contains little order. Isolated information about different elements is given, with one or two explanations.
0 marks	No creditworthy response.
Indicative Content	
<ul style="list-style-type: none"> Sodium to aluminium giant metallic (Relatively) high melting points from sodium to aluminium Due to strong electrostatic attraction between delocalised electrons and positive ions 	
<ul style="list-style-type: none"> Melting point increases from sodium to aluminium Due to increasing charge on ions, giving greater electrostatic attraction 	
<ul style="list-style-type: none"> Silicon exists as a giant covalent lattice Very high melting point Due to network of strong covalent bonds 	
<ul style="list-style-type: none"> Phosphorous, sulfur and chlorine are simple molecular / covalent lattices Low melting point Due to weak dispersion forces between molecules 	
<ul style="list-style-type: none"> Argon exists as atoms Very low melting point Due to weak dispersion forces between atoms 	

- b) Simple covalent / molecular ✓

- c) i) (The enthalpy change when) one mole of gaseous ions is dissolved in water ✓
 ii)



Arrows with correct direction ✓

Correct ΔH labels on arrows (IGNORE state symbols; IGNORE size of arrows; ALLOW hydration steps the other way round) ✓

Correct species on each line (DO NOT ALLOW incorrect state symbols) ✓

- iii) $\Delta H_{\text{hyd}}(\text{Mg}^{2+}(\text{g})) = -2432 + -192 - (2 \times -348)$ ✓
 $\Delta H_{\text{hyd}}(\text{Mg}^{2+}(\text{g})) = -1928 \text{ (kJ mol}^{-1}\text{)}$

- 21 a) +6 ✓ (must specify positive)
 b) Zinc reacting with the acid/ H^+ ✓
 c)

	Titration 1	Titration 2	Titration 3	Titration 4
Start reading (cm^3)	0.00	20.90	0.00	21.30
End reading (cm^3)	20.90	41.05	20.30	41.40
Titre (cm^3)	20.90	20.15	20.30	20.10
Mean titre to 1 d.p. (cm^3)	20.1			

Correct titres calculated ✓

Correct mean titre to 1 d.p. ✓

- d) $\text{Cr}_2\text{O}_7^{2-} + 14\text{H}^+ + 6\text{e}^- \rightleftharpoons 7\text{H}_2\text{O} + 2\text{Cr}^{3+}$ ✓
 $\text{Zn}^{2+} + 2\text{e}^- \rightleftharpoons \text{Zn}$ ✓
 $\text{Cr}_2\text{O}_7^{2-} + 3\text{Zn} + 14\text{H}^+ \rightleftharpoons 2\text{Cr}^{3+} + 3\text{Zn}^{2+} + 7\text{H}_2\text{O}$ ✓
 Moles $\text{Cr}_2\text{O}_7^{2-} = 0.0201 \times 0.005 = 1.005 \times 10^{-4}$ ✓ (Accept ECF from incorrect titre)
 Moles $\text{Zn}^{2+} = 3 \times 1.005 \times 10^{-4} = 3.015 \times 10^{-4}$ ✓ (Moles $\text{Cr}_2\text{O}_7^{2-} \times 3$)
 Mass Zn = $3.015 \times 10^{-4} \times 65.4 = 1.972 \times 10^{-2}$ ✓ (Moles Zn $\times 65.4$)
 $\% \text{ by mass} = \frac{\text{Mass of zinc}}{\text{Mass of tablet}} \times 100 = \frac{1.972 \times 10^{-2}}{2.5 \times 10^{-2}} \times 100 = 78.9 \%$ ✓ (Accept ECF)

- e) (Moles of $\text{Cr}_2\text{O}_7^{2-}$ would be too high) so percentage by mass of zinc would be higher than the actual value ✓
 f) It needs to be repeated by someone else / by a different method ✓