

GCE AS/A Level Chemistry
Specified Practical Work
Exemplar results and analysis

1.6 Gravimetric analysis

Identification of an unknown metal chloride

Results

Mass of unknown chloride / g	0.305
Mass of filter paper / g	0.383
Mass of dry filter paper and silver chloride / g	0.957
Mass of silver chloride formed / g	0.584

Analysis

$$n(\text{AgCl}) = \frac{0.584}{143.5} = 0.00407 \text{ mol}$$

1 mol of AgCl formed for every 1 mol of Cl⁻ ions in the unknown chloride

∴ 0.00407 mol of Cl⁻ ions in 0.305 g of the unknown chloride

$$\text{Molar mass of unknown chloride} = \frac{\text{mass}}{\text{number of moles}} = \frac{0.305}{0.00407} = 74.94 \text{ g}$$

This molar mass suggests that there must be only one chloride ion

$$\text{Molar mass of metal} = 74.94 - 35.5 = 39.44 \text{ g}$$

∴ metal must be potassium and unknown chloride must be KCl

1.7 Back titration

Determination of the percentage of calcium carbonate in limestone

Results

Mass of limestone = 0.54 g

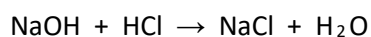
	Rough	Run 2	Run 3	Run 4
Initial volume / cm ³	0.50	0.00	0.35	0.60
Final volume / cm ³	34.50	33.70	34.15	34.30
Titre / cm ³	34.00	33.70	33.80	33.70
Mean titre / cm ³	33.73			

Analysis

33.73 cm³ of NaOH needed to neutralise excess HCl added to the limestone

$$[\text{NaOH}] = 0.098 \text{ M}$$

$$n(\text{NaOH}) = cV = 0.098 \times \frac{33.73}{1000} = 3.31 \times 10^{-3} \text{ mol}$$



$$1 \text{ mol NaOH} : 1 \text{ mol HCl}$$

$$\therefore 3.31 \times 10^{-3} \text{ mol of HCl left over after reaction with CaCO}_3 \text{ in limestone}$$

25.00 cm³ of 0.501 M HCl added initially

$$n(\text{HCl}) = cV = 0.501 \times \frac{25.00}{1000} = 1.25 \times 10^{-2} \text{ mol}$$

Number of moles of HCl that reacted with CaCO₃ = initial amount added – amount left over

$$= 9.19 \times 10^{-3} \text{ mol}$$



$$1 \text{ mol CaCO}_3 : 2 \text{ mol HCl}$$

$$\therefore 4.595 \times 10^{-3} \text{ mol CaCO}_3 \text{ present in limestone sample}$$

$$n(\text{CaCO}_3) = \frac{\text{mass}}{M_r}$$

$$\text{mass} = n(\text{CaCO}_3) \times M_r = 4.595 \times 10^{-3} \times 100 = 0.4595 \text{ g}$$

$$\text{Percentage of CaCO}_3 \text{ in limestone sample} = \frac{0.4595}{0.54} \times 100 = 85.1 \%$$

1.7 Double titration

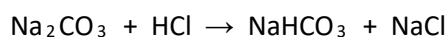
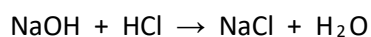
To determine the concentration and the mass of sodium hydroxide and sodium carbonate in a mixed solution

Results

	Rough	Run 2	Run 3	Run 4	Mean
Initial volume / cm ³	0.50	0.10	0.25	0.15	
First end-point volume / cm ³	30.80	29.95	30.05	29.90	
Second end-point volume / cm ³	47.00	46.00	46.20	46.00	
Titre 1 / cm ³	30.30	29.85	29.80	29.75	29.80
Titre 2 / cm ³	16.20	16.05	16.15	16.10	16.10

Analysis

At the first end-point the following reactions are just complete

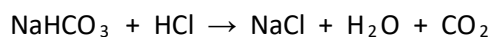


30.00cm³ of 0.100M HCl has reacted

$$n(\text{HCl}) = 0.100 \times \frac{29.80}{1000} = 2.98 \times 10^{-3} \text{ mol}$$

∴ total of 2.98×10^{-3} mol of NaOH and Na₂CO₃ reacted

At the second end-point the following reaction is just complete



16.10cm³ of 0.100M HCl has reacted

$$n(\text{HCl}) = 0.100 \times \frac{16.10}{1000} = 1.61 \times 10^{-3} \text{ mol}$$

∴ 1.61×10^{-3} mol of NaHCO₃ reacted

∴ 1.61×10^{-3} mol of Na₂CO₃ in 25.0cm³ of the original solution

Total of 2.98×10^{-3} mol of NaOH and Na₂CO₃ in 25.0cm³ of the original solution

∴ 1.37×10^{-3} mol of NaOH

$$\text{Concentration of NaOH in mixed solution} = \frac{1.37 \times 10^{-3}}{\frac{25}{1000}} = 0.0548 \text{ M}$$

$$\text{Concentration of Na}_2\text{CO}_3 \text{ in mixed solution} = \frac{1.61 \times 10^{-3}}{\frac{25}{1000}} = 0.0644 \text{ M}$$

$$\begin{aligned} \text{Mass of NaOH in } 1 \text{ dm}^3 \text{ of mixed solution} &= \text{number of moles} \times \text{molar mass} \\ &= 0.0548 \times 40 = 2.19 \text{ g} \end{aligned}$$

$$\begin{aligned} \text{Mass of Na}_2\text{CO}_3 \text{ in } 1 \text{ dm}^3 \text{ of mixed solution} &= \text{number of moles} \times \text{molar mass} \\ &= 0.0644 \times 106 = 6.83 \text{ g} \end{aligned}$$

2.1 Indirect determination of an enthalpy change of reaction

Indirect determination of the enthalpy change of reaction of magnesium oxide and carbon dioxide to form magnesium carbonate

Results

Magnesium oxide

Mass of MgO + weighing boat / g	7.359
Mass of weighing boat / g	6.438
Mass of MgO / g	0.921

Time / min	Temperature / °C
0	23.4
0.5	23.3
1.0	23.2
1.5	23.2
2.0	23.2
2.5	23.2
3.0	
3.5	28.7
4.0	33.2
4.5	36.6

Time / min	Temperature / °C
5.0	36.4
5.5	36.4
6.0	36.3
6.5	36.0
7.0	35.8
7.5	35.7
8.0	35.6
8.5	35.4
9.0	35.4

Magnesium carbonate

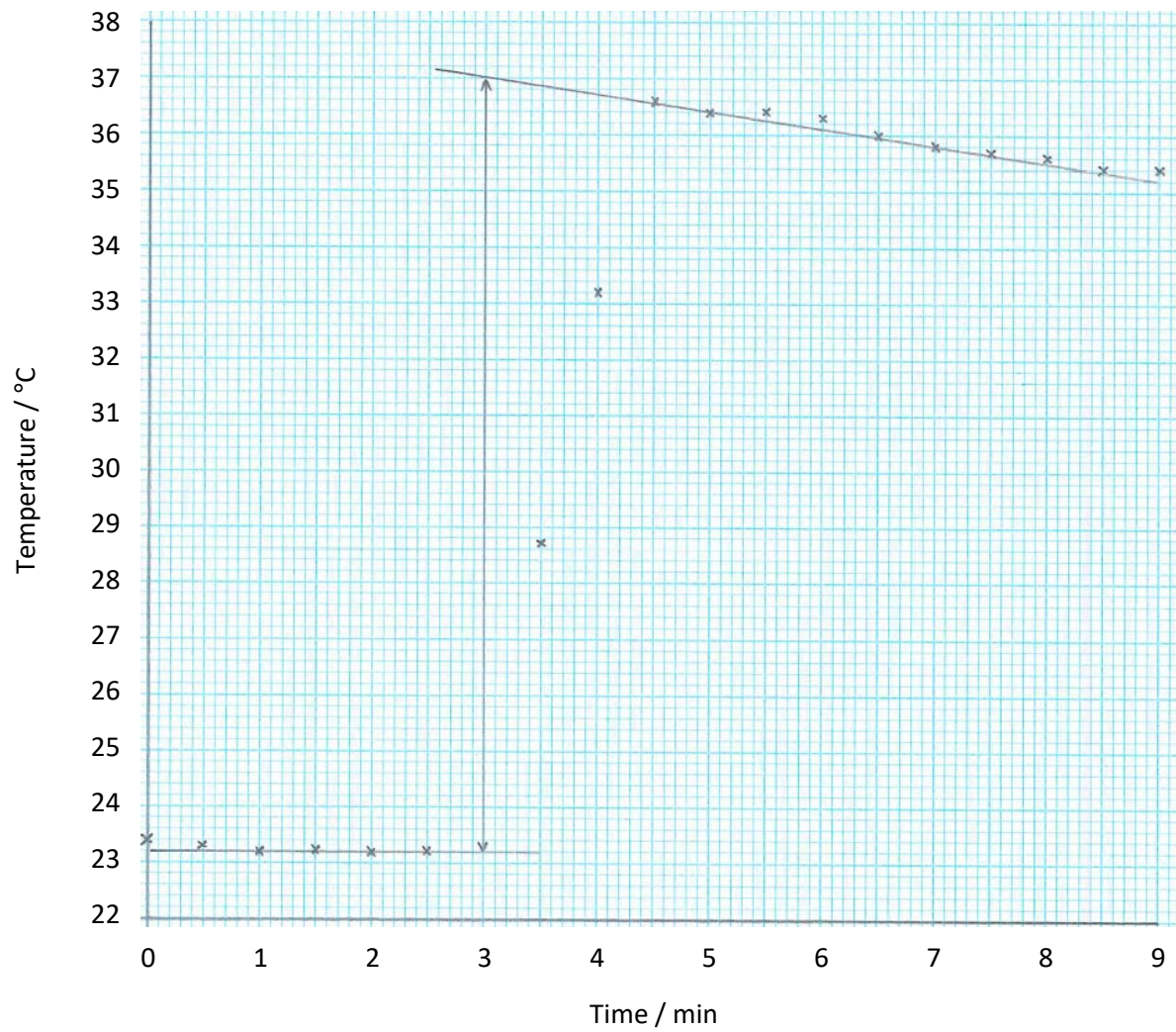
Mass of MgCO ₃ + weighing boat / g	10.015
Mass of weighing boat / g	6.526
Mass of MgCO ₃ / g	3.489

Time / min	Temperature / °C
0	24.2
0.5	24.1
1.0	24.0
1.5	24.0
2.0	24.0
2.5	24.0
3.0	
3.5	32.8
4.0	32.8
4.5	32.3

Time / min	Temperature / °C
5.0	32.2
5.5	32.2
6.0	32.0
6.5	31.9
7.0	31.8
7.5	31.6
8.0	31.4
8.5	31.2
9.0	31.0

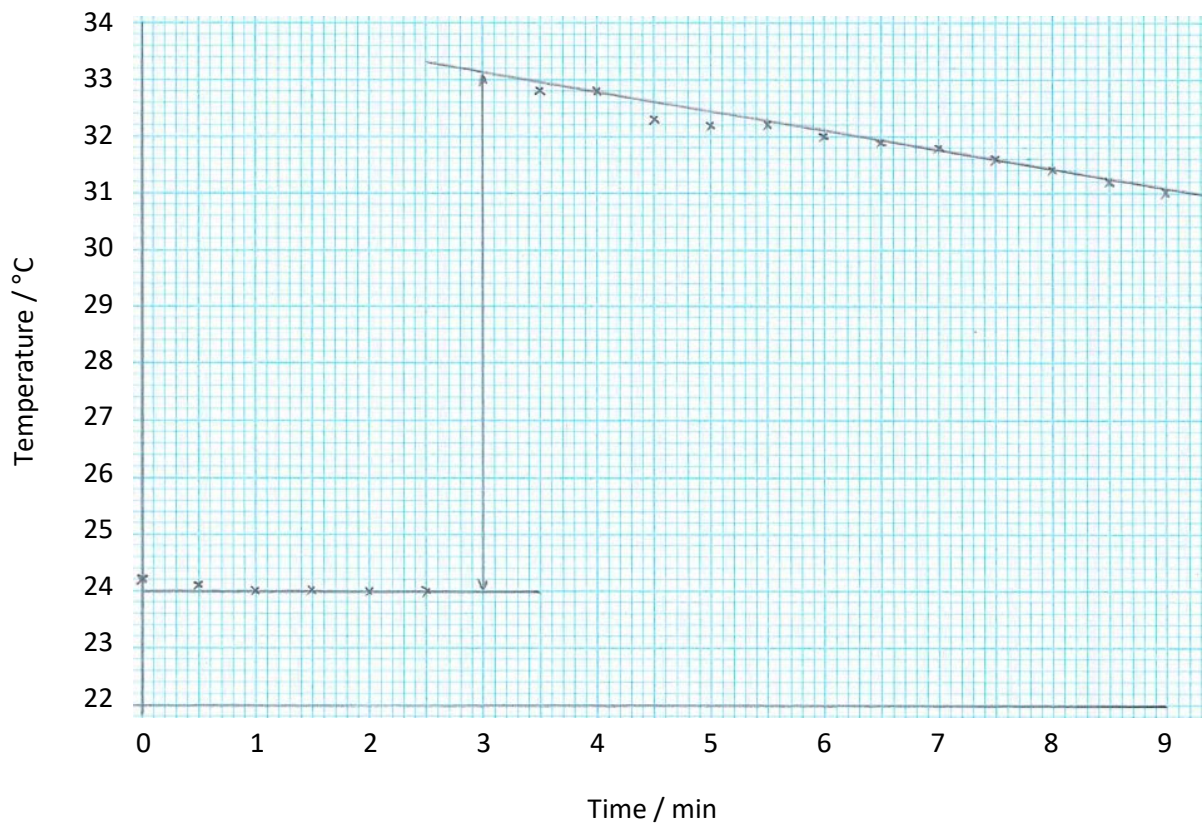
Analysis

Magnesium oxide

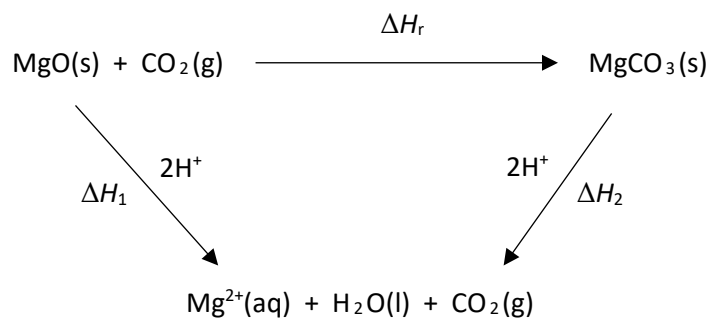


From magnesium oxide graph, the maximum temperature change $\Delta T = 13.8^\circ\text{C}$

Magnesium carbonate



From magnesium carbonate graph, the maximum temperature change $\Delta T = 9.1^{\circ}\text{C}$



Enthalpy of reaction of MgO, ΔH_1

$$\Delta H_1 = -\frac{mc\Delta T}{n}$$

ΔT from graph = 13.8 °C

$$n(\text{MgO}) = \frac{0.921}{40.3} = 0.0229 \text{ mol}$$

$$\Delta H_1 = -\frac{mc\Delta T}{n} = -\frac{50 \times 4.18 \times 13.8}{0.0229} = -126 \text{ kJ mol}^{-1}$$

Enthalpy of reaction of MgCO_3 , ΔH_2

$$\Delta H_2 = -\frac{mc\Delta T}{n}$$

ΔT from graph = 9.1 °C

$$n(\text{MgCO}_3) = \frac{3.489}{84.3} = 0.0414 \text{ mol}$$

$$\Delta H_2 = -\frac{mc\Delta T}{n} = -\frac{50 \times 4.18 \times 9.1}{0.0414} = -45.9 \text{ kJ mol}^{-1}$$

From Hess' cycle

$$\Delta H_r + \Delta H_2 = \Delta H_1$$

$$\Delta H_r = \Delta H_1 - \Delta H_2 = -126 - (-45.9) = -80.1 \text{ kJ mol}^{-1}$$

3.2 Estimation of copper in copper(II) salts

To determine the percentage by mass of copper in copper(II) sulfate crystals

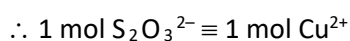
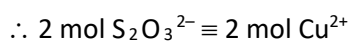
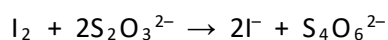
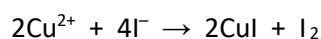
Results

Mass of copper(II) sulfate = 6.01 g

Concentration of $\text{Na}_2\text{S}_2\text{O}_3 = 0.0992 \text{ M}$

	Rough	Run 2	Run 3	Run 4
Initial volume / cm^3	0.50	0.15	24.35	0.10
Final volume / cm^3	25.05	24.35	48.70	24.30
Titre / cm^3	24.55	24.20	24.35	24.20
Mean titre / cm^3	24.25			

Analysis



$$n(\text{Na}_2\text{S}_2\text{O}_3) = 0.0992 \times \frac{24.25}{1000} = 2.406 \times 10^{-3} \text{ mol}$$

$\therefore 2.406 \times 10^{-3} \text{ mol } \text{Cu}^{2+}$ in 25.0 cm^3 of solution

$\therefore 2.406 \times 10^{-2} \text{ mol } \text{Cu}^{2+}$ in 250 cm^3 of solution

Mass of Cu^{2+} ions in 250 cm^3 of solution = $2.406 \times 10^{-2} \times 63.5 = 1.528 \text{ g}$

Percentage by mass of copper in copper(II) sulfate crystals = $\frac{1.528}{6.01} \times 100 = 25.4\%$

3.5 Determination of the order of a reaction

To determine the order of reaction for the oxidation of iodide ions by hydrogen peroxide in acid solution

Results

Volume H ₂ O ₂ / cm ³	Volume H ₂ O / cm ³	*[H ₂ O ₂] / M	Time / s
5	0	0.010	26
4	1	0.008	32
3	2	0.006	42
2	3	0.004	64
1	4	0.002	126

* Concentration of H₂O₂ in total 50 cm³ of solution

Analysis

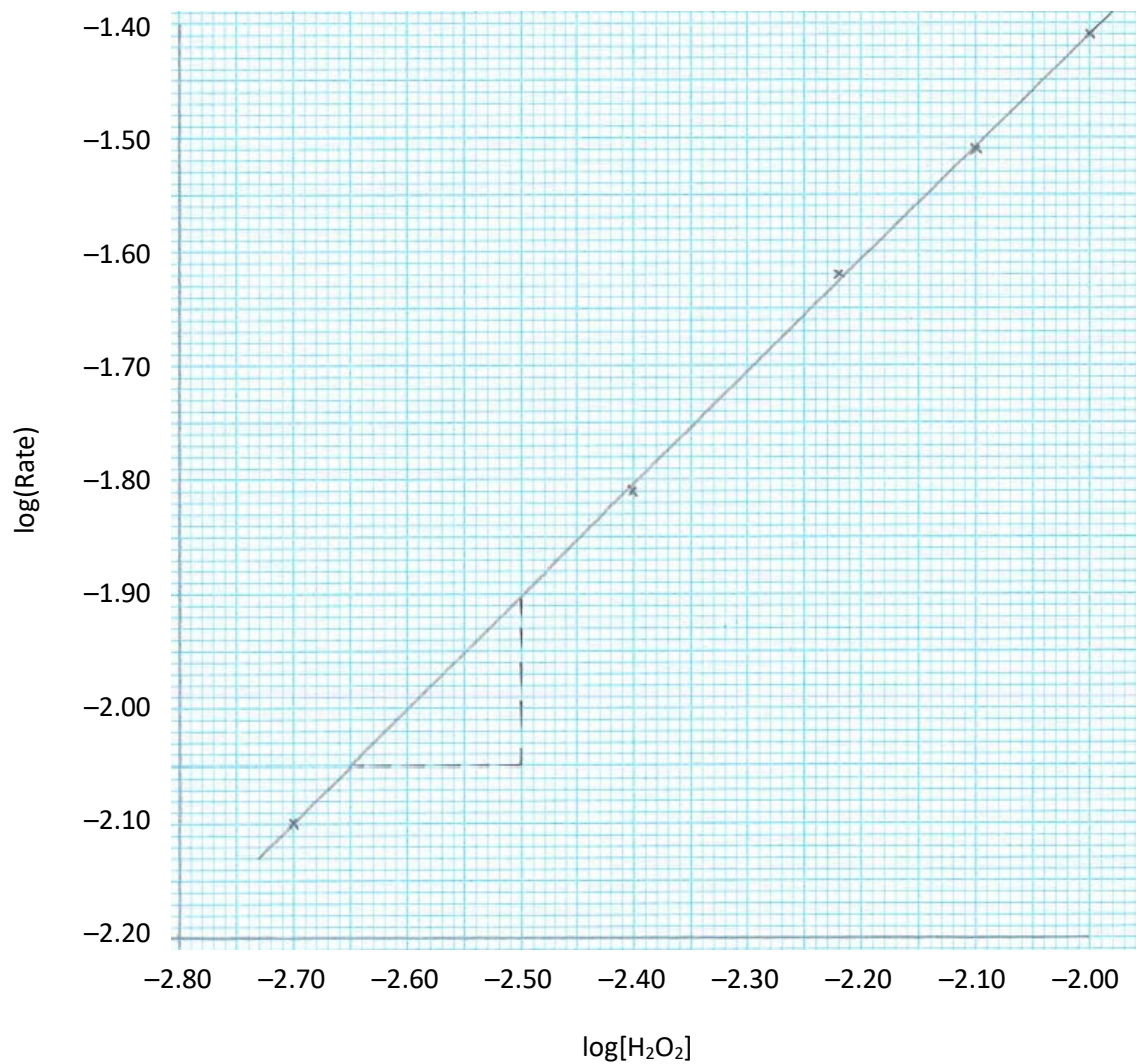
[H ₂ O ₂] / M	log[H ₂ O ₂]	Time / s	Rate / s ⁻¹	log(Rate)
0.010	-2.00	26	0.0385	-1.41
0.008	-2.10	32	0.0313	-1.51
0.006	-2.22	42	0.0238	-1.62
0.004	-2.40	64	0.0156	-1.81
0.002	-2.70	126	0.0079	-2.10

rate = $k \times [\text{H}_2\text{O}_2]^m$ where m is the order of reaction with respect to H₂O₂

$\therefore \log(\text{rate}) = \log k + m \log[\text{H}_2\text{O}_2]$

$$y = c + mx$$

For graph of log(rate) on y-axis against log[H₂O₂] on x-axis, gradient m is the order of reaction



Gradient m

$$m = \frac{\text{change in log (rate)}}{\text{change in log [H}_2\text{O}_2]} = \frac{0.145}{0.15} = 0.97$$

∴ reaction is first order with respect to H₂O₂

3.8 Determination of an equilibrium constant

To determine the equilibrium constant for the reaction of ethanoic acid with ethanol

Part 1

Analysis

Initial number of moles of ethanoic acid, ethanol and water in the reaction mixture

ethanoic acid, CH_3COOH

$$\text{mass} = \text{density} \times \text{volume} = 1.05 \times 6.0 = 6.30 \text{ g}$$

$$M_r = 60$$

$$\text{number of moles} = \frac{\text{mass}}{M_r} = \frac{6.30}{60} = 0.105 \text{ mol}$$

ethanol, $\text{C}_2\text{H}_5\text{OH}$

$$\text{mass} = \text{density} \times \text{volume} = 0.79 \times 6.0 = 4.74 \text{ g}$$

$$M_r = 46$$

$$\text{number of moles} = \frac{\text{mass}}{M_r} = \frac{4.74}{46} = 0.103 \text{ mol}$$

water, H_2O ($6 \text{ cm}^3 + 2 \text{ cm}^3$ from the HCl solution)

$$\text{mass} = \text{density} \times \text{volume} = 1.00 \times 8.0 = 8.00 \text{ g}$$

$$M_r = 18$$

$$\text{number of moles} = \frac{\text{mass}}{M_r} = \frac{8.00}{18} = 0.444 \text{ mol}$$

Part 2

Results

Control mixture titration

	Rough	Run 2	Run 3	Run 4
Initial volume / cm ³	1.20	22.10	0.10	20.75
Final volume / cm ³	22.10	42.85	20.75	41.45
Titre / cm ³	20.90	20.75	20.65	20.70
Mean titre / cm ³		20.70		

Reaction mixture titration

	Rough	Run 2	Run 3	Run 4
Initial volume / cm ³	0.00	0.85	0.25	0.40
Final volume / cm ³	34.70	35.20	34.65	34.70
Titre / cm ³	34.70	34.35	34.40	34.30
Mean titre / cm ³		34.35		

Analysis

Number of moles of HCl in the control mixture

concentration of NaOH = 0.0992 M

$$n(\text{NaOH}) = c \times V = 0.0992 \times \frac{20.70}{1000} = 0.002053 \text{ mol}$$

1 mol NaOH : 1 mol HCl

∴ 0.002053 mol of HCl in the control mixture

Number of moles of H⁺ ions in 1 cm³ of the reaction mixture

concentration of NaOH = 0.0992 M

$$n(\text{NaOH}) = c \times V = 0.0992 \times \frac{34.45}{1000} = 0.00341744 \text{ mol}$$

1 mol NaOH : 1 mol H⁺ ions

∴ 0.00341744 mol of H⁺ ions in 1 cm³ of the reaction mixture

0.0683488 mol of H⁺ ions in 20 cm³ of the reaction mixture

H⁺ ions come from original HCl which remains unchanged and the CH₃COOH present at equilibrium

$$\therefore n(\text{CH}_3\text{COOH}) = 0.0683488 - 0.002053 = 0.0662958 \text{ mol}$$

This information is summarised as follows

Initial concentration	0.105		0.103		0		0.444
	CH ₃ COOH	+	C ₂ H ₅ OH	⇌	CH ₃ COOC ₂ H ₅	+	H ₂ O
Equilibrium concentration	0.0662958		0.0642958		0.0387042		0.4827042

For this equilibrium

$$K_c = \frac{[CH_3COOC_2H_5] \times [H_2O]}{[CH_3COOH] \times [C_2H_5OH]}$$

$$K_c = \frac{[0.0387042] \times [0.4827042]}{[0.0662958] \times [0.0642958]}$$

$$K_c = \frac{0.0186826799}{0.004262541498} = 4.383$$