

**Chemistry**  
**Higher level**  
**Paper 2**

Monday 14 November 2016 (morning)

Candidate session number

2 hours 15 minutes

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**Instructions to candidates**

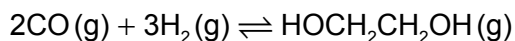
- Write your session number in the boxes above.
- Do not open this examination paper until instructed to do so.
- Answer all questions.
- Write your answers in the boxes provided.
- A calculator is required for this paper.
- A clean copy of the **chemistry data booklet** is required for this paper.
- The maximum mark for this examination paper is **[95 marks]**.



Answer **all** questions. Write your answers in the boxes provided.

1. Ethane-1,2-diol, HOCH<sub>2</sub>CH<sub>2</sub>OH, has a wide variety of uses including the removal of ice from aircraft and heat transfer in a solar cell.

(a) Ethane-1,2-diol can be formed according to the following reaction.



(i) Deduce the equilibrium constant expression,  $K_c$ , for this reaction. [1]

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(ii) State how increasing the pressure of the reaction mixture at constant temperature will affect the position of equilibrium and the value of  $K_c$ . [2]

Position of equilibrium:  
.....  
 $K_c$ :  
.....

(iii) Calculate the enthalpy change,  $\Delta H^\ominus$ , in kJ, for this reaction using section 11 of the data booklet. The bond enthalpy of the carbon–oxygen bond in CO (g) is 1077 kJ mol<sup>-1</sup>. [3]

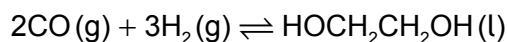
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**(Question 1 continued)**

- (b) (i) Calculate  $\Delta H^\ominus$ , in kJ, for this similar reaction below using  $\Delta H_f^\ominus$  data from section 12 of the data booklet.  $\Delta H_f^\ominus$  of  $\text{HOCH}_2\text{CH}_2\text{OH}(\text{l})$  is  $-454.8 \text{ kJ mol}^{-1}$ . [1]



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- (ii) Deduce why the answers to (a)(iii) and (b)(i) differ. [1]

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- (iii)  $\Delta S^\ominus$  for the reaction in (b)(i) is  $-620.1 \text{ J K}^{-1}$ . Comment on the decrease in entropy. [1]

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- (iv) Calculate the value of  $\Delta G^\ominus$ , in kJ, for this reaction at 298 K using your answer to (b)(i). (If you did not obtain an answer to (b)(i), use  $-244.0 \text{ kJ}$ , but this is not the correct value.) [2]

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**(Question 1 continued)**

- (v) Comment on the statement that the reaction becomes less spontaneous as temperature is increased.

[1]

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- (c) Determine the average oxidation state of carbon in ethene and in ethane-1,2-diol.

[2]

Ethene:  
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Ethane-1,2-diol:  
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- (d) Explain why the boiling point of ethane-1,2-diol is significantly greater than that of ethene.

[2]

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- (e) Ethane-1,2-diol can be oxidized first to ethanedioic acid,  $(\text{COOH})_2$ , and then to carbon dioxide and water. Suggest the reagents needed to oxidize ethane-1,2-diol.

[1]

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**(Question 1 continued)**

- (f) Predict the  $^1\text{H}$ NMR data for ethanedioic acid and ethane-1,2-diol by completing the table.

[2]

	<b>Number of signals</b>	<b>Splitting pattern</b>
<b>Ethanedioic acid:</b>	.....	.....
<b>Ethane-1,2-diol:</b>	.....	Not required



2. The concentration of a solution of a weak acid, such as ethanedioic acid, can be determined by titration with a standard solution of sodium hydroxide, NaOH (aq).

(a) Distinguish between a weak acid and a strong acid.

[1]

Weak acid:  
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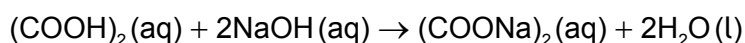
Strong acid:  
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(b) Suggest why it is more convenient to express acidity using the pH scale instead of using the concentration of hydrogen ions.

[1]

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(c) 5.00 g of an impure sample of hydrated ethanedioic acid,  $(\text{COOH})_2 \cdot 2\text{H}_2\text{O}$ , was dissolved in water to make  $1.00 \text{ dm}^3$  of solution.  $25.0 \text{ cm}^3$  samples of this solution were titrated against a  $0.100 \text{ mol dm}^{-3}$  solution of sodium hydroxide using a suitable indicator.



The mean value of the titre was  $14.0 \text{ cm}^3$ .

(i) Suggest a suitable indicator for this titration. Use section 22 of the data booklet.

[1]

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**(Question 2 continued)**

- (e) Outline why all the C–O bond lengths in the ethanedioate ion are the same length and suggest a value for them. Use section 10 of the data booklet. [2]

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- (f) Explain how ethanedioate ions act as ligands. [2]

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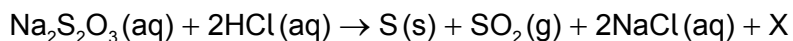
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3. Sodium thiosulfate solution reacts with dilute hydrochloric acid to form a precipitate of sulfur at room temperature.



- (a) Identify the formula and state symbol of X. [1]

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- (b) Suggest why the experiment should be carried out in a fume hood or in a well-ventilated laboratory. [1]

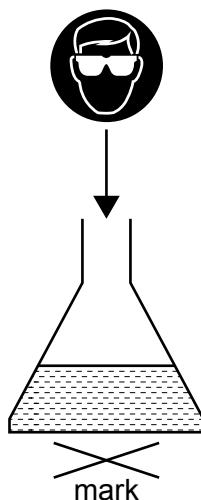
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**(Question 3 continued)**

- (c) The precipitate of sulfur makes the mixture cloudy, so a mark underneath the reaction mixture becomes invisible with time.



10.0 cm<sup>3</sup> of 2.00 mol dm<sup>-3</sup> hydrochloric acid was added to a 50.0 cm<sup>3</sup> solution of sodium thiosulfate at temperature, T<sub>1</sub>. Students measured the time taken for the mark to be no longer visible to the naked eye. The experiment was repeated at different concentrations of sodium thiosulfate.

Experiment	[Na <sub>2</sub> S <sub>2</sub> O <sub>3</sub> (aq)] / mol dm <sup>-3</sup>	Time, t, for mark to disappear / s ± 1 s	$\frac{1^*}{t} / 10^{-3} \text{ s}^{-1}$
1	0.150	23	43.5
2	0.120	27	37.0
3	0.090	36	27.8
4	0.060	60	16.7
5	0.030	111	9.0

\* The reciprocal of the time in seconds can be used as a measure of the rate of reaction.

[Source: Adapted from <http://www.flinnsci.com/>]

Show that the hydrochloric acid added to the flask in experiment 1 is in excess.

[2]

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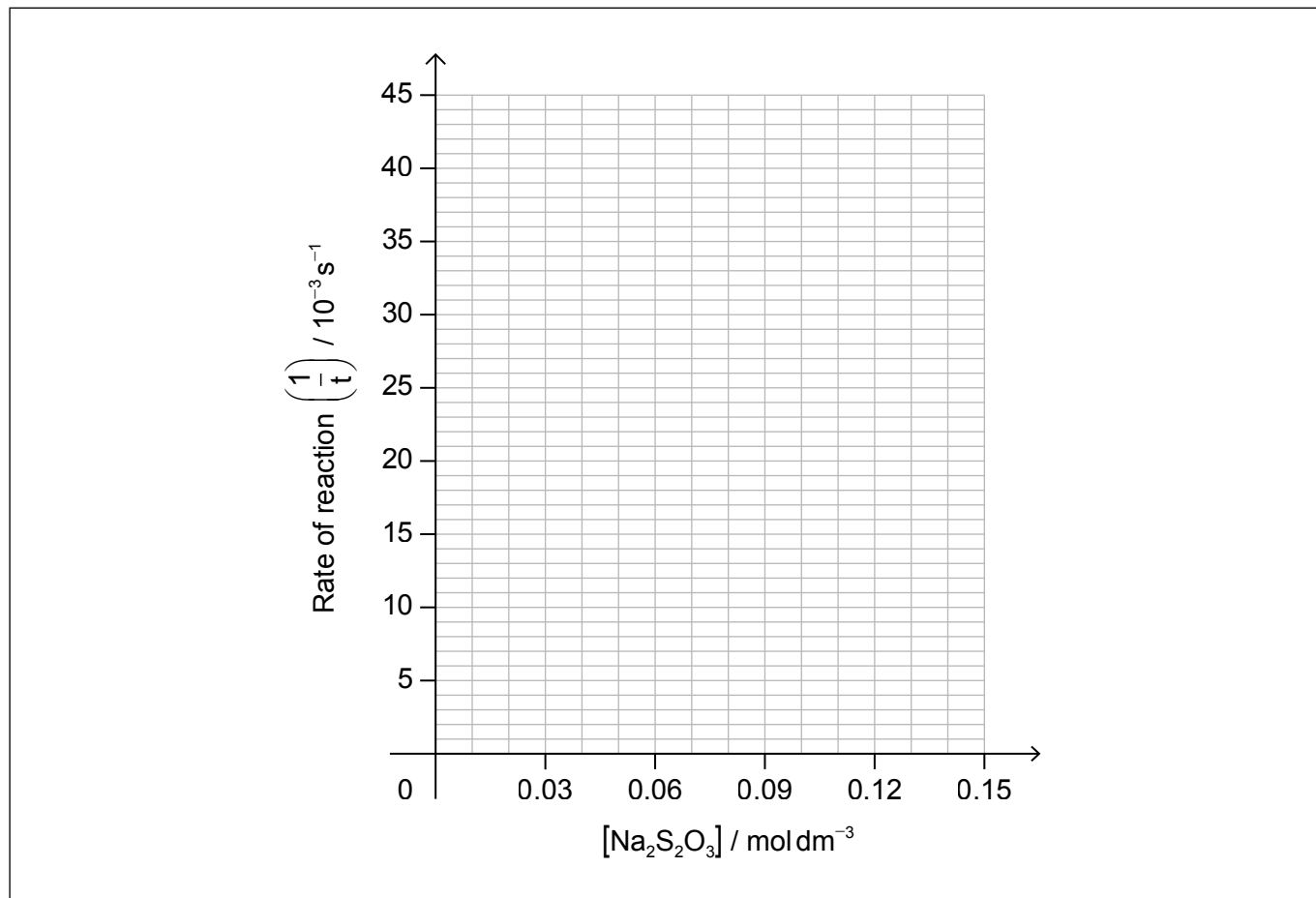
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(Question 3 continued)

- (d) Draw the best fit line of  $\frac{1}{t}$  against concentration of sodium thiosulfate on the axes provided. [2]



- (e) (i) Using the graph, explain the order of reaction with respect to sodium thiosulfate. [2]

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**(Question 3 continued)**

- (ii) In a different experiment, this reaction was found to be first order with respect to hydrochloric acid. Deduce the overall rate expression for the reaction. [1]

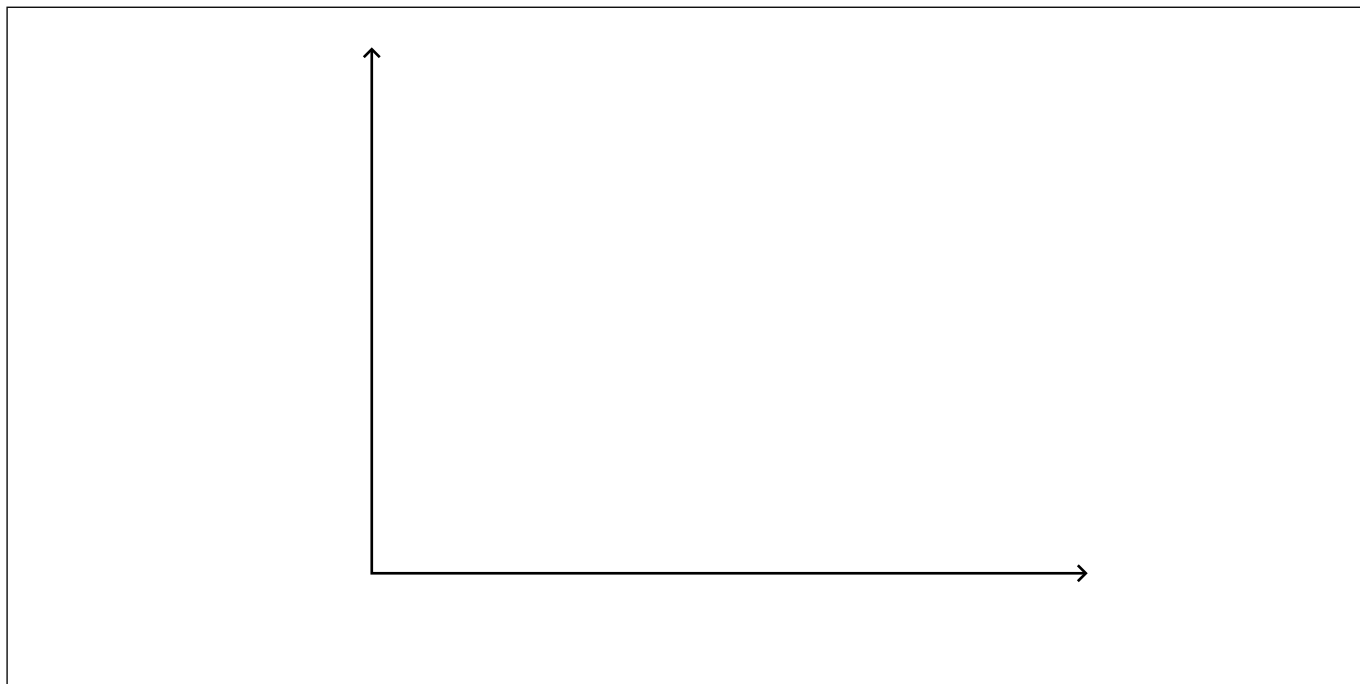
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- (f) A student decided to carry out another experiment using  $0.075 \text{ mol dm}^{-3}$  solution of sodium thiosulfate under the same conditions. Determine the time taken for the mark to be no longer visible. [2]

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- (g) An additional experiment was carried out at a higher temperature,  $T_2$ .

- (i) On the same axes, sketch Maxwell-Boltzmann energy distribution curves at the two temperatures  $T_1$  and  $T_2$ , where  $T_2 > T_1$ . [2]



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**(Question 3 continued)**

(ii) Explain why a higher temperature causes the rate of reaction to increase. [2]

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(h) Suggest one reason why the values of rates of reactions obtained at higher temperatures may be less accurate. [1]

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4. Magnesium is a group 2 metal which exists as a number of isotopes and forms many compounds.

(a) State the nuclear symbol notation,  ${}^A_ZX$ , for magnesium-26. [1]

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(b) Mass spectroscopic analysis of a sample of magnesium gave the following results:

	% abundance
Mg-24	78.60
Mg-25	10.11
Mg-26	11.29

Calculate the relative atomic mass,  $A_r$ , of this sample of magnesium to two decimal places. [2]

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(c) Magnesium ions produce no emission or absorption lines in the visible region of the electromagnetic spectrum. Suggest why most magnesium compounds tested in a school laboratory show traces of yellow in the flame. [1]

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**(Question 4 continued)**

(d) (i) Explain the convergence of lines in a hydrogen emission spectrum. [1]

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(ii) State what can be determined from the frequency of the convergence limit. [1]

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(e) Magnesium burns in air to form a white compound, magnesium oxide. Formulate an equation for the reaction of magnesium oxide with water. [1]

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(f) Describe the trend in acid-base properties of the oxides of period 3, sodium to chlorine. [2]

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(g) In addition to magnesium oxide, magnesium forms another compound when burned in air. Suggest the formula of this compound. [1]

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**(Question 4 continued)**

(h) Describe the structure and bonding in solid magnesium oxide. [2]

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(i) Magnesium chloride can be electrolysed.  
(i) Deduce the half-equations for the reactions at each electrode when **molten** magnesium chloride is electrolysed, showing the state symbols of the products. The melting points of magnesium and magnesium chloride are 922 K and 987 K respectively. [2]

Anode (positive electrode):  
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Cathode (negative electrode):  
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(ii) Identify the type of reaction occurring at the cathode (negative electrode). [1]

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(iii) State the products when a very **dilute** aqueous solution of magnesium chloride is electrolysed. [2]

Anode (positive electrode):  
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Cathode (negative electrode):  
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**(This question continues on the following page)**





**(Question 4 continued)**

- (j) Standard electrode potentials are measured relative to the standard hydrogen electrode. Describe a standard hydrogen electrode. [2]

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- (k) A magnesium half-cell,  $\text{Mg (s)/Mg}^{2+} \text{ (aq)}$ , can be connected to a copper half-cell,  $\text{Cu (s)/Cu}^{2+} \text{ (aq)}$ .

- (i) Formulate an equation for the spontaneous reaction that occurs when the circuit is completed. [1]

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- (ii) Determine the standard cell potential, in V, for the cell. Refer to section 24 of the data booklet. [1]

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- (iii) Predict, giving a reason, the change in cell potential when the concentration of copper ions increases. [2]

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5. Propane and propene are members of different homologous series.

(a) Draw the full structural formulas of propane and propene.

[1]

Propane:

Propene:

(b) (i) Draw diagrams to show how sigma ( $\sigma$ ) and pi ( $\pi$ ) bonds are formed between atoms.

[2]

Sigma ( $\sigma$ ):

Pi ( $\pi$ ):

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**(Question 5 continued)**

(ii) State the number of sigma ( $\sigma$ ) and pi ( $\pi$ ) bonds in propane and propene. [2]

	Number of sigma ( $\sigma$ ) bonds	Number of pi ( $\pi$ ) bonds
<b>Propane</b>	.....	.....
<b>Propene</b>	.....	.....

(c) Both propane and propene react with bromine.

(i) State an equation and the condition required for the reaction of 1 mol of propane with 1 mol of bromine. [2]

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(ii) State an equation for the reaction of 1 mol of propene with 1 mol of bromine. [1]

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(iii) State the type of each reaction with bromine. [1]

Propane:  
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Propene:  
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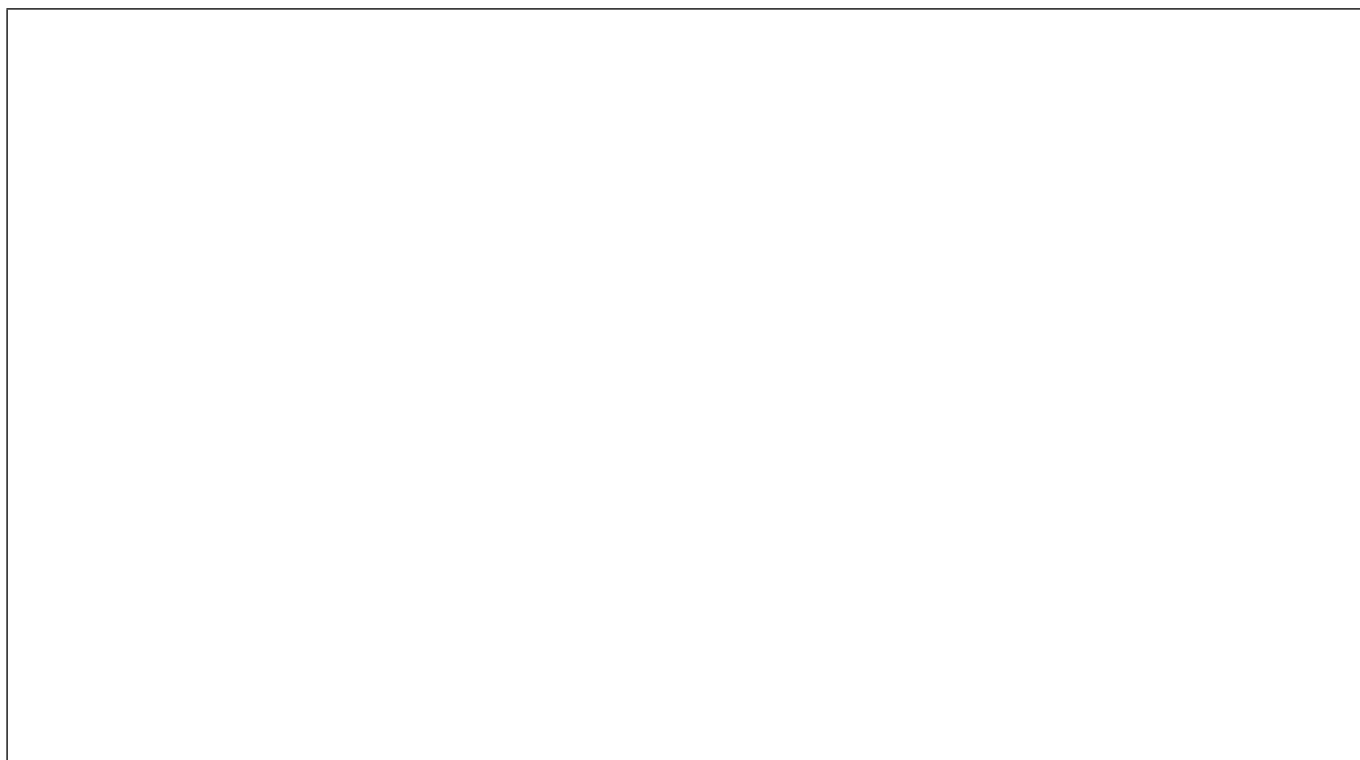
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**(Question 5 continued)**

- (d) Construct the mechanism of the formation of 2-bromopropane from hydrogen bromide and propene using curly arrows to denote the movement of electrons.

[3]



6. One structural isomer of  $C_4H_9Br$  is a chiral molecule.

(a) Draw the three-dimensional shape of each enantiomer of this isomer showing their spatial relationship to each other. [2]

(b) When one enantiomer undergoes substitution by alkaline hydrolysis approximately 75% of the product molecules show inversion of configuration. Comment on the mechanisms that occur. [2]

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(c) Suggest why the rate of alkaline hydrolysis of an enantiomer of iodopropane is greater than that of an enantiomer of bromopropane. [1]

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7. This question is about the weak acid methanoic acid, HCOOH.

- (a) Calculate the pH of  $0.0100 \text{ mol dm}^{-3}$  methanoic acid stating any assumption you make.  
 $K_a = 1.6 \times 10^{-4}$ . [3]

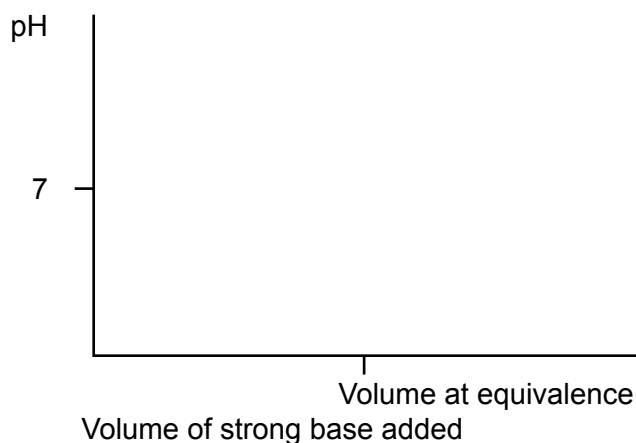
Calculation:

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Assumption:

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- (b) (i) Sketch a graph of pH against volume of a strong base added to a weak acid showing how you would determine  $pK_a$  for the weak acid. [2]



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**(Question 7 continued)**

- (ii) Explain, using an equation, why the pH increases very little in the buffer region when a small amount of alkali is added.

[2]

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Answers written on this page  
will not be marked.



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