## $A Q A D$

Please write clearly in block capitals.

Centre number |  |  |  |  |  |
| :--- | :--- | :--- | :--- | :--- |

Candidate number $\square$

Surname
Forename(s)
Candidate signature
I declare this is my own work.

## A-level

## CHEMISTRY

## Paper 1 Inorganic and Physical Chemistry

Monday 12 June 2023

## Materials

For this paper you must have:

- the Periodic Table/Data Booklet, provided as an insert (enclosed)
- a ruler with millimetre measurements
- a scientific calculator, which you are expected to use where appropriate.


## Instructions

- Use black ink or black ball-point pen.
- Fill in the boxes at the top of this page.
- Answer all questions.
- You must answer the questions in the spaces provided. Do not write outside the box around each page or on blank pages.
- If you need extra space for your answer(s), use the lined pages at the end of this book. Write the question number against your answer(s).
- All working must be shown.
- Do all rough work in this book. Cross through any work you do not want to be marked.


## Information

| For Examiner's Use |  |
| :---: | :---: |
| Question | Mark |
| 1 |  |
| 2 |  |
| 3 |  |
| 4 |  |
| 5 |  |
| 6 |  |
| 7 |  |
| 8 |  |
| 9 |  |
| 10 |  |
| TOTAL |  |

- The marks for questions are shown in brackets.
- The maximum mark for this paper is 105.


| $\mathbf{0}$ | $\mathbf{1}$ | $\mathbf{3}$ | Suggest why the oxidation state of chromium is zero in $\operatorname{Cr}\left(\mathrm{PF}_{3}\right)_{6}$ |
| :--- | :--- | :--- | :--- |

$\qquad$
$\qquad$
$\qquad$

|  |  |  | The compound $\left[\mathrm{Cr}\left(\mathrm{NH}_{3}\right)_{4} \mathrm{Cl} 2\right] \mathrm{Cl}$ contains ammonia molecules. | [1 mark] |
| :---: | :---: | :---: | :---: | :---: |
| 0 | D 1 . 4 |  | Deduce the oxidation state of chromium in $\left[\mathrm{Cr}\left(\mathrm{NH}_{3}\right)_{4} \mathrm{Cl} 2\right] \mathrm{Cl}$ |  |
|  |  |  |  |  |
|  | 0 0 1. 5 |  | Name the type of bond between N and H in ammonia. | [1 mark] |
|  |  |  |  |  |

$\qquad$

| $\mathbf{0}$ | $\mathbf{1}$. | 6 |
| :--- | :--- | :--- | The compound $\left[\mathrm{Cr}\left(\mathrm{NH}_{3}\right)_{4} \mathrm{Cl}_{2}\right] \mathrm{Cl}$ contains a complex ion that shows isomerism.

Draw the two isomers of the complex ion.
State the type of isomerism shown.

## Isomer 1

Isomer 2

Type of isomerism $\qquad$
$\begin{array}{llll}0 & 1 & \mathbf{7} & \text { Complete the equation to show the formation of one complex that contains chromium }\end{array}$ in its +3 oxidation state.
$\mathrm{CrCl}_{3}+5 \mathrm{H}_{2} \mathrm{O} \rightarrow$

| $\mathbf{0}$ | $\mathbf{2}$ | Figure 1 shows a cell used to measure the standard electrode potential for the |
| :--- | :--- | :--- | half-cell

$$
\mathrm{Mg}^{2+}(\mathrm{aq})+2 \mathrm{e}^{-} \rightarrow \mathrm{Mg}(\mathrm{~s})
$$

Figure 1


| $\mathbf{0}$ | $\mathbf{2}$. | $\mathbf{1}$ |
| :--- | :--- | :--- |

Identify an ionic compound that could be used in the salt bridge.

Purpose $\qquad$
$\qquad$
Identity $\qquad$
$\qquad$

| $\mathbf{0}$ | $\mathbf{2} .2$ | $\mathbf{2}$ State how, if at all, the EMF of this cell will change if the surface area of the platinum |
| :--- | :--- | :--- | electrode is increased.

$\qquad$
$\qquad$
The standard electrode potential, $E^{\ominus}$ for the half-cell is shown.
$\mathrm{Mg}^{2+}(\mathrm{aq})+2 \mathrm{e}^{-} \rightarrow \mathrm{Mg}(\mathrm{s}) \quad E^{\ominus}=-2.38 \mathrm{~V}$

| $\mathbf{0}$ | $\mathbf{2}$. | 3 |
| :--- | :--- | :--- | Water is added to the beaker containing the magnesium chloride solution.

What is the effect on the magnitude of the EMF of the cell?
Tick $(\checkmark)$ one box.

EMF increases


EMF stays the same


EMF decreases


Give an equation for the overall reaction that occurs when the cell is operating.
-

## Turn over for the next question

| $\mathbf{0}$ | $\mathbf{3} \quad$ This question is about Period 3 elements and their oxides. |
| :--- | :--- | :--- |


| $\mathbf{0}$ | $\mathbf{3}$. | $\mathbf{1}$ Give an equation for the reaction between phosphorus and an |
| :--- | :--- | :--- | excess of oxygen.



| $\mathbf{0}$ | $\mathbf{3}$. | $\mathbf{3}$ Give the displayed formula for the anion formed when sulfur trioxide reacts with water. |
| :--- | :--- | :--- | [1 mark]


| $\mathbf{0}$ | $\mathbf{3}$. | $\mathbf{4}$ Give an equation for the reaction of magnesium with steam. |
| :--- | :--- | :--- |

State one observation made.

Equation
$\qquad$
Observation $\qquad$
$\qquad$
$\begin{array}{llll}0 & 3 & 5 & \text { Give an equation to show how an excess of magnesium oxide reacts with }\end{array}$ phosphoric acid $\left(\mathrm{H}_{3} \mathrm{PO}_{4}\right)$.
$\qquad$

| 0 | 4 | Nitrogen dioxide decomposes at a high temperature. |
| :--- | :--- | :--- |

$$
2 \mathrm{NO}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{NO}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g}) \quad \Delta H=+113 \mathrm{~kJ} \mathrm{~mol}^{-1}
$$

| 0 | 4 | 1 |
| :--- | :--- | :--- | constant temperature until equilibrium is reached.

At equilibrium, the flask contains 0.120 mol of oxygen.
Calculate the mole fraction of each substance at equilibrium.
[3 marks]

Mole fraction of $\mathrm{NO}_{2}$
Mole fraction of NO $\qquad$
Mole fraction of $\mathrm{O}_{2}$ $\qquad$
$\begin{array}{lll}0 & 4 & 2\end{array}$ The total pressure in the flask in Question 04.1 is 120 kPa at equilibrium.
Calculate the partial pressure, in kPa , of $\mathrm{NO}_{2}$
If you were unable to answer Question $\mathbf{0 4 . 1}$ you should assume that the mole fraction of $\mathrm{NO}_{2}$ is 0.380 . This is not the correct answer.
$\qquad$ kPa


$$
2 \mathrm{NO}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{NO}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g}) \quad \Delta H=+113 \mathrm{~kJ} \mathrm{~mol}^{-1}
$$

Table 1

| Gas | Mole fraction |
| :---: | :---: |
| $\mathrm{NO}_{2}$ | 0.310 |
| NO | 0.460 |
| $\mathrm{O}_{2}$ | 0.230 |

For this equilibrium mixture, $K_{\mathrm{p}}=59.7 \mathrm{kPa}$
Give an expression for $K_{\mathrm{p}}$ for this reaction.
Use your expression and the data in Table 1 to calculate the total pressure, in kPa , in the flask.
$K_{\text {p }}$
$\qquad$ kPa

| 0 | 4 | 4 |
| :--- | :--- | :--- |
| 4 |  |  | The equilibrium mixture in Question 04.3 is compressed into a smaller volume.

Deduce the effect, if any, of this change on the equilibrium yield of oxygen and on the value of $K_{p}$

Effect on yield of oxygen $\qquad$
Effect on $K_{p}$ $\qquad$
$\begin{array}{llll}0 & 4 & 5 & \text { The equilibrium mixture in Question } 04.3 \text { is allowed to reach equilibrium at a lower }\end{array}$ temperature.

Explain why the equilibrium yield of oxygen decreases.
$\qquad$
$\qquad$
$\qquad$

| $\mathbf{0}$ | $\mathbf{5}$ This question is about metal chlorides. |
| :--- | :--- | :--- |


| 0 | 5 |
| :--- | :--- | l Table 2 shows some enthalpy change data.

## Table 2

|  | Enthalpy change $/$ kJ mol $^{-1}$ |
| :--- | :---: |
| $\mathrm{Ca}^{2+}(\mathrm{g}) \rightarrow \mathrm{Ca}^{2+}(\mathrm{aq})$ | -1650 |
| $\mathrm{Cl}^{-}(\mathrm{g}) \rightarrow \mathrm{Cl}^{-}(\mathrm{aq})$ | -364 |
| $\mathrm{Ca}^{2+}(\mathrm{g})+2 \mathrm{Cl}^{-}(\mathrm{g}) \rightarrow \mathrm{CaCl}_{2}(\mathrm{~s})$ | -2237 |

Use the data in Table 2 to calculate the molar enthalpy change when calcium chloride dissolves in water.

Molar enthalpy change $\qquad$ $\mathrm{kJ} \mathrm{mol}{ }^{-1}$

| 0 | 5 | 2 |
| :--- | :--- | :--- | Use your answer to Question 05.1 to deduce how the temperature changes when calcium chloride dissolves in water.

$\qquad$
$\qquad$

| $\mathbf{0}$ | $\mathbf{5}$. | 3 |
| :--- | :--- | :--- | enthalpy of hydration of chloride ions.

$\qquad$
$\qquad$
$\qquad$
$\qquad$

| $\mathbf{0}$ | $\mathbf{5}$ | $\mathbf{4}$ | Figure $\mathbf{2}$ shows an incomplete Born-Haber cycle for calcium chloride. |
| :--- | :--- | :--- | :--- |

Figure 2


Complete the Born-Haber cycle by writing the formulas of the missing species on each of the two blank lines.

| 0 | 5 | 5 |
| :--- | :--- | :--- |

Table 3

|  | Enthalpy change $/ \mathbf{~ k ~}^{2}$ mol $^{-1}$ |
| :--- | :---: |
| Enthalpy of atomisation of calcium | +193 |
| First ionisation energy of calcium | +590 |
| Enthalpy of atomisation of chlorine | +121 |
| Electron affinity of chlorine | -364 |
| Enthalpy of formation of calcium chloride | -795 |
| Enthalpy of lattice formation of calcium chloride | -2237 |

Use Figure 2 and data from Table 3 to calculate the second ionisation energy of calcium.

| 0 | 5 | 6 |
| :--- | :--- | :--- | first ionisation energy of calcium.

$\qquad$
$\qquad$

| $\mathbf{0}$ | $\mathbf{5} .7$ | Table 4 shows lattice enthalpies based on a perfect ionic model and lattice enthalpies |
| :--- | :--- | :--- | :--- | from Born-Haber cycles for three metal chlorides.

Table 4

|  | Lattice enthalpy of dissociation $/$ kJ mol $^{-1}$ |  |
| :--- | :---: | :---: |
|  | Perfect ionic model | Born-Haber cycle |
| Calcium chloride | 2223 | 2237 |
| Potassium chloride | 690 | 701 |
| Silver chloride | 770 | 905 |

Discuss the values in Table 4.
In your answer you should

- compare the three values based on a perfect ionic model
- compare the values based on a perfect ionic model to the values from a Born-Haber cycle for each compound.
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$


| 0 | 6 |
| :--- | :--- | The concentration of dilute hydrochloric acid can be found by titration using a standard solution of barium hydroxide.


| 0 | 6 | 1 |
| :--- | :--- | :--- |
| 1 | Calculate the mass, in g , of solid barium hydroxide $\left(M_{\mathrm{r}}=171.3\right)$ needed to prepare |  | $250 \mathrm{~cm}^{3}$ of $0.100 \mathrm{~mol} \mathrm{dm}^{-3}$ barium hydroxide solution.


| 0 | 6 | 2 | $T h e ~ m a s s ~ o f ~ b a r i u m ~ h y d r o x i d e ~ f r o m ~ Q u e s t i o n ~$ |
| :--- | :--- | :--- | :--- |
| 06.1 |  |  |  | $150 \mathrm{~cm}^{3}$ of distilled water.

Describe how this solution is used to make $250 \mathrm{~cm}^{3}$ of the $0.100 \mathrm{~mol} \mathrm{dm}^{-3}$ barium hydroxide solution.
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$

06 . 3 Before the first titration, the $25 \mathrm{~cm}^{3}$ pipette is rinsed with a small volume of the $0.100 \mathrm{~mol} \mathrm{dm}^{-3}$ barium hydroxide solution.

State why it is good practice to rinse the pipette in this way.
$\qquad$
$\qquad$
$\qquad$

| $\mathbf{0}$ | $\mathbf{6}$. | $\mathbf{4}$ | Hydrochloric acid is added to the burette using a funnel. |
| :--- | :--- | :--- | :--- |

State why it is good practice to remove the funnel from the burette before the titration.
[1 mark]
$\qquad$
$\qquad$
$\qquad$

| 0 | 6 | 5 |
| :--- | :--- | :--- | of standard barium hydroxide solution.

$25.0 \mathrm{~cm}^{3}$ of this barium hydroxide solution reacts with exactly
$24.50 \mathrm{~cm}^{3}$ of hydrochloric acid.
Calculate the concentration of the hydrochloric acid.
$\qquad$ $\mathrm{mol} \mathrm{dm}^{-3}$
 The total uncertainty in the $24.50 \mathrm{~cm}^{3}$ of solution from the burette is $\pm 0.15 \mathrm{~cm}^{3}$

Calculate the total percentage error in using the pipette and burette.

| $\mathbf{0}$ | $\mathbf{7}$ | This question is about complexes containing the aluminium ion. |
| :--- | :--- | :--- |


| $\mathbf{0}$ | $\mathbf{7}$ | $\mathbf{1}$ Give the electron configuration of the $\mathrm{Al}^{3+}$ ion. |
| :--- | :--- | :--- |


| $\mathbf{0}$ | $\mathbf{7}$. | $\mathbf{2}$ When anhydrous aluminium sulfate, $\mathrm{Al}_{2}\left(\mathrm{SO}_{4}\right)_{3}$, is added to water a solution forms that |
| :--- | :--- | :--- | contains the complex aluminium ion, $\left[\mathrm{Al}\left(\mathrm{H}_{2} \mathrm{O}\right)_{6}\right]^{3+}$

Give the equation for the reaction.

| $\mathbf{0}$ | $\mathbf{7}$ | $\mathbf{3}$ Explain why the solution containing $\left[\mathrm{Al}\left(\mathrm{H}_{2} \mathrm{O}\right)_{6}\right]^{3+}$ is acidic. |
| :--- | :--- | :--- | :--- |

$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$

| $\mathbf{0}$ | $\mathbf{7} .4$ | State why the concentration of aluminium sulfate solution can not be determined by |
| :--- | :--- | :--- | :--- | colorimetry.

$\qquad$
$\qquad$
$\qquad$

| $\mathbf{0}$ | $\mathbf{7}$ | $\mathbf{5}$ | An excess of aqueous ammonia is added to a solution containing $\left[\mathrm{Al}\left(\mathrm{H}_{2} \mathrm{O}\right)_{6}\right]^{3+}$ |
| :--- | :--- | :--- | :--- |

Give an ionic equation for the reaction and state one observation.

Equation
$\qquad$

Observation $\qquad$

| 0 | $\mathbf{7} .6$ | An excess of dilute sulfuric acid is added to the products of the reaction in |
| :--- | :--- | :--- | :--- | Question 07.5

Identify the aluminium species produced.

| 0 | $\mathbf{7}$. | $\mathbf{7}$ |
| :--- | :--- | :--- | Figure $\mathbf{3}$ shows the structure of the EDTA ${ }^{4-}$ ion.

Figure 3


Atoms of two different elements in EDTA4- can form co-ordinate bonds with an aluminium ion.

On Figure 3, draw circles around the atoms of two different elements that would link to an aluminium ion by a co-ordinate bond.

| $\mathbf{0}$ | $\mathbf{7}$ | $\mathbf{8}$ | Hydrated aluminium sulfate, $\mathrm{Al}_{2}\left(\mathrm{SO}_{4}\right)_{3} \cdot \mathrm{xH}_{2} \mathrm{O}$, is soluble in water. |
| :--- | :--- | :--- | :--- |

The relative formula mass and value of $x$ can be found from a titration experiment.
Aqueous $\left[\mathrm{Al}\left(\mathrm{H}_{2} \mathrm{O}\right)_{6}\right]^{3+}$ ions react to form a stable complex when treated with an excess of EDTA ${ }^{4-}$ ions.

The excess of EDTA ${ }^{4-}$ ions is determined by titration with $\mathrm{ZnSO}_{4}$ solution.
Method

- Dissolve 1.036 g of $\mathrm{Al}_{2}\left(\mathrm{SO}_{4}\right)_{3} \cdot \mathrm{xH}_{2} \mathrm{O}$ in distilled water and make up to $250 \mathrm{~cm}^{3}$
- Add $25.0 \mathrm{~cm}^{3}$ of this solution to $50.0 \mathrm{~cm}^{3}$ of a solution containing EDTA ${ }^{4-}$ ions of concentration $0.0100 \mathrm{~mol} \mathrm{dm}^{-3}$
- Determine the excess of EDTA4- ions by titrating with $\mathrm{ZnSO}_{4}$ solution in the presence of an indicator.

The excess of EDTA ${ }^{4-}$ ions requires $18.00 \mathrm{~cm}^{3}$ of $0.0105 \mathrm{~mol} \mathrm{dm}^{-3} \mathrm{ZnSO}_{4}$ solution to react completely.

The equations for the reactions are

$$
\begin{aligned}
{\left[\mathrm{Al}\left(\mathrm{H}_{2} \mathrm{O}\right)_{6}\right]^{3+}+\mathrm{EDTA}^{4-} } & \rightarrow[\text { AlEDTA }]^{-}+6 \mathrm{H}_{2} \mathrm{O} \\
{\left[\mathrm{Zn}\left(\mathrm{H}_{2} \mathrm{O}\right)_{6}\right]^{2+}+\mathrm{EDTA}^{4-} } & \rightarrow[\mathrm{ZnEDTA}]^{2-}+6 \mathrm{H}_{2} \mathrm{O}
\end{aligned}
$$

For $\mathrm{Al}_{2}\left(\mathrm{SO}_{4}\right)_{3} \quad M_{\mathrm{r}}=342.3$

Use the information given to calculate the $M_{\mathrm{r}}$ of $\mathrm{Al}_{2}\left(\mathrm{SO}_{4}\right)_{3} \cdot \mathrm{XH}_{2} \mathrm{O}$
Calculate $x$
Give your answer as an integer.
Do not write outside the box
$\qquad$


| $\mathbf{0}$ | 8 | This question is about fuel cells. |
| :--- | :--- | :--- |

In a methanol-oxygen fuel cell, the overall reaction is

$$
\mathrm{CH}_{3} \mathrm{OH}(\mathrm{I})+1 \frac{1}{2} \mathrm{O}_{2}(\mathrm{~g}) \rightarrow \mathrm{CO}_{2}(\mathrm{~g})+2 \mathrm{H}_{2} \mathrm{O}(\mathrm{I}) \quad \mathrm{EMF}=+1.20 \mathrm{~V}
$$

| 0 | 8 | . | $\mathbf{1}$ |
| :--- | :--- | :--- | :--- | At the positive electrode, oxygen reacts with hydrogen ions to form water.

Give a half-equation for this reaction.

| $\mathbf{0}$ | $\mathbf{8}$. | $\mathbf{2}$ At the negative electrode, methanol reacts with water to produce |
| :--- | :--- | :--- | carbon dioxide and hydrogen ions.

Give a half-equation for this reaction.

| $\mathbf{0}$ | $\mathbf{8} .3$ The standard electrode potential for the $\mathrm{CO}_{2} / \mathrm{CH}_{3} \mathrm{OH}$ electrode is +0.03 V |
| :--- | :--- | :--- |

Calculate the standard electrode potential for the $\mathrm{O}_{2} / \mathrm{H}_{2} \mathrm{O}$ electrode.
$\qquad$

| 0 | 8 | 4 | State why a fuel cell does not need to be electrically recharged. |
| :--- | :--- | :--- | :--- |

$\qquad$
$\qquad$

| 0 | 8 | 5 |
| :--- | :--- | :--- |
| 5 |  |  | in cars.

$\qquad$
$\qquad$
$\qquad$

| 0 | 9 |
| :--- | :--- | This is a question about time of flight (TOF) mass spectrometry.


| $\mathbf{0}$ | $\mathbf{9} .1$ | Give the equation, including state symbols, for the formation of $\mathrm{Sr}^{+}$ions from |
| :--- | :--- | :--- | Sr atoms by electron impact.


| $\mathbf{0}$ | $\mathbf{9}$. | $\mathbf{2}$ A sample of strontium is analysed by TOF mass spectrometry. |
| :--- | :--- | :--- |

The sample is ionised using electron impact.
The ions are accelerated to have a kinetic energy (KE) of $7.02 \times 10^{-20} \mathrm{~J}$ An ion takes $9.47 \times 10^{-4} \mathrm{~s}$ to travel along a 95.0 cm flight tube.

$$
K E=\frac{1}{2} m v^{2}
$$

where $m=\operatorname{mass}(\mathrm{kg})$ and $v=$ speed $\left(\mathrm{m} \mathrm{s}^{-1}\right)$
Use the information given to deduce the mass number of this ion.
The Avogadro constant, $L=6.022 \times 10^{23} \mathrm{~mol}^{-1}$
$\qquad$

| 0 | $\mathbf{9}$ | $\mathbf{3}$ Explain how the ions are detected in the TOF mass spectrometer. ${ }^{2}$. |
| :--- | :--- | :--- |

State how the relative abundance of the ions is determined.

How ions are detected $\qquad$
$\qquad$
How relative abundance is determined $\qquad$
$\qquad$
$\qquad$

| 0 | 9 | .4 |
| :--- | :--- | :--- | A sample of strontium contains three isotopes, ${ }^{86} \mathrm{Sr},{ }^{87} \mathrm{Sr}$ and ${ }^{88} \mathrm{Sr}$ $82 \%$ of the sample is ${ }^{88} \mathrm{Sr}$ The other isotopes are in a $1: 2$ ratio of ${ }^{86} \mathrm{Sr}:{ }^{87} \mathrm{Sr}$

Calculate the percentage abundance of ${ }^{87} \mathrm{Sr}$ in this sample.
Use your answer to deduce the relative atomic mass $\left(A_{\mathrm{r}}\right)$ of the sample.
Give your answer to 1 decimal place.

Abundance of ${ }^{87} \mathrm{Sr}$ \%
$A_{r}$ $\qquad$

| $\mathbf{0}$ | $\mathbf{9}$ | $\mathbf{5}$ Electrospray ionisation is used instead of electron impact for the ionisation of a protein |
| :--- | :--- | :--- | :--- | in a mass spectrometry experiment.

Suggest why.
$\qquad$
$\qquad$

| $\mathbf{1}$ | $\mathbf{0}$ | This question is about weak acids. |
| :--- | :--- | :--- |


| 1 | 0 | 1 |
| :--- | :--- | :--- |

## Table 5

| Indicator | pH range |
| :--- | :---: |
| Bromocresol green | $3.8-5.4$ |
| Bromothymol blue | $6.0-7.6$ |
| Thymol blue | $8.0-9.6$ |

Identify the indicator that is most suitable for use in a titration between propanoic acid and sodium hydroxide.
$\qquad$

| $\mathbf{1}$ | $\mathbf{0} .2$ | $\mathbf{2}$ Give the expression for the acid dissociation constant $\left(K_{\mathrm{a}}\right)$ for propanoic acid |
| :--- | :--- | :--- | $\left(\mathrm{CH}_{3} \mathrm{CH}_{2} \mathrm{COOH}\right)$.

$K_{a}$

Give your answer to 2 decimal places.
For propanoic acid, $\mathrm{p} K_{\mathrm{a}}=4.87$
$\qquad$

| 1 | $\mathbf{0}$. | 4 | For butanoic acid, $K_{a}=1.51 \times 10^{-5} \mathrm{~mol} \mathrm{dm}^{-3}$ |
| :--- | :--- | :--- | :--- |

$20.0 \mathrm{~cm}^{3}$ of $0.100 \mathrm{~mol} \mathrm{dm}^{-3}$ sodium hydroxide solution are added to $25.0 \mathrm{~cm}^{3}$ of $0.100 \mathrm{~mol} \mathrm{dm}^{-3}$ butanoic acid solution.

Calculate the pH of the solution formed.

| 1 | 0 | 5 |
| :--- | :--- | :--- |
| 5 |  |  | A student plans to titrate butanoic acid solution with a solution of ethylamine.

Explain why this titration could not be done using an indicator.
$\qquad$
$\qquad$
$\qquad$




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