## CHEMISTRY

## ATAR course examination 2018

## Marking Key

Marking keys are an explicit statement about what the examining panel expect of candidates when they respond to particular examination items. They help ensure a consistent interpretation of the criteria that guide the awarding of marks.

| Question No. | Answer |
| :---: | :---: |
| 1 | c |
| 2 | d |
| 3 | b |
| 4 | a |
| 5 | a |
| 6 | d |
| 7 | d |
| 8 | b |
| 9 | c |
| 10 | All |
| 11 | b |
| 12 | c |
| 13 | d |
| 14 | c |
| 15 | a |
| 16 | c |
| 17 | c |
| 18 | d |
| 19 | b |
| 20 | c |
| 21 | b |
| 22 | d |
| 23 | a |
| 24 | a |
| 25 | a |
|  |  |

* Question 10 - all distractors were deemed correct; all candidates were awarded 1 mark.


## General Notes:

- Answers provided in brackets are desired but not essential for the relevant mark to be allocated.
- Clear and unambiguous demonstration of the candidate's understanding in their response is required rather than the exact wording in the marking key.
- Notes provided within the marking key will indicate which elements of the model answers are required for the allocated mark.


## Question 26

Solid copper(II) hydroxide is added to excess $0.100 \mathrm{~mol} \mathrm{~L}^{-1}$ carbonic acid solution.
(a) Write the balanced equation, with appropriate state symbols, for the reaction that takes place between the copper(II) hydroxide and carbonic acid.
(3 marks)

| Description |  | Marks |
| :---: | :---: | :---: |
| $\begin{aligned} & \mathrm{Cu}(\mathrm{OH})_{2}(\mathrm{~s})+\mathrm{H}_{2} \mathrm{CO}_{3}(\mathrm{aq}) \rightarrow \mathrm{CuCO}_{3}(\mathrm{~s})+2 \mathrm{H}_{2} \mathrm{O}(\ell) \\ & \text { or } \\ & \mathrm{Cu}(\mathrm{OH})_{2}(\mathrm{~s})+2 \mathrm{H}_{2} \mathrm{CO}_{3}(\mathrm{aq}) \rightarrow \mathrm{CuCO}_{3}(\mathrm{~s})+\mathrm{CO}_{2}(\mathrm{~g})+3 \mathrm{H}_{2} \mathrm{O}(\ell) \end{aligned}$ |  |  |
| correct species |  | 1 |
| balancing |  | 1 |
| state symbols |  | 1 |
|  | Total | 3 |

(b) Predict all visible changes that would be observed, if any, while the reactants are mixed together and afterwards.
(3 marks)

| Description | Marks |
| :--- | :---: |
| Any three of the following: <br> - the blue solid disappears (dissolves/reacts) <br> - a green precipitate/solid is produced <br> - once the reaction is complete the solution is colourless <br> effervescence or bubbles formed |  |
|  | $1-3$ |
|  | Total |

Note: Accept colourless, odourless gas instead of effervescence
Note: candidate must provide a minimum of three correct observations for full marks Note: do not accept 'clear' without reference to colour
(c) Predict two observations that would be different if excess $0.100 \mathrm{~mol} \mathrm{~L}^{-1}$ hydrochloric acid was used instead of the $0.100 \mathrm{~mol} \mathrm{~L}^{-1}$ carbonic acid.

| Description |  | Marks |
| :---: | :---: | :---: |
| $\left(\mathrm{Cu}(\mathrm{OH})_{2}(\mathrm{~s})+2 \mathrm{HCl}(\mathrm{aq}) \rightarrow \mathrm{CuCl}_{2}(\mathrm{aq})+2 \mathrm{H}_{2} \mathrm{O}(\ell)\right)$ <br> Any two of the following: <br> - no solid at all is produced <br> - the solution would be blue once the reaction was complete <br> - the reaction would occur faster <br> - no gas produced |  | 1-2 |
|  | Total | 2 |

## Question 26 (continued)

(d) State two personal safety measures the experimenter should take when conducting these experiments.

| Description | Marks |
| :--- | :---: |
| Any two of the following: |  |
| - wear safety glasses |  |
| - wear closed in shoes |  |
| - wear a laboratory coat |  |
| - tie back long hair |  |
| - wear a hair net |  |
| - avoid coming in contact with the chemicals (e.g. wearing gloves) |  |
| - avoid breathing any vapour |  |
| - read materials data sheet |  |
| Accept other relevant answers |  |
| Note: Do not accept PPE |  |
| Note: Do not accept actions taken in response to an accident | $\mathbf{2}$ |

## Question 27

(a) Write the ionisation equation for phosphoric acid in water which shows the second proton of the acid being released into solution.

| Description |  | Marks |
| :---: | :---: | :---: |
| $\mathrm{H}_{2} \mathrm{PO}_{4}^{-}(\mathrm{aq})+\mathrm{H}_{2} \mathrm{O}(\ell) \rightleftharpoons \mathrm{HPO}_{4}^{2-}(\mathrm{aq})+\mathrm{H}_{3} \mathrm{O}^{+}(\mathrm{aq})$ (one minor error) |  | $\begin{gathered} \hline 2 \\ (1) \\ \hline \end{gathered}$ |
|  | Total | 2 |

Note: State symbols not required
Note: Minor errors could include no double arrow, one charge missing
(b) Other than water, list three species (elements, compounds, ions) that would be found in the reacting vessel open to the atmosphere at the completion of the reaction between excess solid magnesium carbonate and an aqueous solution of phosphoric acid.
(3 marks)

| Description | Marks |
| :---: | :---: |
| Only the first three species listed are marked. <br> 1. magnesium carbonate, $\mathrm{MgCO}_{3}$ (s) <br> 2. magnesium phosphate, $\mathrm{Mg}_{3}\left(\mathrm{PO}_{4}\right)_{2}(\mathrm{~s})$ <br> 3. $\mathrm{H}_{2} \mathrm{CO}_{3}(\mathrm{aq})$ (from the reaction of $\mathrm{CO}_{2}$ with $\mathrm{H}_{2} \mathrm{O}$ ) <br> 4. $\mathrm{HCO}_{3}^{-}(\mathrm{aq})$ (from the reaction of $\mathrm{CO}_{2}$ with $\mathrm{H}_{2} \mathrm{O}$ ) <br> 5. $\quad \mathrm{CO}_{3}{ }^{2-}(\mathrm{aq})$ (from the reaction of $\mathrm{CO}_{2}$ with $\mathrm{H}_{2} \mathrm{O}$ ) <br> 6. $\mathrm{H}_{3} \mathrm{O}^{+}(\mathrm{aq})$ ions (from the self-ionisation of water) <br> 7. $\mathrm{OH}^{-}$(from the self-ionisation of water) <br> 8. $\mathrm{CO}_{2}$ <br> 9. $\mathrm{Mg}^{2+} / \mathrm{PO}_{4}{ }^{3-} / \mathrm{HPO}_{4}{ }^{2-} / \mathrm{H}_{2} \mathrm{PO}_{4}-/ \mathrm{O}_{2}$ | 1-3 |
|  |  |
|  |  |
|  |  |
|  |  |
|  |  |
|  |  |
|  |  |
|  |  |
|  |  |
|  | 3 |
| Note: the equation for the reaction is: <br> $3 \mathrm{MgCO}_{3}(\mathrm{~s})+2 \mathrm{H}_{3} \mathrm{PO}_{4}(\mathrm{aq}) \rightarrow \mathrm{Mg}_{3}\left(\mathrm{PO}_{4}\right)_{2}(\mathrm{~s})+3 \mathrm{H}_{2} \mathrm{O}(\ell)+3 \mathrm{CO}_{2}(\mathrm{~g})$ <br> $\mathrm{MgCO}_{3}$ is in excess so there is no $\mathrm{H}_{3} \mathrm{PO}_{4}$ left <br> A small amount of carbon dioxide gas, $\mathrm{CO}_{2}(\mathrm{~g})$, will dissolve and react with water, $\mathrm{H}_{2} \mathrm{O}$, to form $\mathrm{H}_{2} \mathrm{CO}_{3}$ (which will form $\mathrm{HCO}_{3}^{-}(\mathrm{aq})$ and $\mathrm{CO}_{3}{ }^{2-}(\mathrm{aq})$ ions). <br> Note: Only mark the first three listed |  |
|  |  |  |
|  |  |  |
|  |  |  |
|  |  |  |

## Question 27 (continued)

(c) Other than it having too low a molar mass, state two reasons why the concentration of the sodium hydroxide solution cannot be reliably determined by weighing out an amount of solid sodium hydroxide and dissolving it in a known volume of distilled water?
(2 marks)

| Description | Marks |
| :---: | :---: |
| Any two of the following: <br> - hygroscopic - it absorbs atmospheric moisture which changes the mass of the NaOH sample. <br> - deliquescent - it dissolves in the water it absorbs, so no mass of solid can be weighed <br> - absorbs atmospheric carbon dioxide gas, $\mathrm{CO}_{2}(\mathrm{~g})$ to form sodium carbonate, $\mathrm{Na}_{2} \mathrm{CO}_{3}(\mathrm{~s})$, and water, $\mathrm{H}_{2} \mathrm{O}(\ell)$. <br> - Not known purity (\% purity not known) | 1-2 |
| Total | 2 |

(d) Select the acid-base indicator from the table above that would be most suitable for the titration between phosphoric acid, $\mathrm{H}_{3} \mathrm{PO}_{4}(\mathrm{aq})$, and sodium hydroxide solution, $\mathrm{NaOH}(\mathrm{aq})$. Justify your choice of indicator, including one relevant equation.

| Description | Marks |
| :---: | :---: |
| Phenolphthalein | 1-5 |
| Recognition that $\mathrm{PO}_{4}{ }^{3-}$ present in solution at equivalence point. $\left(3 \mathrm{OH}^{-}(\mathrm{aq})+\mathrm{H}_{3} \mathrm{PO}_{4}(\mathrm{aq}) \rightarrow \mathrm{PO}_{4}{ }^{3-}(\mathrm{aq})+3 \mathrm{H}_{2} \mathrm{O}(\ell)\right)$ |  |
| The solution at the equivalence point will be (slightly) basic (with a pH of approximately 9 ) due to the excess of hydroxide ions |  |
| The phosphate ion undergoes hydrolysis to form hydroxide ions. $\mathrm{PO}_{4}{ }^{3-}(\mathrm{aq})+\mathrm{H}_{2} \mathrm{O}(\mathrm{l}) \rightleftharpoons \mathrm{HPO}_{4}{ }^{2-}(\mathrm{aq})+\mathrm{OH}^{-}(\mathrm{aq})$ |  |
| The pH at which the indicator changes colour approximates the pH of the equivalence point |  |
| The $\mathrm{Na}^{+}$ion does not hydrolyse/has no effect on the pH of the solution/is a neutral ion |  |
| Total | 5 |

Note: candidate must give five valid points for full marks, and must include an equation.
Note: no hydrolysis equation, maximum of four marks
Note: If phenol red with explanation above, four maximum
Alternative responses that candidate may provide:
Methyl Orange
The pH of the first equivalence point is around 4.7. If candidate identify this and supplied appropriate logic with equation, up to full marks may be awarded.

If a candidate recognises that the third equivalence point is beyond the end point of phenolphthalein and explains why none of the indicators would be appropriate with sufficient reasoning, up to full marks may be awarded.

## Question 28

(a) Write the half equation to show the reaction at the electrode at which oxidation occurs.

| Description | Marks |
| :--- | :---: |
| $\mathrm{H}_{2}+\mathrm{CO}_{3}{ }^{2-} \rightarrow \mathrm{CO}_{2}+\mathrm{H}_{2} \mathrm{O}+2 \mathrm{e}^{-}$ |  |
| correct equation | 3 |
| correct species but not balanced | 2 |
| Balanced for incorrect species (equation must have merit) | 1 |
|  | $\mathbf{3}$ |

(b) Write the overall equation for the current-producing reaction.

| Description | Marks |  |
| :--- | :---: | :---: |
| $\mathrm{O}_{2}+2 \mathrm{H}^{+}+2 \mathrm{H}_{2}+\mathrm{CO}_{3}{ }^{2-} \rightarrow 3 \mathrm{H}_{2} \mathrm{O}+\mathrm{CO}_{2}$ |  |  |
| or |  |  |
| $2 \mathrm{H}_{2}+\mathrm{O}_{2} \rightarrow 2 \mathrm{H}_{2} \mathrm{O}$ |  |  |
| correct equation | 3 |  |
| correct species but not balanced | 2 |  |
| Balanced for incorrect species (equation must have merit) | 1 |  |
| $\mathbf{3}$ |  |  |

(c) State two reasons why this fuel cell is a more environmentally-friendly alternative to the internal combustion engine.
(2 marks)

| Description | Marks |
| :---: | :---: |
| Any two statements giving an environmental advantage of the fuel cell: <br> - The carbon dioxide gas is recycled so that it doesn't enter the atmosphere (like the waste gases produced by an internal combustion engine) <br> - It uses $\left(\mathrm{O}_{2}\right.$ and $\left.\mathrm{CO}_{2}\right)$ as its fuel sources these are all renewable can be obtained from the atmosphere (whereas the internal combustion engine uses non-renewable fossil fuels.) <br> - In addition to the production of carbon dioxide gas, the by-products of the: <br> combustion engine include pollutant gases including the oxide gases of nitrogen and sulfur. These are significant contributors to acid rain. <br> fuel cell is water. <br> - The process of obtaining (drilling for oil) the fuel for the combustion engine damages the surrounding environments and the refining process (fractional distillation) for this fuel produces its own atmospheric pollutants. | 1-2 |
| Total | 2 |

## Question 29

(a) Write an equation for the buffer system $\left(\mathrm{H}_{3} \mathrm{C}_{6} \mathrm{H}_{5} \mathrm{O}_{7} / \mathrm{H}_{2} \mathrm{C}_{6} \mathrm{H}_{5} \mathrm{O}_{7}^{-}\right)$containing citric acid, $\mathrm{H}_{3} \mathrm{C}_{6} \mathrm{H}_{5} \mathrm{O}_{7}$ and monosodium citrate, $\mathrm{NaH}_{2} \mathrm{C}_{6} \mathrm{H}_{5} \mathrm{O}_{7}$.

| Description |  | Marks |
| :---: | :---: | :---: |
| $\begin{aligned} & \mathrm{H}_{3} \mathrm{C}_{6} \mathrm{H}_{5} \mathrm{O}_{7}(\mathrm{aq})+\mathrm{H}_{2} \mathrm{O}(\ell) \rightleftharpoons \mathrm{H}_{2} \mathrm{C}_{6} \mathrm{H}_{5} \mathrm{O}_{7}^{-}(\mathrm{aq})+\mathrm{H}_{3} \mathrm{O}^{+}(\mathrm{aq}) \\ & \text { or } \\ & \mathrm{H}_{2} \mathrm{C}_{6} \mathrm{H}_{5} \mathrm{O}_{7}^{-}(\mathrm{aq})+\mathrm{H}_{2} \mathrm{O}(\ell) \rightleftharpoons \mathrm{H}_{3} \text { or } \mathrm{H}_{5} \mathrm{O}_{7}(\mathrm{aq})+\mathrm{OH}^{-}(\mathrm{aq}) \\ & \text { (one minor error, include lack of double arrow) } \end{aligned}$ |  | $2$ <br> (1) |
|  | Total | 2 |
| Note: State symbols are not required. |  |  |

(b) Calculate the mass of monosodium citrate, $\mathrm{NaH}_{2} \mathrm{C}_{6} \mathrm{H}_{5} \mathrm{O}_{7}$, that would need to be dissolved in 100.0 mL of distilled water to make the most effective buffer solution.
(3 marks)

| Description |  | Marks |
| :---: | :---: | :---: |
| To form most effective buffer $\mathrm{c}\left(\mathrm{H}_{3} \mathrm{C}_{6} \mathrm{H}_{5} \mathrm{O}_{7}\right)=\mathrm{c}\left(\mathrm{H}_{2} \mathrm{C}_{6} \mathrm{H}_{5} \mathrm{O}_{7}^{-}\right)=0.200 \mathrm{~mol} \mathrm{~L}^{-1}$ |  | 1 |
| $\begin{aligned} \mathrm{V}\left(\mathrm{H}_{2} \mathrm{C}_{6} \mathrm{H}_{5} \mathrm{O}^{-}\right) & =100.0 \mathrm{~mL} \\ \mathrm{c}\left(\mathrm{H}_{2} \mathrm{C}_{6} \mathrm{H}_{5} \mathrm{O}_{7}^{-}\right) & =0.200 \mathrm{~mol} \mathrm{~L} \\ \mathrm{n}\left(\mathrm{H}_{2} \mathrm{C}_{6} \mathrm{H}_{5} \mathrm{O}_{7}^{-}\right) & =\mathrm{c} . \mathrm{V}=0.200 \times 0.100 \\ & =0.0200 \mathrm{~mol} \end{aligned}$ |  | 1 |
| $\begin{aligned} \mathrm{m}\left(\mathrm{NaH}_{2} \mathrm{C}_{6} \mathrm{H}_{5} \mathrm{O}_{7}\right) & =\text { n.M }=0.0200 \times 214.106 \\ & =4.28 \mathrm{~g} \end{aligned}$ |  | 1 |
|  | Total | 3 |

(c) If a citric acid buffer was prepared to a pH of 3.5, what would be the concentration of the hydroxide ion at $25.0^{\circ} \mathrm{C}$ ?
(3 marks)

| Description |  | Marks |
| :---: | :---: | :---: |
| $\begin{aligned} & \mathrm{pH}=-\log \left[\mathrm{H}^{+}\right] \\ & -3.5=\log \left[\mathrm{H}^{+}\right] \\ & {\left[\mathrm{H}^{+}\right]=3.1622777 \times 10^{-4}\left(\text { or } 10^{-3.5}\right)} \end{aligned}$ |  | 1 |
| So $\left[\mathrm{OH}^{-}\right]=1 \times 10^{-14} / 3.16 \times 10^{-4}$ |  | 1 |
| $=3.16 \times 10^{-11} \mathrm{~mol} \mathrm{~L}^{-1}$ |  | 1 |
|  | Total | 3 |
| Note: alternative answer $\begin{aligned} \mathrm{pOH} & =14-3.5(1 \text { mark }) \\ & =10.5 \\ {\left[\mathrm{OH}^{-}\right] } & =10^{-10.5}(1 \text { mark }) \\ & =3.16 \times 10^{-11} \mathrm{~mol} \mathrm{~L}^{-1}(1 \text { mark }) \end{aligned}$ <br> (must express in this form for three marks) |  |  |

(d) Explain why only a small change in pH is observed in this buffer solution when a small amount of sodium hydroxide solution is added, compared to adding a similar amount of sodium hydroxide solution to a system that is not a buffer solution. Your answer should refer to the buffer equilibrium in part (a).
(4 marks)

| Description | Marks |
| :--- | :---: |
| The added $\mathrm{OH}^{-}$ions are removed by reaction with citric acid <br> $\left(\mathrm{H}_{3} \mathrm{C}_{6} \mathrm{H}_{5} \mathrm{O}_{7}(\mathrm{aq})+\mathrm{OH}^{-}(\mathrm{aq}) \rightarrow \mathrm{H}_{2} \mathrm{C}_{6} \mathrm{H}_{5} \mathrm{O}_{7}^{-}(\mathrm{aq})+\mathrm{H}_{2} \mathrm{O}(\ell)\right)$ | 1 |
| This increases the concentration of citrate ion $\left(\mathrm{H}_{2} \mathrm{C}_{6} \mathrm{H}_{5} \mathrm{O}_{7}^{-}\right)$and decreases <br> concentration of citric acid $\left(\mathrm{H}_{3} \mathrm{C}_{6} \mathrm{H}_{5} \mathrm{O}_{7}\right)$ so the buffer equilibrium favours <br> production of citric acid | 1 |
| This in turn reduces the concentration of $\mathrm{H}_{3} \mathrm{O}^{+}$ions (This in turn increases <br> concentration of $\left.\mathrm{OH}^{-}\right)$ | 1 |
| Recognition that when $\mathrm{OH}^{-}$is added to a non-buffer solution, there is no <br> weak acid/species for it to react with and so contributes fully to pH change | 1 |
| Total | $\mathbf{4}$ |


| Alternative marking key for Question $29(\mathrm{~d})$ |  |
| :--- | :---: |
| Description | Marks |
| $\mathrm{H}_{3} \mathrm{C}_{6} \mathrm{H}_{5} \mathrm{O}_{7}(\mathrm{aq})+\mathrm{H}_{2} \mathrm{O}(\ell) \rightleftharpoons \mathrm{H}_{2} \mathrm{C}_{6} \mathrm{H}_{5} \mathrm{O}_{7}{ }^{-}(\mathrm{aq})+\mathrm{H}_{3} \mathrm{O}^{+}(\mathrm{aq})$ <br> The OH reacts with the $\mathrm{H}_{3} \mathrm{O}^{+}$and reduces its concentration | 1 |
| The reduction in $\mathrm{H}_{3} \mathrm{O}^{+}$causes the equilibrium to shift to the right <br> or <br> The reduction of the $\mathrm{H}_{3} \mathrm{O}^{+}$reduces the rate of the reverse reaction more <br> than the forward reaction resulting in increase in the amount of $\mathrm{H}_{3} \mathrm{O}^{+}$ <br> present when equilibrium is re-established. <br> or <br> Therefore, the forward reaction rate is greater relative to the reverse until <br> equilibrium is re-established and the $\left[\mathrm{H}_{3} \mathrm{O}^{+}\right]$increases again as the system <br> adjusts. | 1 |
| As the concentration of $\mathrm{H}_{3} \mathrm{O}^{+}$increases to close to before the addition of <br> OH- |  |
| Recognition that when $\mathrm{OH}^{-}$is added to a non-buffer solution, there is no <br> weak acid/species for it to react with and so contributes fully to pH change | 1 |
| Total | $\mathbf{4}$ |

(e) Increasing the concentration of this buffer solution will increase its buffering capacity. Explain this statement.

| Description | Marks |
| :--- | :---: |
| Recognition that increasing the concentration involves increasing the <br> concentrations of both the citric acid and citrate ions. | 1 |
| The increased concentrations of citric acid and citrate ions will be able to <br> react with more hydrogen ions, $\mathrm{H}^{+}(\mathrm{aq})$, or hydroxide ions, $\mathrm{OH}^{-}(\mathrm{aq})$, that are <br> added to the system. | 1 |
| Therefore more acid or base could be added before there is significant <br> change in the pH of the system | 1 |
| Total | $\mathbf{3}$ |

(a) In the space below, draw the structural formula of PCTG, showing two repeating units.

| Description | Marks |
| :---: | :---: |
|  | 2 <br> (1) |
| Total | 2 |
| Note: One complete unit maximum of one mark Minor errors include missing hydrogens, terminating, not including ring in benzen Incorrect or missing ester links is a major error, 0 marks |  |

(b) State the name or give the formula of the by-product of this polymerisation process.
(1 mark)

| Description | Marks |  |
| :---: | :---: | :---: |
| water or $\mathrm{H}_{2} \mathrm{O}$ | Total | $\mathbf{1}$ |

(c) Name two reagents that could be used to synthesise (C) from (B) in Step $3 . \quad$ (2 marks)

| Description | Marks |  |
| :--- | :---: | :---: |
| - acidified (potassium/sodium) dichromate (solution) and acidified <br> (potassium) permanganate (solution) | 2 |  |
| orsulfuric acid with either (potassium) permanganate solution or <br> (potassium/sodium) dichromate solution |  |  |
| (potassium/sodium) dichromate solution and/or |  |  |
| - (potassium/sodium) permanganate solution | 1 |  |
| Note: No acid mentioned, maximum of one mark |  | $\mathbf{2}$ |

(d) Write a balanced half equation to show (B) reacting to form (C).


## Question 31

(a) Write the equilibrium constant expression for this reaction.

| Description | Marks |
| :--- | :---: |
| $\mathrm{K}=\frac{\left[\mathrm{H}_{3} \mathrm{AsO}_{4}\right]^{2}\left[\mathrm{O}_{2}\right]^{5}}{\left[\mathrm{O}_{3}\right]^{5}}$ | 2 |
| (one minor error) | Total |
| Note: Minor errors - not including K=, single transcription error | $\mathbf{2}$ |
|  |  |

(b) The arsenate ion, $\mathrm{HAsO}_{4}{ }^{2-}(\mathrm{aq})$, is amphoteric, meaning it can act as an acid and as a base.
(i) With the aid of equations, describe the amphoteric nature of $\mathrm{HAsO}_{4}{ }^{2-}$ in this aqueous solution.

| Description | Marks |
| :--- | :---: |
| Recognition that $\mathrm{HAsO}_{4}{ }^{2-}$ has the capacity to both gain and lose a <br> proton or indicates which equation represents $\mathrm{HAsO}_{4}{ }^{2-}$ acting as an <br> acid or acting as a base | 1 |
| Equation 1 $\mathrm{HAsO}_{4}{ }^{2-}+\mathrm{H}_{2} \mathrm{O} \rightleftharpoons \mathrm{H}_{2} \mathrm{AsO}_{4}+\mathrm{OH}^{-}$ | 1 |
| Equation 2 $\mathrm{HAsO}_{4}{ }^{2-}+\mathrm{H}_{2} \mathrm{O} \rightleftharpoons \mathrm{AsO}_{4}{ }^{3-}+\mathrm{H}_{3} \mathrm{O}^{+} \quad$ Total | 1 |
|  | $\mathbf{3}$ |

(ii) State why an aqueous solution containing $\mathrm{HAsO}_{4}{ }^{2-}$ is found to have a $\mathrm{pH}>7$ at $25^{\circ} \mathrm{C}$.

| Description | Marks |
| :--- | :---: |
| Given the solution is basic, reaction 1 must occur to a greater <br> extent than reaction 2. <br> or <br> The $K$ value of reaction 1 in (i) is larger than the $K$ value of reaction <br> 2 in (ii). |  |
|  | 1 |

Use the following equations to explain why an increasing concentration of atmospheric carbon dioxide will decrease the formation of calcium carbonate.

| $\mathrm{CO}_{2}(\mathrm{~g})$ | $\rightleftharpoons \mathrm{CO}_{2}(\mathrm{aq})$ |  | Equation 1 |
| ---: | :--- | ---: | :--- |
| $\mathrm{CO}_{2}(\mathrm{aq})+\mathrm{H}_{2} \mathrm{O}(\ell)+\mathrm{CO}_{3}{ }^{2-}(\mathrm{aq})$ | $\rightleftharpoons 2 \mathrm{HCO}_{3}(\mathrm{aq})$ |  | Equation 2 |
| $\mathrm{Ca}^{2+}(\mathrm{aq})+\mathrm{CO}_{3}{ }^{2-}(\mathrm{aq})$ | $\rightleftharpoons \mathrm{CaCO}_{3}(\mathrm{~s})$ |  | Equation 3 |


| Description | Marks |  |
| :--- | :---: | :---: |
| - $\quad$ increased concentration of $\mathrm{CO}_{2}($ aq $)$ shifts equilibrium forward (equation 2) |  |  |
| - thus reducing $\left[\mathrm{CO}_{3}{ }^{2-}(\mathrm{aq})\right]$ | $1-4$ |  |
| - decreased $\mathrm{CO}_{3}{ }^{2-}$ shifts equilibrium left in equation 3 |  |  |
| - therefore the formation of $\mathrm{CaCO}_{3}$ is decreased (equation 3) | Total | $\mathbf{4}$ |

## Question 33

(a) Given that the molecular formulas indicate that the compounds contain the same number of carbon atoms and differ only in the number of one or two hydrogen or oxygen atoms, propose an hypothesis for why there is a variation in the boiling points of these compounds.

| Description | Marks |
| :--- | :---: |
| A detailed statement based on the information available and other relevant <br> knowledge that can be tested experimentally | $\mathbf{2}$ |
| A brief statement based on available information that can be tested <br> experimentally | 1 |
|  | Total |

Possible answers for two marks include:

- Variation in boiling points can be accounted for by the different arrangements of the oxygen and hydrogen atoms (functional groups) attached to the carbon chain.
- Increasing boiling points can be accounted for by increasing intermolecular forces which occur because of different arrangements of atoms/functional groups.
- The boiling point is affected to a greater extent by the presence of one oxygen atom rather than the absence of two hydrogen atoms.
Possible answers for one mark include:
- Boiling points increase with increasing strength of intermolecular forces
- Boiling points increase with increasing solubility
(b) Explain why these organic compounds have very different solubilities in water. (6 marks)

| Description | Marks |
| :--- | :---: |
| The strength of the dispersion forces between butane and water is weak <br> compared to the forces of attraction between butane molecules dispersion <br> and water molecules dispersion, (dipole-dipole) and hydrogen bonding and <br> so there is insufficient energy available to disrupt the butane-butane and <br> water-water forces of attraction and so solubility is low. | 1 |
| The strength of the dispersion, dipole-dipole and hydrogen bonding <br> between butanone and water is greater than the forces of attractions <br> between the molecules in each substance, dispersion/dipole in butanone, <br> and so sufficient energy is released during bond formation to disrupt more <br> forces of attraction in the solute and solvent and so has higher solubility. | $1-2$ |
| The strength of the forces of attractions between butan-1-ol and water, <br> dispersion, dipole-dipole and hydrogen bonding, is greater than the forces <br> of attractions between the molecules in each substance, dispersion, dipole- <br> dipole and hydrogen bonding in butan-1-ol, and so sufficient energy is <br> released, and dissolving occurs. | $1-2$ |
| As there are more sites for hydrogen bonding between water and butan-1- <br> ol, solubility will be greater than for butanone and water. | 1 |
|  | $\mathbf{6}$ |

Question 33 (continued)

| Alternative marking key for Question 33 (b) |  |
| :--- | :---: |
| Description | Marks |
| Recognition that dissolving occurs when the energy released during the <br> forming of solute-solvent forces of attraction is sufficient to disrupt the <br> solute-solute and solvent-solvent forces of attraction. | 1 |
| Recognition that force of attraction between water and butane are weak <br> due to only dispersion forces and reference stronger dispersion and <br> hydrogen bonding in water and so solubility is poor. | 1 |
| Recognition that there is hydrogen bonding, dispersion and dipole-dipole <br> forces of attraction between butanone and water and are stronger than the <br> forces of attraction between molecules in butanone and between <br> molecules in water, and so dissolving occurs. | $1-2$ |
| Recognition that there is hydrogen, dispersion and dipole-dipole forces of <br> attraction between butan-1-ol and water and are stronger than the forces of <br> attraction between the molecules in butan-1-ol and between the molecules <br> in water and so dissolving occurs. | 1 |
| Recognition that there is more hydrogen bonding between butan-1-ol and <br> water compared to butanone and water and so the solubility is higher | 1 |
|  | $\mathbf{T o t a l}$ |

(c) Complete the table below to describe a chemical test that could be used to distinguish between butan-1-ol and butanoic acid by stating the reagent/s used and the distinguishing observations.

| Reagent used | acidified (potassium) permanganate (solution) |  |
| :--- | :--- | :--- |
| Substance being tested | Butan-1-ol | Butanoic acid |
| Observation | purple solution decolourises | no observable change |
| Note: accept $\mathrm{H}^{+} / \mathrm{MnO}_{4}^{-}$ |  |  |

or

| Reagent used | acidified (potassium) dichromate (solution) |  |  |
| :--- | :---: | :---: | :---: |
| Substance being tested | Butan-1-ol | Butanoic acid |  |
| Observation | orange solution turns green | no observable change |  |
| Note: accept $\mathrm{H}^{+} / \mathrm{Cr}_{2} \mathrm{O}_{7}^{2-}$ |  |  |  |

or

| Reagent used | carbonate/hydrogencarbonate (salt) solution |  |  |
| :--- | :---: | :---: | :---: |
| Substance being tested | Butan-1-ol | Butanoic acid |  |
| Observation | no observable change | effervescence/bubbling |  |
| Note: accept $\mathrm{CO}_{3}{ }^{2-}(\mathrm{aq})$ |  |  |  |

or

| Reagent used | add an acidified alcohol - specify alcohol |  |
| :--- | :--- | :--- |
| Substance being tested | Butan-1-ol | Butanoic acid |
| Observation | no observable change | fruity odour detected |

or

| Reagent used | add an acidified carboxylic acid - specify acid |  |
| :--- | :--- | :--- |
| Substance being tested | Butan-1-ol | Butanoic acid |
| Observation | fruity odour detected | no observable change |

or

| Reagent used | magnesium |  |
| :--- | :---: | :---: |
| Substance being tested | Butan-1-ol | Butanoic acid |
| Observation | No observable change | effervescence/bubbling |


| Description | Marks |
| :--- | :---: |
| reagent used | 1 |
| observation with butan-1-ol | 1 |
| observation with butanoic acid | $\mathbf{1}$ |
| Accept other relevant answers |  |

For the molecular formula $\mathrm{C}_{6} \mathrm{H}_{12} \mathrm{O}$ draw two different structural isomers, one which can be readily oxidised by acidified dichromate solution and one which cannot be readily oxidised by acidified dichromate solution. Show all atoms.

Isomer that can be readily oxidised by acidified dichromate solution.
Isomer that cannot be readily oxidised by acidified dichromate solution.

| Description |  |  |  |  |  |  |  | Marks |
| :--- | :--- | :--- | :--- | :---: | :---: | :---: | :---: | :---: |
| Isomer which can be readily oxidised by acidified dichromate solution |  |  |  |  |  |  |  |  |
| Readily oxidized isomer (aldehyde) |  |  |  |  |  |  |  |  |

## Question 35

(16 marks)
A chemical, commonly called iopromide (IOP), is used to enhance the images produced by a medical procedure called a CT scan. It contains carbon, hydrogen, iodine, nitrogen and oxygen, $\mathrm{C}_{\mathrm{v}} \mathrm{H}_{w} \mathrm{~N}_{\mathrm{X}} \mathrm{NO}_{\mathrm{z}}$.

Use the following information to determine the molecular formula of IOP.

- The molar mass of IOP is $791.102 \mathrm{gmol}^{-1}$.
- A 5.62 g sample of IOP contained 0.2986 g of nitrogen, N .
- A 3.54 g sample of IOP is fully combusted to produce;
1.72 L of carbon dioxide gas, $\mathrm{CO}_{2}(\mathrm{~g})$, at $125^{\circ} \mathrm{C}$ and 155.3 kPa . 0.967 g of water vapour, $\mathrm{H}_{2} \mathrm{O}(\mathrm{g})$.
- All of the iodine contained in a 2.523 g sample of IOP is converted to iodide, $\mathrm{I}^{-}$. This sample is then dissolved in water and excess lead(II) nitrate solution, $\mathrm{Pb}\left(\mathrm{NO}_{3}\right)_{2}(\mathrm{aq})$, is added to precipitate the iodine as lead(II) iodide, $\mathrm{Pbl}_{2}(\mathrm{~s})$. This produced 2.21 g of lead(II) iodide.

| Description | Marks |
| :---: | :---: |
| Carbon |  |
| $\begin{aligned} & \mathrm{V}\left(\mathrm{CO}_{2}\right)=1.72 \mathrm{~L} \\ & \mathrm{~T}=125+273.15=398.15 \mathrm{~K} \\ & \mathrm{P}=155.3 \mathrm{kPa} \end{aligned}$ |  |
| $\begin{aligned} & P V=n R T \\ & 155.3 \times 1.72=n \times 8.314 \times 398.15 \end{aligned}$ | 1 |
| $\begin{aligned} & \mathrm{n}(\mathrm{C})=\mathrm{n}\left(\mathrm{CO}_{2}\right)=0.08069 \mathrm{~mol} \\ & \mathrm{~m}(\mathrm{C})=0.08069 \times 12.01=0.9691 \mathrm{~g} \end{aligned}$ | 1 |
| Percentage of C by mass in IOP $=0.9691 / 3.54 \times 100=27.37 \%$ | 1 |
| Hydrogen |  |
| $\begin{aligned} & \mathrm{m}\left(\mathrm{H}_{2} \mathrm{O}\right)=0.967 \mathrm{~g} \\ & \mathrm{n}\left(\mathrm{H}_{2} \mathrm{O}\right)=0.967 / 18.016 \quad=0.05367 \mathrm{~mol} \end{aligned}$ | 1 |
| $\begin{array}{lll} \mathrm{n}(\mathrm{H}) & =2 \times 0.053674511 & =0.1073 \mathrm{~mol} \\ \mathrm{~m}(\mathrm{H}) & =0.107349023 \times 1.008 & =0.1082 \mathrm{~g} \end{array}$ | 1 |
| Percentage of H by mass in IOP $=0.1082 / 3.54 \times 100=3.05 \%$ | 1 |
| Nitrogen |  |
| Percentage of N by mass in IOP $=0.2986 / 5.62 \times 100=5.313 \%$ | 1 |
| lodine |  |
| $\begin{aligned} & \mathrm{m}\left(\mathrm{PbI}_{2}\right)=2.21 \mathrm{~g} \\ & \mathrm{M}\left(\mathrm{Pbl}_{2}\right)=461 \mathrm{~g} \mathrm{~mol}^{-1} \end{aligned}$ | 1 |
| $\begin{aligned} & \mathrm{n}\left(\mathrm{PbI}_{2}\right)=2.21 / 461=4.793926247 \times 10^{-3} \mathrm{~mol} \\ & \mathrm{n}(\mathrm{I}) \quad=2 \times 4.793 \times 10^{-3}=0.009587 \mathrm{~mol} \\ & \mathrm{~m}(\mathrm{I}) \quad=0.009587 \times 126.9=1.216 \mathrm{~g} \end{aligned}$ | 1 |
| Percentage of I by mass in IOP $=1.216 / 2.523 \times 100=48.22$ \% | 1 |
| Oxygen (add all \% to work out \% oxygen) |  |
| $\begin{gathered} \text { \% oxygen }=100-(27.37 \% \mathrm{C}+3.056 \% \mathrm{H}+5.313 \% \mathrm{~N}+ \\ 48.22 \% \mathrm{I}) \end{gathered}$ | 1 |
| $=100-83.97=16.03 \%$ | 1 |

Question 35 (continued)

|  | C | H | N | 1 | 0 |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| \% | 27.37 | 3.056 | 5.313 | 48.22 | 16.03 |  |
| Moles | 2.279 | 3.032 | 0.3792 | 0.3800 | 1.002 | 1 |
| Mole ratio | 6.0107 | 7.996 | 1 | 1 | 2.64 | 1 |
|  | $\times 3$ | $\times 3$ | x 3 | x 3 | x $3=7.92$ |  |
|  | 18 | 24 | 3 | 3 | 8 | 1 |
| this gives the formula as $\mathrm{C}_{18} \mathrm{H}_{24} \mathrm{I}_{3} \mathrm{~N}_{3} \mathrm{O}_{8}$ |  |  |  |  |  |  |
| the empirical formula mass $\left(791.102 \mathrm{~g} \mathrm{~mol}^{-1}\right)$ is identical to the molar mass given so this is also the molecular formula mass ( $791.102 \mathrm{~g} \mathrm{~mol}^{-1}$ ) |  |  |  |  |  | 1 |
|  |  |  |  |  | Total | 16 |


| Alternative marking key for Question 35 (for proportion method) |  |
| :---: | :---: |
| Description | Marks |
| Carbon in 3.54 g |  |
| $\begin{aligned} & \text { PV = nRT } \\ & 155.3 \times 1.72=n \times 8.314 \times 398.15 \end{aligned}$ | 1 |
| $\begin{aligned} & n(C)=n\left(\mathrm{CO}_{2}\right)=0.08069 \mathrm{~mol} \\ & m(\mathrm{C})=0.080694356 \times 12.01=0.9691 \mathrm{~g} \end{aligned}$ | $\begin{aligned} & 1 \\ & 1 \end{aligned}$ |
| Hydrogen in 3.54 g |  |
| $\begin{array}{ll} m\left(\mathrm{H}_{2} \mathrm{O}\right)=0.967 \mathrm{~g} & \\ \mathrm{n}\left(\mathrm{H}_{2} \mathrm{O}\right)=0.967 / 18.016 \quad=0.05367 \mathrm{~mol} \end{array}$ | 1 |
| $\begin{array}{lll} \hline \mathrm{n}(\mathrm{H}) & =2 \times 0.053674511 & =0.10735 \mathrm{~mol} \\ \mathrm{~m}(\mathrm{H}) & =0.107349023 \times 1.008 & =0.1082 \mathrm{~g} \end{array}$ | 1 |
| Nitrogen in 3.54 g |  |
| Mass N in $3.54 \mathrm{~g}=0.2986 / 5.62 \times 3.54=0.1881 \mathrm{~g}$ | 1 |
| lodine in 3.54 g |  |
| $\begin{aligned} & \mathrm{m}\left(\mathrm{PbI}_{2}\right)=2.21 \mathrm{~g} \\ & \mathrm{M}\left(\mathrm{Pbl}_{2}\right)=461 \mathrm{~g} \mathrm{~mol}^{-1} \end{aligned}$ | 1 |
| $\begin{aligned} & \mathrm{n}\left(\mathrm{PbI}_{2}\right)=2.21 / 461=4.793926247 \times 10^{-3} \mathrm{~mol} \\ & \mathrm{n}(\mathrm{I}) \\ & \mathrm{m}(\mathrm{I}) \\ & =9 \times 4.793926247 \times 10^{-3}=9.588 \times 10^{-3} \mathrm{~mol} \\ & \end{aligned}$ | 1 |
| Mass I in $3.54 \mathrm{~g}=1.217 / 2.523 \times 3.54=1.707 \mathrm{~g}$ | 1 |
| Oxygen (add all mass to work out mass of oxygen) |  |
| m oxygen $=3.54-(\mathrm{mC}+\mathrm{mH}+\mathrm{m} \mathrm{N}+\mathrm{ml})$ | 1 |
| $=3.54-2.972=0.5675 \mathrm{~g}$ | 1 |


|  | C | H | N | 1 | 0 |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| mass | 0.9691 g | 0.1082 g | 0.1881 g | 1.7071 g | 0.5675 g |  |
| Moles | 0.0807 | 0.1073 | 0.0134 | 0.01345 | 0.0355 | 1 |
| Mole ratio | 6 | 8.01 | 1 | 1 | 2.65 | 1 |
|  | $\times 3$ | $\times 3$ | $\times 3$ | x 3 | x $3=7.92$ | 1 |
|  | $18$ | $24$ | 3 | 3 | 8 |  |
| this gives | the formula | $\mathrm{C}_{18} \mathrm{H}_{24} \mathrm{I}_{3} \mathrm{~N}_{3}$ |  |  |  |  |
| the empirical formula mass $\left(791.102 \mathrm{~g} \mathrm{~mol}^{-1}\right)$ is identical to the molar mass given so this is also the molecular formula mass ( $791.102 \mathrm{~g} \mathrm{~mol}^{-1}$ ) molecular formula is $\mathrm{C}_{18} \mathrm{H}_{24} \mathrm{I}_{3} \mathrm{~N}_{3} \mathrm{O}_{8}$ |  |  |  |  |  | 1 |
|  |  |  |  |  | Total | 16 |

## Question 36

(a) Outline the difference between random and systematic errors. Give an example of each that might be encountered in this investigation.

| Description | Marks |
| :---: | :---: |
| Random errors are fluctuations in measured data due to limitations of the measurement device/technique, they occur unpredictably and to varying extents throughout the experiment. Non-directional effect on results. | 1 |
| Examples <br> - concentration of heavy metals may vary at depth <br> - judgement made by the sample collector about depth <br> - different collectors at different sites | 1 |
| Systematic errors are repeatable errors that occur due to faulty equipment or poor experiment design, they occur consistently to the same extent throughout the experiment. Results are skewed in one direction. | 1 |
| Examples <br> - faulty timer <br> - concentration of the nitric acid may not be exactly the same as labelled <br> - water bottles and whether or not 100 mL is marked accurately on them (glassware uncertainty). | 1 |
| Total | 4 |

Water samples were collected from 59 of the 180 possible locations.
(b) (i) Calculate the sample size as a percentage of the testable locations.
(1 mark)

| Description | Marks |
| :---: | :---: |
| sample size $=(59 / 180) \times 100=32.8 \%$ | 1 |

(ii) Propose two reasons why samples were not collected from all locations.
(2 marks)

|  | Description | Marks |
| :--- | :---: | :---: |
| $\mathbf{\bullet}$ too expensive |  |  |
| $\bullet$ too time consuming |  | $1-2$ |
| $\bullet$ | accessibility |  |
| - sample size deemed to be sufficient (statistical) |  |  |
|  | Total | $\mathbf{2}$ |

Note: reason to be related to the context of the investigation
(iii) Predict the effect of using a smaller sample size on the reliability of the overall results. Justify your answer.

| Description | Marks |
| :--- | :---: |
| Prediction: less reliable | 1 |
| Justification: <br> - smaller sample size means less locations sampled which <br> means less chance of collecting samples with the full range of <br> heavy metal concentrations | 1 |
| or  <br> $\bullet$ Increases the effect of random errors |  |

Question 36 (continued)
(c) Calculate the minimum mass of sodium phosphate needed to remove all the cadmium(II) ions from the water. Give your final answer to the appropriate number of significant figures. Assume that there are no species other than cadmium(II) ions present in the dam that will react with the phosphate ions.

(a) Explain the effect of temperature on reaction rate by applying collision theory. Support your explanation with an appropriate and clearly-labelled diagram.

| Description | Marks |
| :---: | :---: |
| - increased temperature increases the average kinetic energy of the particles | 1 |
| - thus more particles have energy higher than the $\mathrm{E}_{\mathrm{A}}$ (or clearly annotated on diagram) | 1 |
| - therefore, a greater proportion of the collisions are successful | 1 |
| - the particles are also moving faster so collide more frequently | 1 |
| - therefore the overall rate of reaction is higher | 1 |
| Clearly labelled diagram as below: (Boltzmann diagram) |  |
| For copyright reasons this image cannot be reproduced in the online version of this document, but may be viewed at https://blogs.glowscotland.org.uk/gc/hchemunit/files/2015/06/Picture27.png | 1-2 |
| Total | 7 |

## Question 37 (continued)

This graph shows how the amount of gallium extracted from red mud varies over time at a hydrochloric acid concentration of $1.00 \mathrm{~mol} \mathrm{~L}^{-1}$.

(b) Sketch on the graph above the result that would be obtained if the hydrochloric acid concentration was changed to $2.00 \mathrm{~mol} \mathrm{~L}^{-1}$.

| Description | Marks |
| :--- | :---: |
| Line drawn clearly with steeper slope | 1 |
| Line drawn clearly reaching same plateau/horizontal earlier | 1 |
|  | Total |

(c) Use collision theory to justify the position and shape of the graph you sketched in part (b). Assume that all other reaction conditions were kept constant.

| Description | Marks |
| :--- | :---: |
| -higher concentration means (more particles present in the same <br> volume hence) greater frequency of collisions therefore the rate of <br> reaction increases (hence the steeper slope for $2.00 \mathrm{~mol} \mathrm{~L}^{-1} \mathrm{HCl}$ ) | 1 |
| the faster reaction rate explains why the reaction is finished faster <br> (reaches a plateau) | 1 |
|  | Total |

(d) State one reason why the hydrochloric acid used in this analysis needed to be in excess.

| Description | Marks |  |  |
| :--- | :---: | :---: | :---: |
| Any one of the following: |  |  |  |
| $\bullet$ to ensure that all of the gallium was extracted from the red mud |  |  |  |
| $\bullet$ so it doesn't limit the reaction | 1 |  |  |
| $\bullet$ so all of the gallium(III) oxide reacts |  |  |  |
| To ensure maximum gallium yield |  |  | $\mathbf{1}$ |

## Question 38

(a) With reference to rates of reaction, equilibrium and economic considerations, explain why the above conditions are considered optimal for the Sabatier reaction. In your response, address any compromises in conditions.
(7 marks)

| Description | Marks |
| :---: | :---: |
| Temperature <br> - high temperature gives a fast rate of reaction <br> - reaction is exothermic so a low temperature increases the product yield <br> - so temperature conditions to maximise rate and yield are opposite therefore a compromise is needed. <br> - hence the moderate temperature of $\left(300-400^{\circ} \mathrm{C}\right)$ that is used | 1-3 |
| Pressure <br> - high pressure increases the rate of the reaction <br> - in this reaction high pressure increases the product yield | 1-2 |
| Economics <br> - using high temperatures and pressures increases the operating costs (high energy requirements, cost of specialised equipment) | 1 |
| - use of catalyst which increases the reaction rate, means lower pressures and temperatures can be used to achieve reasonable reaction rates, improving the economics of the process | 1 |
| Total | 7 |

(b) Predict the effect of each of the following changes on the methane yield in the Sabatier reaction.

| Description |  | Marks |
| :---: | :---: | :---: |
| - a suitable catalyst is added | no effect | 1 |
| - the volume of the reaction vessel is increased | decrease | 1 |
| - the temperature is increased | decrease | 1 |
| - methane is removed through a special valve as soon as it forms | increase | 1 |
| - the partial pressure of carbon dioxide is decreased | decrease |  |
|  | Total | 5 |


| Imposed change | Effect on methane yield <br> (circle your answer) |  |  |
| :--- | :--- | :--- | :--- |
| a suitable catalyst is added | increase | decrease | no effect |
| the volume of the reaction <br> vessel is increased | increase | deecrease | no effect |
| the temperature is increased | increase | decrease | no effect |
| methane is removed through <br> a special valve as soon as it <br> forms | increase | decrease | no effect |
| the partial pressure of carbon <br> dioxide is decreased | increase | ddecrease | no effect |

## Question 38 (continued)

Graphs can be drawn to show the effects of imposed changes on equilibrium systems. The graph below shows the effects of some changes that might be made to the reacting system in a flexible vessel.

(c) With reference to the above graph, answer the following questions.
(i) What happened at $T_{1}$ ?

| Description | Marks |
| :--- | :---: |
| Any one of the following:  <br> - the system has reached equilibrium  <br> - addition of a catalyst  <br>  no change |  |
|  | Total |

(ii) Identify the change imposed at each time in the table below.

| Description | Marks |
| :--- | :---: |
| $\mathrm{T}_{2}:$ the volume of the reaction vessel is decreased <br> (Accept: Pump in same number of moles of each reactant at the <br> same time) | 1 |
| $\mathrm{~T}_{3}:$ some $\mathrm{H}_{2} \mathrm{O}$ was removed from the reaction vessel | 1 |
|  | $\mathbf{2}$ |

(iii) The temperature of the reaction vessel was decreased at $\mathrm{T}_{4}$. Sketch on the graph above to show how this affected the partial pressures of all species present. Include any changes to scale and continue until a new equilibrium is established.

| Description | Marks |
| :--- | :---: |
| $\mathrm{CO}_{2}$ and $\mathrm{H}_{2}$ curve down | 1 |
| $\mathrm{CH}_{4}$ and $\mathrm{H}_{2} \mathrm{O}$ curve up | 1 |
| $\mathrm{H}_{2}$ changes 4 x as much as $\mathrm{CO}_{2}$ and $\mathrm{H}_{2} \mathrm{O}$ changes 2 x as much as | 1 |
| $\mathrm{CH}_{4}$ |  |$|$| Total |
| :--- |
| Note: focus on whether lines represent increase or decrease (and by what <br> factor) |

(a) Draw the full structural formula of this section of the reflectin. Show all hydrogen atoms.
(3 marks)

|  | Marks |
| :--- | :---: | :---: |
| Note: circles relate to part (b) |  |

(b) Circle one peptide bond in the structure that you drew in part (a).

| Description | Marks |
| :---: | :---: |
| correctly circled on diagram as below | 1 |
| Total | 1 |
| Note other acceptable alternatives as shown in diagram above in (a) or as | ws: |

(c) Draw the full structural formula of leucine, Leu, in each of the conditions specified below.

Show all hydrogen atoms.
(4 marks)

|  | Description | Marks |
| :---: | :---: | :---: |
| Low pH (acidic) <br> (one minor error) |  | 2 <br> (1) |
|  | Subtotal | 2 |
| High pH (basic) |  | 2 |
| (one minor error) |  | (1) |
|  | Subtotal | 2 |
|  | Total | 4 |

(d) Explain why the structure of Leu is pH dependent.

| Description | Marks |  |  |  |
| :--- | :---: | :---: | :---: | :---: |
| at neutral pH (it exists as zwitterion) and has $\mathrm{NH}_{3}{ }^{+}$and $\mathrm{COO}^{-}$ | 1 |  |  |  |
| amine group (is basic and so) is protonated at low (acidic) pH | 1 |  |  |  |
| carboxylic acid group (is acidic and so) is deprotonated at high (basic) pH | 1 |  |  |  |
| Total |  |  |  | $\mathbf{3}$ |


| Alternative marking key for Question 39 (d) if considered as zwitterion (name not required) |  |  |
| :---: | :---: | :---: |
| Description |  | Marks |
| Neutral pH contains $\mathrm{NH}_{3}{ }^{+}$and $\mathrm{COO}^{-}$groups |  | 1 |
| $\mathrm{NH}_{3}^{+}$( is acidic and) is deprotonated at high pH |  | 1 |
| $\mathrm{COO}^{-}$(is basic and) is protonated at low pH |  | 1 |
|  | Total | 3 |

Question 39 (continued)
(e) Identify the pair most strongly attracted to each other. Justify your choice.

| Description | Marks |
| :--- | :---: |
| Cys and Cys (this can be demonstrated by circling the diagram) | 1 |
| Cys and Cys have disulfide bridges/bonds which are covalent. <br> The disulfide bridges/bonds are much stronger than the other forces of <br> attraction (intermolecular forces) present between the other pairs. | $\mathbf{1 - 2}$ |
| Total | $\mathbf{3}$ |

## Question 40

The molar mass of salicylic acid is $138.121 \mathrm{~g} \mathrm{~mol}^{-1}$.
The molar mass of aspirin is $180.158 \mathrm{~g} \mathrm{~mol}^{-1}$.
This reaction is an equilibrium reaction with a K value of approximately 5 .
(a) In the synthesis of aspirin, 45.0 g of salicylic acid was reacted with excess acetic anhydride. This produced 50.2 g of aspirin. What was the percentage yield of this reaction?

|  | Description | Marks |
| :--- | :---: | :---: |
| n (aspirin) $=\mathrm{n}$ (salicylic acid) | $=45 / 138.121$ | 1 |
|  | $=0.3258 \mathrm{~mol}$ | 1 |
| m (aspirin) | $=0.3258 \times 180.158$ | 1 |
|  | $=58.695 \mathrm{~g}$ | 1 |
| $\%$ yield | $=50.2 / 58.695 \times 100$ | 1 |
|  | $=85.5 \%$ | $\mathbf{T o t a l}$ |

(b) Complete the equation for the reaction between salicylic acid and sodium hydroxide solution.

(c) Calculate the mass of salicylic acid in the sample and therefore the percentage purity of the sample.
(5 marks)

|  | Description | Marks |
| :--- | :--- | :---: |
| $\mathrm{n}\left(\mathrm{OH}^{-}\right)$in 20.00 mL | $=0.0966 \times 0.01845$ | 1 |
|  | $=1.78 \times 10^{-3} \mathrm{~mol}$ | 1 |
|  | $=\mathrm{n}($ acid in 20.00 mL$)$ | 1 |
| $\mathrm{n}($ acid in 250 mL$)$ | $=250 / 20 \times 1.78 \times 10^{-3}$ |  |
|  | $=2.23 \times 10^{-2} \mathrm{~mol}$ | 1 |
| $\mathrm{~m}($ acid $)$ | $=138.121 \times\left(2.23 \times 10^{-2}\right)$ |  |
| $\%$ purity | $=3.08 \mathrm{~g}$ | 1 |
|  | $=(3.08 / 3.55) \times 100$ | Total |
|  | $=86.7 \%$ | $\mathbf{5}$ |

Question 40 (continued)
(d) Identify two incorrect procedures from the list above, select the effect on the calculated concentration of salicylic acid and give the reason for the effect.

| Description | Marks |  |
| :--- | :--- | :---: |
| Number 3 | 1 |  |
| Effect | decrease | 1 |
| Explanation | there will be a lower titre of NaOH | 1 |
|  | Subtotal | $\mathbf{3}$ |
| Number | 4 | 1 |
| Effect | decrease | 1 |
| Explanation | the acid will be diluted in the pipette due to residual water <br> from rinsing | $\mathbf{1}$ |
|  | Subtotal | $\mathbf{3}$ |
|  | Total | $\mathbf{6}$ |

Note: If Number 6 identified as an error, the explanation must be appropriated for the effect identified.

## ACKNOWLEDGEMENTS

Question 37(a) Diagram from: Hyndland Science Faculty. (n.d.). Kinetic energy distributions and temperature [Diagram]. Retrieved October, 2018, from https://blogs.glowscotland.org.uk/gc/hchemunit/files/2015/06/Picture27.pn g

Question 39(d) Text under ‘Description’ information from: Leucine. (2018). In Wikipedia. Retrieved October, 2018, from https://en.wikipedia.org/wiki/Leucine

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