4 Colourless solutions of $\mathbf{X}(\mathrm{aq})$ and $\mathbf{Y}(\mathrm{aq})$ react to form an orange solution of $\mathbf{Z}(\mathrm{aq})$ according to the following equation.

$$
\mathbf{X}(\mathrm{aq})+2 \mathbf{Y}(\mathrm{aq}) \rightleftharpoons \mathbf{Z}(\mathrm{aq}) \quad \Delta H=-20 \mathrm{~kJ} \mathrm{~mol}^{-1}
$$

A student added a solution containing 0.50 mol of $\mathbf{X}(\mathrm{aq})$ to a solution containing 0.50 mol of $\mathrm{Y}(\mathrm{aq})$ and shook the mixture.

After 30 seconds, there was no further change in colour.
The amount of $\mathbf{Z}(\mathrm{aq})$ at equilibrium was 0.20 mol .

| 0 | 4 | 1 | Deduce the amounts of $\mathbf{X}(\mathrm{aq})$ and $\mathbf{Y}(\mathrm{aq})$ at equilibrium. |
| :--- | :--- | :--- | :--- |

Amount of $\mathbf{X}(\mathrm{aq})=$ $\qquad$ mol

Amount of $\mathbf{Y}(\mathrm{aq})=$ mol

| 0 | 4 | 2 |
| :--- | :--- | :--- | time of initial mixing until 60 seconds had elapsed.

[3 marks]


| $\mathbf{0}$ | $\mathbf{4}$. |
| :--- | :--- | concentrations of $\mathbf{X}$ and $\mathbf{Z}$ were:

$\mathbf{X}(\mathrm{aq})=0.40 \mathrm{~mol} \mathrm{dm}^{-3}$ and $\mathbf{Z}(\mathrm{aq})=0.35 \mathrm{~mol} \mathrm{dm}^{-3}$.
For this reaction, the equilibrium constant $K_{\mathrm{c}}=2.9 \mathrm{~mol}^{-2} \mathrm{dm}^{6}$.
Calculate a value for the concentration of $\mathbf{Y}$ at equilibrium.
Give your answer to the appropriate number of significant figures.
$[\mathrm{Y}]=$ $\qquad$ $\mathrm{mol} \mathrm{dm}^{-3}$
 $\mathbf{Z}(\mathrm{aq})$ in Question 4.3.

Suggest how the colour of the mixture changed. Give a reason for your answer.

Colour change $\qquad$
Reason $\qquad$
$\qquad$
$\qquad$
$\qquad$

| $\mathbf{0}$ | $\mathbf{4}$ | $\mathbf{5}$ The student warmed the equilibrium mixture from Question 4.3. |
| :--- | :--- | :--- | :--- |

Predict the colour change, if any, when the equilibrium mixture was warmed.
$\qquad$
$\qquad$

| $\mathbf{0}$ | $\mathbf{5}$ |
| :--- | :--- |$\quad$ This question is about equilibrium.

Sulfur trioxide decomposes to form sulfur dioxide and oxygen at temperature $\boldsymbol{T}_{\mathbf{1}}$ according to the equilibrium shown.

$$
2 \mathrm{SO}_{3}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{SO}_{2}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g}) \quad \Delta H=+196 \mathrm{~kJ} \mathrm{~mol}^{-1}
$$

The graph in Figure 4 shows the concentrations of sulfur trioxide and of oxygen over a period of 6 minutes at temperature $\boldsymbol{T}_{1}$

Figure 4

 Explain your answer.

Time $\qquad$ minutes

Explanation $\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$

| $\mathbf{0}$ | $\mathbf{5}$ | .2 | Sketch on the graph in Figure $\mathbf{4}$ how the concentration of sulfur dioxide changes over |
| :--- | :--- | :--- | :--- | these 6 minutes at temperature $\boldsymbol{T}_{1}$


| $\mathbf{0}$ | $\mathbf{5}$ | $\mathbf{3}$ The temperature of the mixture was changed to $\boldsymbol{T}_{\mathbf{2}}$ and the mixture left to establish a |
| :--- | :--- | :--- | new equilibrium.

In the new equilibrium mixture the concentration of sulfur trioxide was found to be $0.07 \mathrm{~mol} \mathrm{dm}^{-3}$

Deduce which of $\boldsymbol{T}_{1}$ and $\boldsymbol{T}_{2}$ is the higher temperature.
Explain your deduction.

Higher temperature $\qquad$
Explanation $\qquad$
$\qquad$
$\qquad$
$\qquad$

| 0 | 8 | $M e t h a n o l$ |
| :--- | :--- | :--- |

$$
\mathrm{CO}(\mathrm{~g})+2 \mathrm{H}_{2}(\mathrm{~g}) \rightleftharpoons \mathrm{CH}_{3} \mathrm{OH}(\mathrm{~g})
$$

| $\mathbf{0}$ | $\mathbf{8}$ | $\mathbf{1}$ State and explain the effect of using a catalyst on the yield of methanol in this |
| :--- | :--- | :--- | :--- | equilibrium.

[2 marks]
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$

| $\mathbf{0}$ | $\mathbf{8}$. | $\mathbf{2}$ Give an expression for the equilibrium constant $\left(K_{\mathrm{c}}\right)$ for this reaction. |
| :--- | :--- | :--- |


| $\mathbf{0}$ | $\mathbf{8}$. | 3 |
| :--- | :--- | :--- | A mixture of carbon monoxide and hydrogen was allowed to reach equilibrium in a container of volume $250 \mathrm{~cm}^{3}$ at temperature $T$.

At equilibrium, the mixture contained 0.340 mol of carbon monoxide, 0.190 mol of hydrogen and 0.0610 mol of methanol.

Calculate the value of the equilibrium constant $\left(K_{\mathrm{c}}\right)$ for this reaction at temperature $T$.

| 0 | $\mathbf{8}$ | $\mathbf{4}$ Methanol decomposes on heating in a reaction that is the reverse of that used in its |
| :--- | :--- | :--- | :--- | manufacture.

$$
\mathrm{CH}_{3} \mathrm{OH}(\mathrm{~g}) \rightleftharpoons \mathrm{CO}(\mathrm{~g})+2 \mathrm{H}_{2}(\mathrm{~g})
$$

Use your answer from Question $\mathbf{0 8 . 3}$ to determine the value of $K_{\mathrm{c}}$ for this equilibrium at temperature $T$.
State the units for this value of $K_{c}$
(If you were unable to complete the calculation in Question 08.3, assume a value of $K_{\mathrm{c}}=0.825 \mathrm{~mol}^{-2} \mathrm{dm}^{6}$. This is not the correct value.)

Value of $K_{c}$ $\qquad$

Units of $K_{c}$

## Section A

Answer all questions in this section.
1 Ethene reacts with steam in the presence of an acid catalyst to form ethanol.

$$
\mathrm{CH}_{2}=\mathrm{CH}_{2}(\mathrm{~g})+\mathrm{H}_{2} \mathrm{O}(\mathrm{~g}) \rightleftharpoons \mathrm{CH}_{3} \mathrm{CH}_{2} \mathrm{OH}(\mathrm{~g})
$$

 Deduce the units of $K_{c}$.

Expression $\qquad$
$\qquad$
$\qquad$
Units $\qquad$

| $\mathbf{0}$ | $\mathbf{1}$ | $\mathbf{2}$ An equilibrium mixture was found to contain 0.700 mol of ethene, 1.20 mol of |
| :--- | :--- | :--- | :--- | steam and 4.40 mol of ethanol at a temperature $\boldsymbol{T}$. The volume of the container was $2.00 \mathrm{dm}^{3}$.

Calculate a value of $K_{\mathrm{c}}$ for this equilibrium at this temperature.
Give your answer to an appropriate number of significant figures.
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$
$\qquad$

| $\mathbf{0}$ | $\mathbf{4} \quad$ Compounds $\mathbf{A}$ and $\mathbf{B}$ react together to form an equilibrium mixture containing |
| :--- | :--- | compounds $\mathbf{C}$ and $\mathbf{D}$ according to the equation

$$
2 \mathbf{A}+\mathbf{B} \rightleftharpoons 3 \mathbf{C}+\mathbf{D}
$$

| 0 | 4 | 1 |
| :--- | :--- | :--- | A beaker contained $40 \mathrm{~cm}^{3}$ of a $0.16 \mathrm{~mol} \mathrm{dm}^{-3}$ aqueous solution of $\mathbf{A}$. $9.5 \times 10^{-3} \mathrm{~mol}$ of $\mathbf{B}$ and $2.8 \times 10^{-2} \mathrm{~mol}$ of $\mathbf{C}$ were added to the beaker and the mixture was left to reach equilibrium.

The equilibrium mixture formed contained $3.9 \times 10^{-3} \mathrm{~mol}$ of $\mathbf{A}$.
Calculate the amounts, in moles, of $\mathbf{B}, \mathbf{C}$ and $\mathbf{D}$ in the equilibrium mixture.

Amount of D mol

| $\mathbf{0}$ | $\mathbf{4} \cdot \mathbf{2}$ Give the expression for the equilibrium constant $\left(K_{\mathrm{c}}\right)$ for this equilibrium and its units. |
| :--- | :--- | :--- | [2 marks]

$K_{\text {c }}$

Units $\qquad$

| $\mathbf{0}$ | $\mathbf{4} .3$ A different equilibrium mixture of these four compounds, at a different temperature, |
| :--- | :--- | :--- | contained 0.21 mol of $\mathbf{B}, 1.05 \mathrm{~mol}$ of $\mathbf{C}$ and 0.076 mol of $\mathbf{D}$ in a total volume of $5.00 \times 10^{2} \mathrm{~cm}^{3}$ of solution.

At this temperature the numerical value of $K_{\mathrm{c}}$ was 116
Calculate the concentration of $\mathbf{A}$, in $\mathrm{mol} \mathrm{dm}^{-3}$, in this equilibrium mixture. Give your answer to the appropriate number of significant figures.

| 0 | $\mathbf{4} .4$ | Justify the statement that adding more water to the equilibrium mixture in |
| :--- | :--- | :--- | Question 04.3 will lower the amount of $\mathbf{A}$ in the mixture.

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

| Question | Marking guidance | Mark | AO | Comments |
| :---: | :---: | :---: | :---: | :---: |
| 04.1 | amount of $X=0.50-0.20=0.30(\mathrm{~mol})$ <br> amount of $\mathrm{Y}=0.50-2 \times 0.20=0.10(\mathrm{~mol})$ | $\begin{aligned} & 1 \\ & 1 \end{aligned}$ | $\begin{aligned} & \text { AO2h } \\ & \text { AO2h } \end{aligned}$ |  |
| 04.2 | Axes labelled with values, units and scales that use over half of each axis <br> Curve starts at origin <br> Then flattens at 30 seconds at 0.20 mol | $\begin{aligned} & 1 \\ & 1 \\ & 1 \end{aligned}$ | $\begin{aligned} & \mathrm{AO} 2 \mathrm{~h} \\ & \mathrm{AO} 2 \mathrm{~h} \\ & \mathrm{AO} 2 \mathrm{~h} \end{aligned}$ | All three of values, units and scales are required for the mark |
| 04.3 | $\begin{aligned} & \text { Expression }=K_{\mathrm{c}}=\frac{[\mathrm{Z}]}{[\mathrm{X}] \mathrm{Y}]^{2}} \\ & {[\mathrm{Y}]^{2}=\frac{[\mathrm{Z}]}{[\mathrm{X}] K_{\mathrm{c}}}} \\ & {[\mathrm{Y}]=(0.35 / 0.40 \times 2.9)^{0.5}=0.5493=0.55\left(\mathrm{~mol} \mathrm{dm}^{-3}\right)} \end{aligned}$ | 1 <br> 1 <br> 1 | AO1a <br> AO2b <br> AO1b | Answer must be to 2 significant figures |
| 04.4 | Darkened / went more orange <br> The equilibrium moved to the right <br> To oppose the increased concentration of $Y$ | $\begin{aligned} & 1 \\ & 1 \\ & 1 \end{aligned}$ | $\begin{aligned} & \mathrm{AO} 2 \mathrm{~g} \\ & \mathrm{AO} 2 \mathrm{~g} \\ & \mathrm{AO} 2 \mathrm{~g} \end{aligned}$ |  |
| 04.5 | The orange colour would fade | 1 | AO3 1a |  |


| Qu | Marking Guidance | Additional Comments | Mark |
| :---: | :--- | :--- | :--- |
| 5.1 | 3 minutes | M2 dependent on M1 or near miss | 1 |
| 5.2 | (At equilibrium, rate ${ }_{\text {fwd }}=$ rate $_{\text {back }}$ so) concentrations (of $\mathrm{O}_{2}$ and $\mathrm{SO}_{3}$ ) <br> remain constant | Not concentrations are the same/equal <br> Allow (after this point) gradient is zero / <br> curve flattens out | 1 |


| Question | Marking Guidance | Mark | Comments |
| :---: | :---: | :---: | :---: |
| 8.1 | M1 no effect (on yield) <br> M2 increases rate / speed of both / forward and reverse reactions equally / by the same amount | $1$ | CE $=0$ if yield changes <br> If no reference to effect on yield, could still score M2 <br> Ignore reference to no change in position of equilibrium, and reference to lowering activation energies <br> M2 allow changes rate of both / forward and reverse reactions equally / by the same amount |

8.2

$$
\left(K_{c}=\right) \frac{\left[\mathrm{CH}_{3} \mathrm{OH}\right]}{[\mathrm{CO}]\left[\mathrm{H}_{2}\right]^{2}}
$$

Must be square brackets Ignore state symbols Ignore units

M1 divides moles by volume ( 0.250 or $\frac{250}{1000}$ )
M2 $\quad K_{C}=\frac{\frac{0.0610}{0.250}}{\left[\frac{0.340}{0.250}\right]\left[\frac{0.190}{0.250}\right]^{2}}\left(=\frac{0.244}{1.36 \times 0.76^{2}}\right)$
M3 $\quad 0.311$

Correct answer scores 3; M3 to at least 2sf (0.3106159 ...) ; ignore units
Allow ECF from M1 to M2 if an attempt to calculate concentration has been made by dividing by some factor of $250 \mathrm{~cm}^{3}$
Allow ECF from M2 to M3 for use of an expression containing each reagent in a correctly substituted $\mathrm{K}_{\mathrm{c}}$ expression
If volume not used, then allow M3 only for 4.97 (4.96985 ... to at least 2sf)

| 8.4 | $\mathbf{M 1} \frac{1}{\text { Answer to } 8.3}=3.22$ | 1 | $\mathbf{M 1}$ to at least $2 \mathrm{sf}(0.31$ gives $3.2(258)$ ) |
| :---: | :--- | :---: | :--- |
| $\mathbf{M 2} \mathrm{mol}^{2} \mathrm{dm}^{-6}$ | 1 | $\mathbf{M 1}=1.21$ if alternative answer to 8.3 used |  |
| If an error was made in 8.3, but the candidate produced an |  |  |  |
| answer in 8.4 that did fit the inverted calculation from 8.3, |  |  |  |
| then candidate could score $\mathbf{M 1}$ |  |  |  |
| (if volumes are not used, then candidate would get $0.20(12)$. |  |  |  |


| Question | Marking Guidance | Mark | Comments |
| :---: | :---: | :---: | :---: |
| 01.1 | M1 $\quad\left(\mathrm{K}_{\mathrm{c}}=\right) \frac{\left[\mathrm{CH}_{3} \mathrm{CH}_{2} \mathrm{OH}\right]}{\left[\mathrm{CH}_{2}=\mathrm{CH}_{2}\right]\left[\mathrm{H}_{2} \mathrm{O}\right]}$ <br> M2 $\mathrm{mol}^{-1} \mathrm{dm}^{3}$ | 1 <br> 1 | M1 penalise missing brackets or use of (); allow correct molecular formulae in correct expression (and allow $\mathrm{CH}_{2} \mathrm{CH}_{2}$ ); <br> ignore powers shown as 1 <br> M2 units must be in simplest form on one line (or $\mathrm{dm}^{3} \mathrm{~mol}^{-1}$ ) <br> M2 units are consequential on expression in M1 ( $\mathrm{mol}^{-1} \mathrm{dm}^{3}$ only scores if it is the units for the expression in M1) |


| 01.2 | M1 $\frac{\left[\frac{4.40}{2.00}\right]}{\left[\frac{0.70}{2.00}\right] \times\left[\frac{1.20}{2.00}\right]}$ or <br> M2 10.5 (must be 3sf) | $\frac{2.20}{0.35 \times 0.60}$ |  | $\frac{4.40}{0.70 \times 1.20}$ | $\times 2.00$ | 1 | 10.5 (3sf) scores both marks; <br> correct value to 2 sf (10) or 4sf or more (10.476...) scores 1 mark <br> Volume not used is $C E=0$ <br> If use incorrect expression for $K_{c}$ in 1.2 then no marks in 1.2 <br> If a value from the question is copied incorrectly into the expression, could still score M2 if then used correctly in calculation (AE -1) <br> Ignore units |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |


| Question | Answers | Mark | Additional Comments/Guidance |
| :---: | :---: | :---: | :---: |
| 04.1 | $\begin{aligned} & \text { Initial amount of } A=6.4 \times 10^{-3} \\ & \text { Equ } A=6.4 \times 10^{-3}-2 x \quad \therefore x=1.25 \times 10^{-3} \\ & B=9.5 \times 10^{-3}-x=8.25 \times 10^{-3} \\ & C=2.8 \times 10^{-2}+3 x=0.0318 \\ & D=x=1.25 \times 10^{-3} \end{aligned}$ | $\begin{aligned} & \text { M1 } \\ & \text { M2 } \\ & \text { M3 } \\ & \text { M4 } \\ & \text { M5 } \end{aligned}$ | If M1 wrong can score max 3 <br> If incorrect $x$ can score max 3 <br> Allow 2 or more sig figs |
| 04.2 | $\begin{aligned} & K_{\mathrm{c}}=\frac{[C]^{3}[D]}{[A]^{2}[B]} \\ & \text { Units }=\mathrm{mol} \mathrm{dm}^{-3} \end{aligned}$ | 1 <br> 1 | Penalise ( ) but mark on in $4.2 \& 4.3$ <br> If $K_{\mathrm{c}}$ wrong no mark for units |
| $\begin{gathered} 04.3 \\ \text { Can see } \\ 4.2 \end{gathered}$ | M1 for correct rearrangement $[A]^{2}=\frac{[C]^{3}[D]}{K_{c}[B]}$ or $[A]=\sqrt{ } \frac{[C]^{3}[D]}{K_{c}[B]}$ M2 for division of mol of $B, C$ and $D$ by correct volume $[A]^{2}=\frac{\left.\left.\left[^{1.05}\right] / 0.5^{3}\right]^{0.076} / 0.5\right]}{116 \times\left[^{0.21 / 0.5}\right]} \text { or } 0.0289 \text { or } 0.0290$ <br> M3 for final answer: $[\mathrm{A}]=\underline{0.17}$ (must be 2 sfs ) | M1 <br> M2 <br> M3 | If $K_{\mathrm{c}}$ wrong in 4.2 can score 1 for dividing by correct volume <br> If $K_{\mathrm{c}}$ correct but incorrect rearrangement can score 1 for dividing by correct volume |
| 04.4 | (All) conc fall: (ignore dilution) <br> Equm moves to side with more moles <br> To oppose the decrease in conc | $1$ | OR $K_{\mathrm{c}}=$ mole ratio $\times 1 / \mathrm{V}$ <br> If vol increases, mole ratio must increase <br> To keep $K_{c}$ constant <br> If only conc of A falls CE=0 <br> If pressure falls $\mathrm{CE}=0$ |
| Total |  | 13 |  |

