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NSW Education Standards Authority

Student Number

2023 HIGHER SCHOOL CERTIFICATE EXAMINATION

Chemistry

General Instructions

- Reading time 5 minutes
- Working time 3 hours
- Write using black pen
- Draw diagrams using pencil
- Calculators approved by NESA may be used
- · A formulae sheet, data sheet and Periodic Table are provided at the back of this paper
- · Write your Centre Number and Student Number at the top of this page

Total marks: 100

Section I – 20 marks (pages 2–11)

- Attempt Questions 1–20
- Allow about 35 minutes for this section

Section II - 80 marks (pages 13-40)

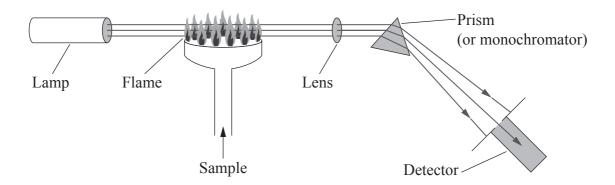
- Attempt Questions 21–37
- · Allow about 2 hours and 25 minutes for this section

Section I

20 marks Attempt Questions 1–20 Allow about 35 minutes for this section

Use the multiple-choice answer sheet for Questions 1–20.

- 1 What is the safest method for disposing of a liquid hydrocarbon after an experiment?
 - A. Pour it down the sink
 - B. Place it in a garbage bin
 - C. Burn it by igniting with a match
 - D. Place it in a separate waste container
- 2 The technique illustrated is used to analyse chemical substances in a sample.



What is the technique shown?

- A. Flame test
- B. Mass spectrometry
- C. Atomic absorption spectroscopy
- D. Ultraviolet-visible spectrophotometry

3 The structural formula of a compound is given.

What is the preferred IUPAC name of this compound?

- A. Pent-2-ene
- B. Pent-2-yne
- C. Pent-3-ene
- D. Pent-3-yne

4 Sodium chloride dissolves in water according to the following equation.

$$NaCl(s) \rightleftharpoons Na^{+}(aq) + Cl^{-}(aq)$$

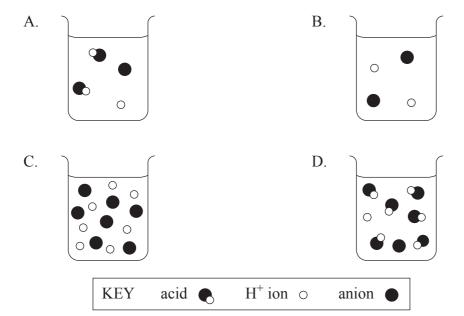
A saturated solution of NaCl in water contains sodium and chloride ions at the following concentrations.

Ion	Concentration (mol L ⁻¹)
Na ⁺	6.13
Cl ⁻	6.13

What is the K_{sp} of sodium chloride?

- A. 2.65×10^{-2}
- B. 8.16×10^{-2}
- C. 12.26
- D. 37.6

5 Which diagram represents the most concentrated weak acid?



6 The pH of a solution changes from 8 to 5.

What happens to the concentration of hydrogen ions during this change of pH?

- A. It increases by a factor of 3.
- B. It decreases by a factor of 3.
- C. It increases by a factor of 1000.
- D. It decreases by a factor of 1000.

A mixture of 0.8 mol of CO(g) and 0.8 mol of $H_2(g)$ was placed in a sealed 1.0 L container. The following reaction occurred.

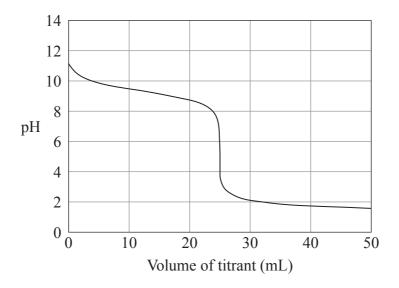
$$\mathrm{CO}(g) \,+\, 2\mathrm{H}_2(g) \ensuremath{\rightleftharpoons}\xspace \mathrm{CH}_3\mathrm{OH}(g)$$

When equilibrium was established, the mixture contained 0.5 mol of CO(g).

What amount of $H_2(g)$ was present at equilibrium?

- A. 0.2 mol
- B. 0.4 mol
- C. 0.6 mol
- D. 1.0 mol

- 8 How many structural isomers have the molecular formula $C_3H_6F_2$?
 - A. 2
 - B. 3
 - C. 4
 - D. 5
- **9** A titration was performed using two solutions of equal concentration, producing the following titration curve.



Which combination of solutions does the titration curve represent?

- A. Addition of a weak base to a weak acid
- B. Addition of a weak base to a strong acid
- C. Addition of a strong acid to a weak base
- D. Addition of a strong acid to a strong base

- Which of the following correctly lists the compounds in order of increasing boiling point?
 - A. Heptane < heptan-2-one < heptan-1-o1 < heptanoic acid
 - B. Heptane < heptan-1-o1 < heptan-2-one < heptanoic acid
 - C. Heptanoic acid < heptan-2-one < heptan-1-o1 < heptane
 - D. Heptanoic acid < heptan-1-o1 < heptan-2-one < heptane
- 11 An indicator solution was obtained by boiling a flower in water.

Flower water indicator chart

Colour	Red			Purple			Blue		Blue-green		Green- yellow	
pН	1	2	3	4	5	6	7	8	9	10	11	12

Two solutions were tested with this indicator.

Which row of the table correctly identifies the colour of each solution?

	$H_2SO_4 (1 \times 10^{-5} \text{ mol L}^{-1})$	NaOH (5 \times 10 ⁻⁵ mol L ⁻¹)
A.	Red	Green-yellow
B.	Red	Blue-green
C.	Purple	Blue-green
D.	Purple	Green-yellow

12 The industrial production of ammonia is represented by the Haber process reaction shown.

$$N_2(g) + 3H_2(g) \rightleftharpoons 2NH_3(g)$$
 $\Delta H = -91.8 \text{ kJ mol}^{-1}$

Factors such as temperature and pressure need to be considered in order to maximise yield.

Which of the following is correct?

- A. A lower pressure would result in a higher yield.
- B. A higher pressure would result in a higher yield.
- C. A lower temperature would result in a lower yield.
- D. A higher temperature would result in a higher yield.
- 13 The table shows four separate tests used to identify a dilute, aqueous sample of a compound.

Test number	Test	Observations
1	Test with red litmus	Stays red
2	Add Ba ²⁺ ions to a sample	White precipitate formed
3	Add OH ⁻ ions to a sample	Brown precipitate formed
4	Add Cl ⁻ ions to a sample	White precipitate formed

Which compound would produce the observations shown?

- A. Silver sulfate
- B. Lead(II) acetate
- C. Iron(II) bromide
- D. Magnesium carbonate

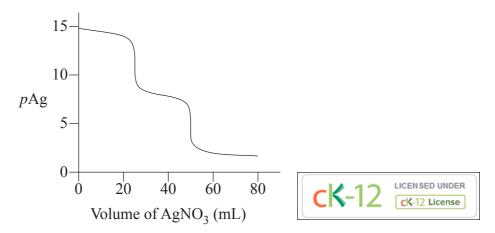
- What volume of 0.540 mol L⁻¹ hydrochloric acid will react completely with 1.34 g of sodium carbonate?
 - A. 11.7 mL
 - B. 23.4 mL
 - C. 29.9 mL
 - D. 46.8 mL
- 15 The table gives the heat of combustion of three different alcohols at 25°C.

Alcohol	Heat of combustion (kJ g ⁻¹)
Methanol	22.68
Ethanol	29.67
Butan-1-ol	36.11

Which of the following gives the best approximation for the molar heat of combustion of propan-1-ol, expressed in $kJ g^{-1}$?

- A. $\left(\frac{22.68 + 29.67 + 36.11}{3}\right)$
- B. $\left(\frac{29.67 + 36.11}{2}\right)$
- C. $\left(\frac{22.68 + 29.67}{2}\right)$
- D. $\left(\frac{3 \times 36.11}{4}\right)$

A solution contains potassium iodide and potassium chloride. It was analysed by performing a precipitation titration using silver nitrate. The titration curve for this reaction is shown, where $pAg = -log_{10}[Ag^+]$.



Why is this a valid and correct procedure for quantifying the amount of each anion present in the mixture?

- A. AgCl would precipitate out first, followed by AgI.
- B. AgI would precipitate out first, followed by AgCl.
- C. Both AgI and AgCl precipitate out of the solution together.
- D. Neither AgCl nor AgI would precipitate out of the solution.

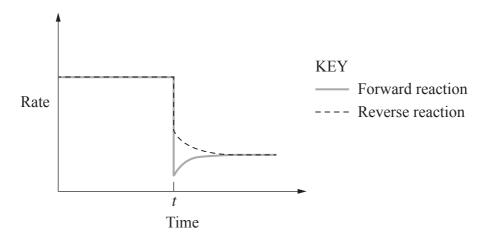
What mass of lead(II) iodide $(MM = 461 \text{ g mol}^{-1})$ will dissolve in 375 mL of water?

- A. 0.233 g
- B. 0.293 g
- C. 0.369 g
- D. 0.621 g

18 Carbon dioxide reacts with hydrogen gas to form carbon monoxide and water vapour in a sealed flask, according to the following equation.

$$CO_2(g) + H_2(g) \rightleftharpoons CO(g) + H_2O(g)$$

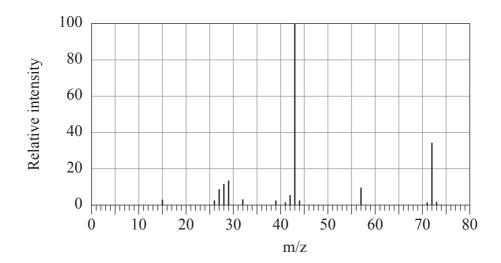
A temperature change was imposed on the equilibrium system at time *t* and the rates of both the forward and reverse reactions were monitored.



Which row of the table correctly identifies the nature of both temperature change at time t and the ΔH of the forward reaction?

	Temperature change at time t	ΔH of the forward reaction
A.	Decrease	+
B.	Decrease	_
C.	Increase	+
D.	Increase	-

19 The diagram shows a simplified mass spectrum for butan-2-one.



Which equation best represents the process that produces the particle responsible for the peak at m/z 43?

- A. $CH_3COCH_2CH_3^+ \rightarrow CH_3CO + {}^+CH_2CH_3$
- B. $CH_3COCH_2CH_3^+ \rightarrow CH_3CO^+ + CH_2CH_3$
- C. $CH_3COCH_2CH_3^+ \rightarrow CH_3CH_2CH_2 + {}^+CHO$
- D. $CH_3COCH_2CH_3^+ \rightarrow CH_3CH_2CH_2^+ + CHO$

20 Nitrogen monoxide and oxygen combine to form nitrogen dioxide, according to the following equation.

$$2NO(g) + O_2(g) \rightleftharpoons 2NO_2(g)$$
 $K_{eq} = 2.47 \times 10^{12}$

A 2.00 L vessel is filled with 1.80 mol of $NO_2(g)$ and the system is allowed to reach equilibrium.

What is the equilibrium concentration of NO(g)?

- $A. \hspace{0.5cm} 0.00 \hspace{0.1cm} \text{mol} \hspace{0.1cm} L^{-1}$
- B. $4.34 \times 10^{-5} \text{ mol L}^{-1}$
- C. $6.90 \times 10^{-5} \text{ mol L}^{-1}$
- D. $8.69 \times 10^{-5} \text{ mol L}^{-1}$

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2023 HIGHER SCHOOL CERTIFICATE EXAMINATION							
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Chemistry							
Section II Answer Booklet			,	Stuc	dent	Nun	nber

80 marks
Attempt Questions 21–37
Allow about 2 hours and 25 minutes for this section

Instructions

- Write your Centre Number and Student Number at the top of this page.
- Answer the questions in the spaces provided. These spaces provide guidance for the expected length of response.
- Show all relevant working in questions involving calculations.
- Extra writing space is provided at the back of this booklet.
 If you use this space, clearly indicate which question you are answering.

Please turn over

Do NOT write in this area.

Question 21 (2 marks)

Some isomers with the formula C₄H₈O are shown.

butan-2-one
$$\begin{array}{c} O \\ H_3C - CH_2 - C - CH_3 \\ \end{array}$$
butanal
$$\begin{array}{c} O \\ \parallel \\ H_3C - CH_2 - CH_2 - C - H \\ \end{array}$$

$$\begin{array}{c} CH_3 & O \\ \parallel & \parallel \\ H_3C - CH - C - H \\ \end{array}$$

Name ONE pair of functional group isomers and ONE pair of chain isomers from the structures above.

Type of isomer	Pair of isomers
Functional group	and
Chain	and

Question 22 (4 marks)

Explain how the following substances would be classified under the Arrhenius and Brønsted–Lowry definitions of acids. Support your answer with relevant equations.

4

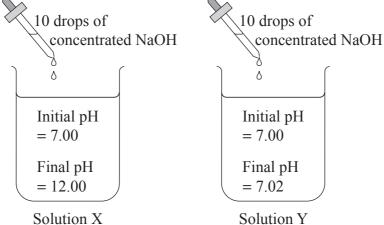
)	HCI(aq	1)					
1	NH ₄ Cl	(aq)					
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Question 23 (3 marks)

The pH of two solutions, X and Y, were measured before and after 10 drops of concentrated NaOH was added to each.

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explain the pri changes that occurred in solutions A and 1.	

Question 24 (2 marks)

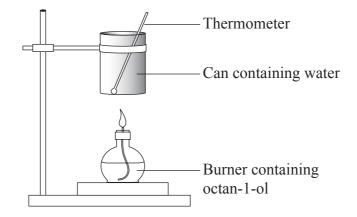
The hydrogen oxalate ion $\left(\mathrm{HC_2O_4}^-\right)$ is classified as amphiprotic.	2
Describe, using chemical equations, how this ion is amphiprotic.	

Please turn over

3

Question 25 (5 marks)

A student used the apparatus shown to investigate the combustion of octan-1-ol.



The following results were obtained by the student.

Mass of water heated = 205 gInitial temperature of water = 23.7°C Final temperature of water = 60.4°C

The following data are given.

Molar enthalpy of combustion of octan-1-ol = $-5294 \text{ kJ mol}^{-1}$ Molar mass of octan-1-ol = $130.23 \text{ g mol}^{-1}$

(a)	Assuming that no energy released by this combustion is lost to the surroundings, calculate the mass of octan-1-ol burnt.			

Question 25 continues on page 19

Question 25	(continued)
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(b) Explain ONE advantage of using a biofuel compared to fossil fuels.					

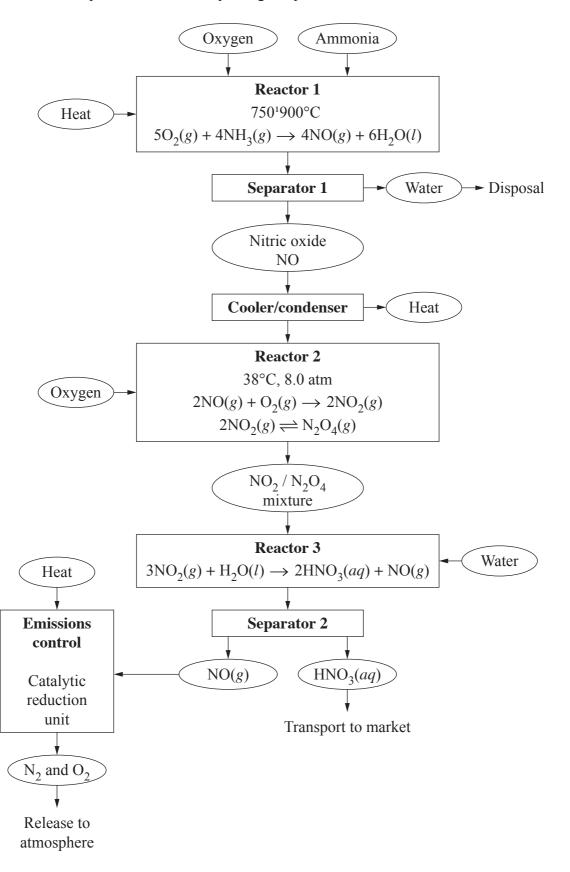
End of Question 25

Please turn over

2

Question 26 (5 marks)

Nitric acid can be produced industrially using the process shown.



Question 26 continues on page 21

Question 26 (continued)

A mixture of NO_2 and N_2O_4 enters Reactor 3, where only NO_2 is consumed by the reaction with water.
Explain, with respect to Le Chatelier's principle, what happens to the $\rm N_2O_4$.
Explain TWO improvements that can be made to the design of the process shown.
shown.
shown.
shown.

End of Question 26

Question 27 (4 marks)

A student has been asked to produce 185 mL of ethanol ($MM = 46.068 \text{ g mol}^{-1}$) by fermenting glucose using yeast, as shown in the equation.

 $\mathrm{C_6H_{12}O_6}(aq) \rightarrow 2\mathrm{C_2H_5OH}(aq) + 2\mathrm{CO_2}(g)$

Given that the densi dioxide gas produced	•	0 kPa.	calculate the	

•••••	 	

Question 28 (5 marks)

Alkene Q undergoes an addition reaction with chlorine gas to form compound R.

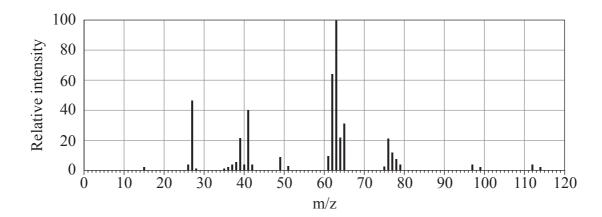
(a)	Describe a chemical test that could be done in a school laboratory to confirm
	that Q is an alkene. Include expected observations in your answer.

2

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(b) Compound R was analysed and found to contain approximately 32% carbon by mass. The mass spectrum of compound R is shown.

3



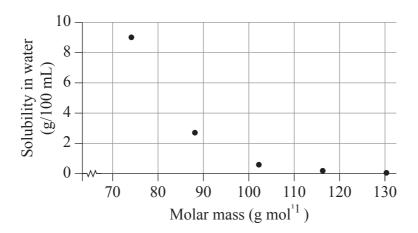
Provide a structural formula for compound R. Support your answer with calculations.

Structural formula of compound R:

3

Question 29 (3 marks)

The following graph shows the solubility of some alkan-1-ols in water at 20°C.



Explain the relationship between the trend shown in the graph and the relevant intermolecular forces.

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 	•••••	 	

Question 30 (4 marks)

A water sample contains at least one of the following anions at concentrations of $1.0 \text{ mol } L^{-1}$.

4

- bromide (Br⁻)
- carbonate (CO_3^{2-})

Outline a sequence of tests that could be performed in a school laboratory to confirm the identity of the anion or anions present. Include expected observations an TWO balanced chemical equations in your answer.

 	 	 ••••••

7

Question 31 (7 marks)

Copper(II) ions (Cu^{2+}) form a complex with lactic acid $(C_3H_6O_3)$, as shown in the equation.

 $Cu^{2+}(aq) + 2C_3H_6O_3(aq) \rightleftharpoons [Cu(C_3H_6O_3)_2]^{2+}(aq)$

This complex can be detected by measuring its absorbance at 730 nm. A series of solutions containing known concentrations of $\left[\mathrm{Cu}(\mathrm{C_3H_6O_3})_2\right]^{2^+}$ were prepared, and their absorbances measured.

Concentration of $\left[\mathrm{Cu}(\mathrm{C_3H_6O_3})_2\right]^{2+}$ (mol L^{-1})	Absorbance
0.000	0.00
0.010	0.13
0.020	0.28
0.030	0.43
0.040	0.57
0.050	0.72

Two solutions containing Cu²⁺ and C₃H₆O₃ were mixed. The initial concentrations of each in the resulting solution are shown in the table.

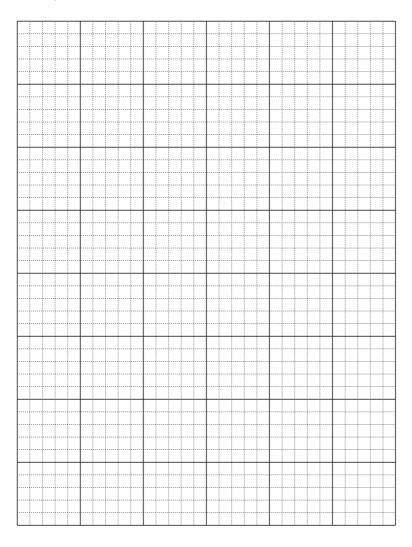
Species	Initial concentration (mol L ⁻¹)
Cu ²⁺	0.056
C ₃ H ₆ O ₃	0.111

When the solution reached equilibrium, its absorbance at 730 nm was 0.66.

You may assume that under the conditions of this experiment, the only species present in the solution are those present in the equation above, and that $\left[\text{Cu}\left(\text{C}_{3}\text{H}_{6}\text{O}_{3}\right)_{2}\right]^{2+}$ is the only species that absorbs at 730 nm.

With the support of a line graph, calculate the equilibrium constant for the reaction.

Question 31 continues on page 27



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End of Question 31

5

Question 32 (5 marks)

The ammonium ion content of mixtures can be determined by boiling the mixture with a known excess of sodium hydroxide. This converts the ammonium ions into gaseous ammonia, which is removed from the system.

$$\mathrm{NH_4}^+(aq) + \mathrm{OH}^-(aq) \rightarrow \mathrm{NH_3}(g) + \mathrm{H_2O}(l)$$

The excess sodium hydroxide can then be titrated with an acid solution of known concentration.

A fertiliser containing ammonium ions was analysed as follows.

- A sample of fertiliser was treated with 50.00 mL of 1.124 mol L⁻¹ sodium hydroxide solution and the solution boiled.
- After all of the ammonia was removed, the resulting solution was transferred to a 250.0 mL volumetric flask and made up to the mark with deionised water.
- 20.00 mL aliquots of this solution were titrated with 0.1102 mol L⁻¹ hydrochloric acid, giving the following results.

Titration	Volume HCl (mL)
1	22.65
2	22.05
3	22.00
4	21.95

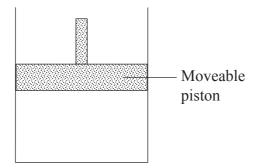
Question 32 continues on page 29

Question 32 (continued)
Calculate the mass of ammonium ions in the sample of fertiliser.

End of Question 32

Question 33 (6 marks)

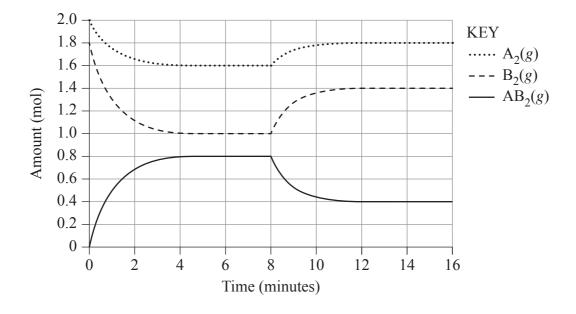
Gases A2 and B2 are placed in a closed container of variable volume, as shown.



The reaction between these substances is as follows.

$$A_2(g) + 2B_2(g) \rightleftharpoons 2AB_2(g)$$
 $\Delta H = -10 \text{ kJ mol}^{-1}$

The following graph shows changes in the amounts (in mol) of these three substances over time in this container.



Question 33 continues on page 31

Question 33 (continued)

a)	Explain what is happening in this system between 6 minutes and 8 minutes.	2
)	Explain TWO different factors that could result in the disturbance at 8 minutes.	4

End of Question 33

5

Question 34 (5 marks)

precipitates. The K_{sp} of magnesium fluoride is 5.16×10^{-11} .
Calculate the equilibrium concentration of magnesium ions in this solution.

When 125 mL of a magnesium nitrate solution is mixed with 175 mL of a 1.50 mol L^{-1} sodium fluoride solution, 0.6231 g of magnesium fluoride ($MM = 62.31 \text{ g mol}^{-1}$)

Question 35 (6 marks)

The following data apply to the ionisation of acetic acid (CH ₃ COOH) and ichloroacetic acid (CCl ₃ COOH). $ \begin{array}{c ccccccccccccccccccccccccccccccccccc$				
Fichloroacetic acid (CCl ₃ COOH).				
Fichloroacetic acid (CCl ₃ COOH).				••••••
Fichloroacetic acid (CCl ₃ COOH).				
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Fichloroacetic acid (CCl ₃ COOH).				
Fichloroacetic acid (CCl ₃ COOH).				
Fichloroacetic acid (CCl ₃ COOH).				
$\begin{array}{c cccc} & CH_{3}COOH & CCl_{3}COOH \\ \hline pK_{a} & 4.76 & 0.51 \\ \hline \Delta H^{\circ} \text{ (kJ mol}^{-1}\text{)} & -0.1 & +1.2 \\ \hline \Delta S^{\circ} \text{ (J K}^{-1} \text{ mol}^{-1}\text{)} & -91.6 & -5.8 \\ \hline -T\Delta S^{\circ} \text{ (kJ mol}^{-1}\text{)} & +27.3 & +1.7 \\ \hline \end{array}$	ving data apply to the	ionisation of ace	etic acid (CH ₃ CC	OOH) and
pK_a 4.76 0.51 ΔH° (kJ mol ⁻¹) -0.1 +1.2 ΔS° (J K ⁻¹ mol ⁻¹) -91.6 -5.8 $-T\Delta S^{\circ}$ (kJ mol ⁻¹) +27.3 +1.7	etic acid (CCl_3COOH).		` -	,
pK_a 4.76 0.51 ΔH° (kJ mol ⁻¹) -0.1 +1.2 ΔS° (J K ⁻¹ mol ⁻¹) -91.6 -5.8 $-T\Delta S^{\circ}$ (kJ mol ⁻¹) +27.3 +1.7		CH ₃ COOH	CCl ₃ COOH	
$\Delta S^{\circ} (J K^{-1} \text{ mol}^{-1})$ -91.6 -5.8 - $T\Delta S^{\circ} (kJ \text{ mol}^{-1})$ +27.3 +1.7	pK_a			
$-T\Delta S^{\circ} \text{ (kJ mol}^{-1}\text{)} +27.3 +1.7$		0.1	.1.2	1
	$\Delta H^{\circ} (\text{kJ mol}^{-1})$	-0.1	+1.2	
$\Delta G^{\circ} \text{ (kJ mol}^{-1}\text{)} +27.2 +2.9$	` '			_
	$\Delta S^{\circ} (J K^{-1} mol^{-1})$	-91.6	-5.8	
	$\Delta S^{\circ} (J K^{-1} mol^{-1})$ $-T\Delta S^{\circ} (kJ mol^{-1})$	-91.6 +27.3	-5.8 +1.7	
	$\Delta S^{\circ} (J K^{-1} mol^{-1})$ $-T\Delta S^{\circ} (kJ mol^{-1})$ $\Delta G^{\circ} (kJ mol^{-1})$	-91.6 +27.3 +27.2	-5.8 +1.7 +2.9	
	$\Delta S^{\circ} (J K^{-1} mol^{-1})$ $-T\Delta S^{\circ} (kJ mol^{-1})$	-91.6 +27.3 +27.2	-5.8 +1.7 +2.9	
	$\Delta S^{\circ} (J K^{-1} mol^{-1})$ $-T\Delta S^{\circ} (kJ mol^{-1})$ $\Delta G^{\circ} (kJ mol^{-1})$	-91.6 +27.3 +27.2	-5.8 +1.7 +2.9	
		etic acid (CCl ₃ COOH).	etic acid (CCl ₃ COOH). CH ₃ COOH	etic acid (CCl ₃ COOH). CH ₃ COOH CCl ₃ COOH

9

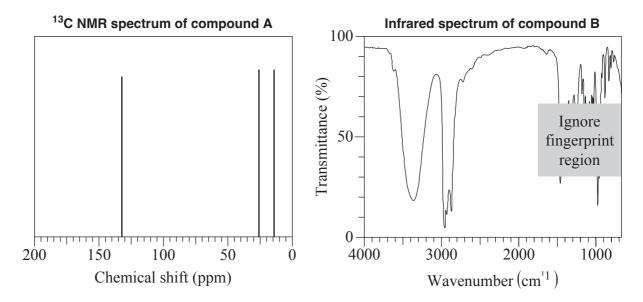
Question 36 (9 marks)

An organic reaction pathway involving compounds A, B and C is shown in the flow chart.

 $\mathbf{A} \xrightarrow{\text{Dilute H}_2\text{SO}_4} \mathbf{B} \xrightarrow{\text{KMnO}_4} \mathbf{C}$

The molar mass of A is $84.156 \,\mathrm{g \, mol}^{-1}$.

A chemist obtained some spectral data for the compounds as shown.



Data from ¹H NMR spectrum of compound C

Chemical shift (ppm)	Relative peak area	Splitting pattern
1.01	3	Triplet
1.05	3	Triplet
1.65	2	Multiplet
2.42	2	Triplet
2.46	2	Quartet

¹H NMR chemical shift data

Type of proton	δ/ppm
$R-C\mathbf{H}_3$, $R-C\mathbf{H}_2-R$	0.7–1.7
H_3C-CO- (aldehydes, ketones, - CH_2-CO-) carboxylic acids or esters)	2.0–2.6
R—С Н О	9.4–10.0
R—COOH	9.0–13.0

Question 36 continues on page 35

Ouestion	36	(continued)
O GCSGOII	20	Communaca

Identify the functional group present in each of compounds A to C and draw the structure of each compound. Justify your answer with reference to the information provided.

Compound A	Functional group:
Compound B	Functional group:
Compound C	Functional group:

Question 36 continues on page 36

Question 36 (continued)

End of Question 36

Question 37 (5 marks)

When performing industrial reductions with CO(g), the following equilibrium is of great importance.

$$2CO(g) \rightleftharpoons CO_2(g) + C(s)$$
 $K_{eq} = 10.00 \text{ at } 1095 \text{ K}$

A 1.00 L sealed vessel at a temperature of 1095 K contains CO(g) at a concentration of 1.10×10^{-2} mol L⁻¹, $CO_2(g)$ at a concentration of 1.21×10^{-3} mol L⁻¹, and excess solid carbon.

Is the system at equilibrium? Support your answer with calculations.
Carbon dioxide gas is added to the system above and the mixture comes to equilibrium. The equilibrium concentrations of $CO(g)$ and $CO_2(g)$ are equal. Excess solid carbon is present and the temperature remains at 1095 K.
Calculate the amount (in mol) of carbon dioxide added to the system.

End of paper

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Chemistry

FORMULAE SHEET

$n = \frac{m}{MM}$	$c = \frac{n}{V}$	PV = nRT
$q = mc\Delta T$	$\Delta G^{\circ} = \Delta H^{\circ} - T \Delta S^{\circ}$	$pH = -\log_{10}[H^+]$
$pK_a = -\log_{10}[K_a]$	$A = \varepsilon lc = \log_{10} \frac{I_o}{I}$	
Avogadro constant, N_A		$6.022 \times 10^{23} \text{ mol}^{-1}$
Volume of 1 mole ideal gas: at		
_	at 0°C (273.15 K)	22.71 L
	at 25°C (298.15 K)	24.79 L
Gas constant		$1.8.314 \text{ J mol}^{-1} \text{ K}^{-1}$
Ionisation constant for water a	t 25°C (298.15 K), K _w	1.0×10^{-14}

DATA SHEET

Solubility constants at 25°C

Compound	K_{sp}	Compound	K_{sp}
Barium carbonate	2.58×10^{-9}	Lead(II) bromide	6.60×10^{-6}
Barium hydroxide	2.55×10^{-4}	Lead(II) chloride	1.70×10^{-5}
Barium phosphate	1.3×10^{-29}	Lead(II) iodide	9.8×10^{-9}
Barium sulfate	1.08×10^{-10}	Lead(II) carbonate	7.40×10^{-14}
Calcium carbonate	3.36×10^{-9}	Lead(II) hydroxide	1.43×10^{-15}
Calcium hydroxide	5.02×10^{-6}	Lead(II) phosphate	8.0×10^{-43}
Calcium phosphate	2.07×10^{-29}	Lead(II) sulfate	2.53×10^{-8}
Calcium sulfate	4.93×10^{-5}	Magnesium carbonate	6.82×10^{-6}
Copper(II) carbonate	1.4×10^{-10}	Magnesium hydroxide	5.61×10^{-12}
Copper(II) hydroxide	2.2×10^{-20}	Magnesium phosphate	1.04×10^{-24}
Copper(II) phosphate	1.40×10^{-37}	Silver bromide	5.35×10^{-13}
Iron(II) carbonate	3.13×10^{-11}	Silver chloride	1.77×10^{-10}
Iron(II) hydroxide	4.87×10^{-17}	Silver carbonate	8.46×10^{-12}
Iron(III) hydroxide	2.79×10^{-39}	Silver hydroxide	2.0×10^{-8}
Iron(III) phosphate	9.91×10^{-16}	Silver iodide	8.52×10^{-17}
		Silver phosphate	8.89×10^{-17}
		Silver sulfate	1.20×10^{-5}

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Infrared absorption data

Bond	Wavenumber/cm ⁻¹
N—H (amines)	3300–3500
O—H (alcohols)	3230–3550 (broad)
С—Н	2850–3300
O—H (acids)	2500–3000 (very broad)
C≡N	2220–2260
c=o	1680–1750
c=c	1620–1680
С—О	1000–1300
С—С	750–1100

¹³C NMR chemical shift data

Type of carbon		δ/ppm								
$\begin{array}{ c c c c c c c c c c c c c c c c c c c$	5–40									
$\begin{bmatrix} R - C - Cl \ o \end{bmatrix}$	10–70									
$ \begin{array}{c c} R - C - C - \\ \parallel & \mid \\ O \end{array} $										
R-C-N		25–60								
- C - O -	alcohols, ethers or esters	50–90								
C = C		90–150								
$R-C \equiv N$		110–125								
		110–160								
R — C — O	esters or acids	160–185								
R — C — O	aldehydes or ketones	190–220								

UV absorption

(This is not a definitive list and is approximate.)

Chromophore	λ_{\max} (nm)
С—Н	122
С—С	135
C=C	162

Chromophore	λ_{\max} (nm)				
C≡C	173 178				
	196 222				
C—Cl	173				
C CI	175				
C—Br	208				
СВ	200				

Some standard potentials

		F	
$K^+ + e^-$	\rightleftharpoons	K(s)	-2.94 V
$Ba^{2+} + 2e^{-}$	\rightleftharpoons	Ba(s)	-2.91 V
$Ca^{2+} + 2e^{-}$	\rightleftharpoons	Ca(s)	-2.87 V
$Na^+ + e^-$	\rightleftharpoons	Na(s)	-2.71 V
$Mg^{2+} + 2e^{-}$	\rightleftharpoons	Mg(s)	-2.36 V
$Al^{3+} + 3e^{-}$	\rightleftharpoons	Al(s)	-1.68 V
$Mn^{2+} + 2e^-$	\rightleftharpoons	Mn(s)	-1.18 V
$H_2O + e^-$	\rightleftharpoons	$\frac{1}{2}\mathrm{H}_2(g) + \mathrm{OH}^-$	-0.83 V
$Zn^{2+} + 2e^{-}$	\rightleftharpoons	Zn(s)	-0.76 V
$Fe^{2+} + 2e^{-}$	\rightleftharpoons	Fe(s)	-0.44 V
$Ni^{2+} + 2e^{-}$	\rightleftharpoons	Ni(s)	-0.24 V
$\mathrm{Sn}^{2+} + 2\mathrm{e}^{-}$	\rightleftharpoons	Sn(s)	-0.14 V
$Pb^{2+} + 2e^{-}$	\rightleftharpoons	Pb(s)	-0.13 V
$H^{+} + e^{-}$	\rightleftharpoons	$\frac{1}{2}$ H ₂ (g)	0.00 V
$SO_4^{2-} + 4H^+ + 2e^-$	\rightleftharpoons	$SO_2(aq) + 2H_2O$	0.16 V
$Cu^{2+} + 2e^{-}$	\rightleftharpoons	Cu(s)	0.34 V
$\frac{1}{2}$ O ₂ (g) + H ₂ O + 2e ⁻	\rightleftharpoons	2OH ⁻	0.40 V
$Cu^+ + e^-$	\rightleftharpoons	Cu(s)	0.52 V
$\frac{1}{2}I_2(s) + e^-$	\rightleftharpoons	I-	0.54 V
$\frac{1}{2}I_2(aq) + e^-$	\rightleftharpoons	I-	0.62 V
$\mathrm{Fe^{3+} + e^{-}}$	\rightleftharpoons	Fe^{2+}	0.77 V
$Ag^+ + e^-$	\rightleftharpoons	Ag(s)	0.80 V
$\frac{1}{2}\mathrm{Br}_2(l) + \mathrm{e}^-$	\rightleftharpoons	Br ⁻	1.08 V
$\frac{1}{2}\mathrm{Br}_2(aq) + \mathrm{e}^{-}$	\rightleftharpoons	Br ⁻	1.10 V
$\frac{1}{2}$ O ₂ (g) + 2H ⁺ + 2e ⁻	\rightleftharpoons	H_2O	1.23 V
$\frac{1}{2}\operatorname{Cl}_2(g) + \mathrm{e}^-$	\rightleftharpoons	Cl¯	1.36 V
$\frac{1}{2}$ Cr ₂ O ₇ ²⁻ + 7H ⁺ + 3e ⁻	\rightleftharpoons	$Cr^{3+} + \frac{7}{2}H_2O$	1.36 V
$\frac{1}{2}\text{Cl}_2(aq) + e^-$	\rightleftharpoons	Cl¯	1.40 V
$MnO_4^- + 8H^+ + 5e^-$	\rightleftharpoons	$Mn^{2+} + 4H_2O$	1.51 V
$\frac{1}{2}F_2(g) + e^-$	\rightleftharpoons	F^-	2.89 V

Aylward and Findlay, *SI Chemical Data* (5th Edition) is the principal source of data for the standard potentials. Some data may have been modified for examination purposes.

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	——————————————————————————————————————	4.C				+							_				+				Ra	1	<u> </u>	
			6	Ц	19.00	17	⊃:	35.45	Chlorine	35	Br	79.90	Bromine	53	Ι	126.9	lodine	\$	At		Astatine	117	Ts	Tennessine
			8	0	16.00	Oxygen 16	2 v	32.07	Sulfur	34	Se	78.96	Selenium	52	Te	127.6	Tellurium	8 4 4	S		Polonium	116	Lv	Moscovium Livermorium
			7	Z	14.01	INIU OBCII	J	30.97	Phosphorus	33	As	74.92	Arsenic	51	Sp	121.8	Antimony	£ 33	<u>P</u>	209.0	Bismuth	115	Mc	Moscovium
			9	ر ر	12.01	11/1	‡ :Z	28.09	Silicon	32	Ge	72.64	Germanium	50	Sn	118.7	Tin	7 8 8	Pb	207.2	Lead	114	丘	Flerovium
			2	В	10.81	13	SE	26.98	Aluminium	31	Сa	69.72	Gallium	49	In	114.8	Indium	<u>∞</u> i	=	204.4	Thallium	113	N	Nihonium
FLEMENTS										30	Zn	65.38	Zinc	48	g	112.4	Cadmium	08 1	БH	200.6	Mercury	112	Cn	Copernicium
										29	Cn	63.55	Copper	47	Ag	107.9	Silver	6/.	Au	197.0	Gold	1111	Rg	Meitnerium Darmstadtium Roentgenium Copernicium
OF THE										28	Z	58.69	Nickel	46	Pd	106.4	Palladium	∞,	K	195.1	Platinum	110	Ds	Darmstadtium
TARLE (KEY	62	Au	197.0	Cold				27	ට	58.93	Cobalt	45	Rh	102.9	Rhodium	<u></u>	Iľ	192.2	Iridium	109	Mt	Meitnerium
			Atomic Number	Symbol	mic Weight	Name				26	Æ	55.85	Iron	44	Ru	101.1	Ruthenium	9,0	S	190.2	Osmium	108	Hs	Hassium
PERIODIC			Ator		Standard Atomic Weight																			Bohrium
										24	Ċ	52.00	Chromium	42	Mo	95.96	Molybdenum	4	>	183.9	Tungsten	106	Sg	Seaborgium
										23	>	50.94	Vanadium	41	Nb	92.91	Niobium	£ E	Ia	180.9	Tantalum	501	Db	Dubnium
										22	Ξ	47.87	Titanium	40	Zr	91.22	Zirconium	7.5	HI	178.5	Hafnium	104	Rf	Rutherfordium
										21	Sc	44.96	Scandium	39	×	88.91	Yttrium	57–71			Lanthanoids	89–103		Actinoids
			4	Be	9.012	1.0	7 <u>Σ</u>	24.31	Magnesium	20	Ca	40.08	Calcium	38	Sr	87.61	Strontium	56	Ба	137.3	Barium	88	Ra	Radium
,	П Н	1.008 Hydrogen	3	Ľ.	6.941	11	Za	22.99	Sodium	19	×	39.10	Potassium	37	Rb	85.47	Rubidium	52	S	132.9	Caesium	87	H H	Francium

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57	58	59	09	19	62	63	64	65	99	<i>L</i> 9	89	69	70	71
La	Ce	Pr	pN	Pm	Sm	En	PS	Tb	Dy	Ho	Ë	Tm	ΛP	Γn
138.9	140.1	140.9	144.2		150.4	152.0	157.3	158.9	162.5	164.9	167.3	168.9	173.1	175.0
Lanthanum	Cerium	Praseodymium	Neodymium	Promethium	Samarium	Europium	Gadolinium	Terbium	Dysprosium	Holmium	Erbium	Thulium	Ytterbium	Lutetium

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102	Š		Nobelium
101	РW		Mendelevium
100	Fm		Fermium
66	Es		Einsteinium
86	Cf		Californium
67	Bk		Berkelium
96	Cm		Curium
95	Am		Americium
94	Pu		Plutonium
93	ď	•	Neptunium
92	n	238.0	Uranium
91	Pa	231.0	Protactinium
06	Th	232.0	Thorium
68	Ac		Actinium

Lawrencium

103 Lr

Standard atomic weights are abridged to four significant figures. Elements with no reported values in the table have no stable nuclides.

Information on elements with atomic numbers 113 and above is sourced from the International Union of Pure and Applied Chemistry Periodic Table of the Elements (November 2016 version). The International Union of Pure and Applied Chemistry Periodic Table of the Elements (February 2010 version) is the principal source of all other data. Some data may have been modified.