# **Queenstown Secondary School**



# Preliminary Examination 2024 Secondary Four Express Chemistry 6092/02

20 August 2024 Tuesday Time: 1115 – 1300h Duration: 1 hour 45 minutes

Setter:

Additional Materials:

Candidates answer on the Question Paper. No Additional Materials are required.

# READ THESE INSTRUCTIONS FIRST

Write your name, class and index number on all the work you hand in. Write in dark blue or black pen. You may use an HB pencil for any diagrams or graphs. Do not use staples, paper clips, glue or correction fluid.

#### Section A

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

#### Section B

Answer **one** question. Write your answers in the spaces provided.

The number of marks is given in brackets [] at the end of each question. A copy of the Periodic Table is printed on page 24.

The use of an approved scientific calculator is expected, where appropriate.

Examiner's Use			
Section A	/70		
Section B	/10		
B9			
B10			
TOTAL	/80		

### Section A

#### Answer **all** questions.

A1 Use the list of substances to answer the questions.

calcium hydroxide	carbon monoxide	copper		
hydrogen	magnesium	nitrogen		
nitrogen dioxide	sulfur dioxide	zinc oxide		

Each substance can be used once, more than once or not at all.

Name the substance(s) which

(a)	is added to soil to increase the pH,	
		[1]
(b)	helps prevent iron in underwater pipes from rusting,	
		[1]
(c)	lead to the formation of acid rain that corrodes limestone buildings,	
	and	[1]
(d)	react in a 1:3 ratio in the Haber Process to produce ammonia.	
	and	[1]
	[Total: 4]	

- A2 Esters are commercially used as solvents for cosmetics, perfumes and glue.
  - (a) Propyl ethanoate has a characteristic odour of pears and is commonly used in fragrances. A bottle of perfume containing propyl ethanoate is left opened in a laboratory and it diffused throughout the laboratory.
    - (i) Give the names and draw the full structural formulae of the carboxylic acid and the alcohol that react to form propyl ethanoate.

(ii) The temperature of the laboratory is increased.

Explain, in terms of collisions between reacting particles, what effect would this have on the rate of diffusion of propyl ethanoate.

.....[2]

(b) The table shows some information about different esters.

Table 2

name	formula	relative molecular mass
methyl methanoate	HCO <sub>2</sub> CH <sub>3</sub>	60
ethyl methanoate	$HCO_2C_2H_5$	74
propyl methanoate	HCO <sub>2</sub> C <sub>3</sub> H <sub>7</sub>	88
butyl methanoate	$HCO_2C_4H_9$	102
pentyl methanoate	$HCO_2C_5H_{11}$	116

101

(i) Use information in the table to give one piece of evidence that suggests that the esters belong to the same homologous series.

.....

- ......[1]
- (ii) Which ester has the greatest rate of diffusion at room temperature and pressure? Explain your answer.

......[2]

A3 Seawater constitutes a rich source of various commercially important elements.

(a) Magnesium bromide is an important salt found in seawater.

Table 3 shows some information about two isotopes of bromine.

Table 3

sym	nbol	number of protons	number of neutrons	number of electrons		
79 35	Br	35		35		
$^{81}_{35}$	Br⁻		46			
(i)	Cor	nplete the table.			[1]	
(!!)		dete forme the table to	and the the terms for the			
(ii)	Use	e data from the table to	explain the term isotop	Des.		
	[1]					
	[1]					
(iii)	Chl	orine is treated with a s	sample of seawater cor	ntaining magnesium		
	bromide.					
	Write the ionic equation for the reaction that occurs.					
	vvri	te the ionic equation to		15.		
	[1]					

[Total: 7]

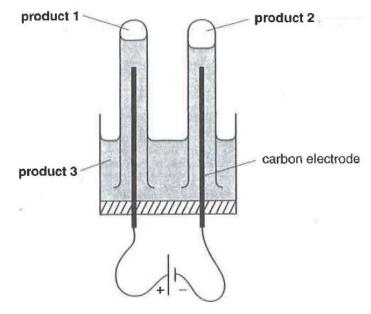
(iv) Explain why astatine does not react with aqueous magnesium bromide. ..... [1] ..... The distinctive smell of the seaside was thought to be due to ozone  $(O_3)$ . (b) Ozone reacts with aqueous potassium iodide as shown in the equation below.  $2I^{-}(aq) + O_{3}(g) + 2H^{+}(aq) \rightarrow I_{2}(aq) + O_{2}(g) + H_{2}O(l)$ (i) Describe what is observed when ozone is bubbled into aqueous potassium iodide. [1] ..... (ii) Explain, in terms of oxidation states, why this is a redox reaction. ..... ..... [2] ..... (c) It is now known that the smell of the seaside is due to the presence of dimethyl sulfide, (CH<sub>3</sub>)<sub>2</sub>S. Draw a 'dot-and-cross' diagram for dimethyl sulfide,  $(CH_3)_2S$ . (i) [2] (ii) Explain, in terms of bonding and structure, why dimethyl sulfide is a gas at room temperature and pressure.

> > [Total: 10]

Table 4.1

1					effect on	1
	electrolyte	state	electrode	half-equation at anode	electrolyte	
	silver nitrate	aqueous			no change	
	copper(II)				fully	
	chloride		graphite		decomposed	[2]

(b) The diagram below shows the electrolysis of concentrated aqueous sodium chloride.





(i) Write the half equation for the reaction that occurs at the negative electrode.

[1]
-----

(ii) The volumes of products 1 and 2 should theoretically be the same.

Explain why the volume of product 1 is lesser.

[1	1]	]	
----	----	---	--

(iii) What happens to the resulting solution during the electrolysis? Explain your reasoning. ..... ..... [1] ..... (iv) The same apparatus can be used to electrolyse dilute aqueous sodium chloride. Give one similarity and one difference between the products of the electrolysis of concentrated and dilute aqueous sodium chloride. similarity ..... ..... . . . . difference ..... ..... ..... [2] ..... [Total: 7]

A5 Ethanol is the intoxicating ingredient of many alcoholic beverages such as beer.

(a) Ethanol is manufactured by the reaction between ethene and steam.

$$C_2H_4$$
 (g) +  $H_2O$  (g)  $\rightarrow C_2H_5OH$  (l)

#### Table 5

bond	C-C	C=C	C-H	O-H	C-O	C=O	0-0	O=0
bond energy in kJ/mol	346	610	414	463	358	804	144	498

(i) Using the bond energies provided in Table 5.1, calculate the enthalpy change for this reaction.

Your diagram should show:

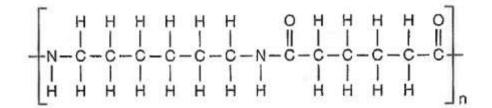
- the products of the reaction
- the activation energy for the reaction
- the enthalpy change of reaction,  $\Delta H$

		energy $C_2H_4 + H_2O$	
(b)	Ethar	progress of reaction nol is also manufactured by the fermentation of glucose.	[3]
	(i)	Write a balanced chemical equation for this fermentation.	
			[1]
	(ii)	Briefly describe this process.	
		Include in your answer the conditions needed for fermentation and how	
		the ethanol is purified.	
			[0]
			[3]

	(c)	Ethanol is oxidised by oxygen in the air to form ethanoic acid.	
		Ethanol and ethanoic acid can be distinguised by chemical tests.	
		Describe two of these chemical tests and the observations that allow you to	
		make the distinctions.	
		test 1	
		observation	
		test 2	
		observation	
			[2]
		[Total: 14]	
<b>A6</b>	Poly	(propene) and nylon are both used to make strong, waterproof ropes.	
	Poly	(propene) is an addition polymer. Nylon is a condensation polymer.	
	(a)	Describe one difference between addition polymers and condensation	
		polymers.	
			[1]
	(h)	Prove the structure of poly(propose), showing two report units	

**(b)** Draw the structure of poly(propene), showing two repeat units.

(c) There are several different types of nylon. One type of nylon is nylon-6,6.This is the repeating unit of nylon-6,6.



(i) Draw the structures of the two monomers that react to form nylon-6,6.

(ii) During the manufacturing process, the chain length of the nylon is controlled so that the nylon polymer molecules have an average relative molecular mass in the range 12 000 to 20 000.

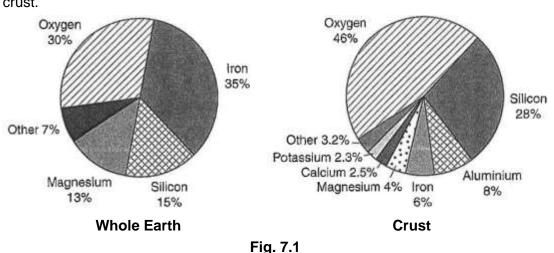
What is the range of the average number of repeating units in the nylon-6,6 molecules? Show your working.

> > [Total: 6]

[2]

**A7** Read the information about elements and compounds in the Earth.

The Earth's crust is the thin outer layer of the Earth. The pie charts show a comparison of the percentages of elements in the whole Earth and in the Earth's crust.



In the Earth's crust, silicon and oxygen are the most abundant elements. Rocks such as quartz are made of covalently bonded compounds of silicon and oxygen. Typically, quartz contains 46.7% silicon and 53.3% oxygen by mass.

Some rocks such as feldspars contain ionic silicate compounds. These contain metal ions ionically bonded to silicate ions.

Examples of naturally occurring silicates are shown in the table below.

name of silicate compound	formula
forsterite	Mg <sub>2</sub> SiO <sub>4</sub>
phenacite	Be <sub>2</sub> SiO <sub>4</sub>
anorthite	CaAl <sub>2</sub> Si <sub>2</sub> O <sub>8</sub>
microcline	KA/Si <sub>3</sub> O <sub>8</sub>

Table 7.2

The formulae of the silicate compounds are not simple. Some silicate compounds contain one type of metal ion, others contain more than one. All silicate ions contain silicon and oxygen, but the numbers of the atoms and the charges on the ions vary. For example,

- phenacite (Be<sub>2</sub>SiO<sub>4</sub>) contains only Be<sup>2+</sup> metal ions and the formula of its silicate ion is SiO<sub>4</sub><sup>4-</sup>,
- microcline (KA/Si<sub>3</sub>O<sub>8</sub>) contains K<sup>+</sup> and Al<sup>3+</sup> metal ions and the formula of its silicate ion is Si<sub>3</sub>O<sub>8</sub><sup>4-</sup>.

(a) Scientists believe that the centre of the Earth is made from an inner core made mainly of iron.

What evidence from the pie charts supports this idea?

......[1]

(b) (i) Use the information to work out the empirical formula and hence the name of the main compound in quartz.

empirical formula .....

(ii) There are other compounds of oxygen found in the Earth's crust as well as quartz. One such example is potassium oxide.

The table shows some differences between the properties of potassium oxide and the compound in **(b)(i)**.

Table 7.3

compound	melting point / °C	electrical conductivity	
compound		under room conditions	
potassium oxide	740	does not conduct	
compound in <b>(b)(i)</b>	2230	does not conduct	

Explain, in terms of bonding and structure, why potassium oxide and the compound in **(b)(i)** have different properties.

[4] Give the formulae and charges of the ions present in anorthite.

(d) Beryllium and silicon can both be extracted from the mineral phenacite.
 Show by calculation that 1 kg of phenacite contains a larger mass of silicon than beryllium but a larger number of moles of beryllium atoms than silicon atoms.

[4]

[Total: 12]

(c)

**A8** A series of experiments was carried out to investigate the effect of different catalysts on the rate of a reaction.

The table shows the time taken for the reaction to finish when different metal compounds were used as catalysts.

The metal compounds contained Group 1 metals, Group 2 metals or transition metals.

ovporiment	ootolyot	temperature at start	time taken for reaction to finish
experiment	catalyst	/ °C	/ s
1	NaC <i>l</i>	19	45
2	FeCl <sub>2</sub>	20	22
3	$CoCl_2$	19	26
4	MgCl <sub>2</sub>	20	46
5	NaNO <sub>3</sub>	19	45
6	Fe(NO <sub>3</sub> ) <sub>2</sub>	20	22
7	Fe(NO <sub>3</sub> ) <sub>3</sub>	19	15
8	Co(NO <sub>3</sub> ) <sub>2</sub>	19	26
9	Mg(NO <sub>3</sub> ) <sub>2</sub>	19	46

Table 8

(a) Explain, in terms of collisions and energy, the effect of a catalyst on the rate of

a reaction.

.....[2]

Group 1 and Group 2 metal compounds are less effective than transition (b) metal compounds as catalysts. Explain how the information in the table supports this statement. ..... ..... ..... [2] Iron is a transition metal. (C) Two different iron ions were used in the experiments. (i) Give the formulae of the two ions. [1] ..... and ..... (ii) Which iron ion appears to be the more effective catalyst? Explain your reasoning. ..... ..... [1] ..... State one other property of transition metals. (iii) [1] .....

(d) A student wrote this conclusion from the results in the table.

The type of anion in the catalyst compound does not affect the rate of reaction.

(i) Do you agree with this conclusion?

Use the results to explain your reasoning.

.....[2]

(ii) Predict the time taken for the reaction to finish if iron(III) chloride was

used as a catalyst.

.....

[1]

[Total: 10]

#### **Section B**

Answer **one** question from this section.

**B9** A factory is suspected of releasing non-biodegradable toxic waste such as calcium ions, aluminium ions, zinc ions and nitrate ions into the nearby river.

A scientist tested two samples of the river water by adding excess aqueous sodium hydroxide and aqueous ammonia respectively, followed by warming with aluminium foil. He recorded his observations in Table 9 below.

	on adding		on warming with
test	Ũ	on adding excess	6
	a few drops		aluminium foil
		white precipitate,	
aqueous	white precipitate	soluble in excess	
sodium	formed	giving a colourless	no visible reaction
hydroxide	lonnou		
		solution	
		white precipitate,	
		some soluble in	
aqueous	white precipitate	excess giving a	gas evolved turns
ammonia	formed	colourless solution,	damp litmus paper
annonia	Ionnea		blue
		some remained	
		insoluble in excess	

#### Table 9

Based on this information, the scientist claimed that the factory had indeed been releasing aluminium ions, zinc ions and nitrate ions into the river. However, the factory owner argued that this data showed that the factory was only releasing aluminium ions and zinc ions, but not nitrate ions.

(a) (i) Explain how the data shows that calcium ions are absent, but both aluminium ions and zinc ions are present in the river water.

19

(ii) The scientist claims that nitrate ions are detected in the river water as the test with aqueous ammonia produced ammonia gas. Do you agree? Explain your answer.

(b) The river water also contained trace amounts of chloride ions. Describe how you would carry out a test to show the presence of chloride ions in a solution. Include the observations you would expect.

.....

[2]

(c) The salt zinc chloride found in the river water can be prepared in the laboratory.

Describe how a pure sample of zinc chloride crystals can be prepared from insoluble zinc carbonate.

 [3]

[Total: 10]

- **B10** The reactivity series summarises information about the reactions of metals with acids and water, displacement reactions and the extraction of metals from their ores.
  - (a) A student carried out some experiments to place four metals, W, X, Y and Z in order of reactivity. The table shows the results.

	metal W	metal X	metal Y	metal <b>Z</b>
solution of <b>W</b> nitrate		×	×	×
solution of <b>X</b> nitrate	~		✓	✓
solution of Y nitrate	~	×		√
solution of <b>Z</b> nitrate	~	×	*	

Table	10	.1
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[key:  $\checkmark$  = shows a reaction happened;  $\varkappa$  = shows no reaction happened]

(i) Place the metals in order of reactivity, starting with the most reactive.
 (ii) Metal Z reacts with hydrochloric acid.

What would you see when metal Z reacts with hydrochloric acid?

Explain your reasoning.

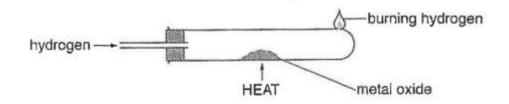
(iii) The student carried out further experiments to place metal M in the list.
 She used dilute hydrochloric acid and samples of the metals.
 She found out that metal M is the fourth most reactive metal.

Describe the experiments that the student carried out.

Your answer should include

- the experiments that she carried out using dilute hydrochloric acid and samples of the metals,
- the measurements that she made,
- how the results showed that metal M is the fourth most reactive metal.

(b) The student also carried out some experiments to investigate the reduction of three metal oxides by hydrogen.



[3]

The table shows the appearance of each metal oxide when cold.

Table 10.2	2
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experiment	metal oxide	colour when cold
1	calcium oxide	white
2	copper oxide	red
3	lead oxide	yellow

(i) What would you expect to see happen in each experiment?

Explain your answer.

(ii) In which tube would you expect to see the fastest reaction?

Explain your answer.

[2] [Total: 10] [insert Periodic Table]

# MARK SCHEME

# Section A [70 m]

A1	(a) (b)		ium hydroxic nesium		<u>r dioxide</u> and <u>nitroge</u> gen and <u>hydrogen</u>	en dioxide	[2] [2]	
A2	(a)	(i) (ii)	ethanoic acid, CH <sub>3</sub> COOH; propanol, C <sub>3</sub> H <sub>7</sub> OH At higher temperatures, the reacting particles move faster leading to a <u>higher frequency of effective collisions</u> between the particles, resulting in a <u>higher rate of diffusion</u> .					
	(b)	(i)	The esters	have the <u>same ger</u>	neral formula, HCO <sub>2</sub> ctional group, HCO <sub>2</sub>		[1]	
		(ii)	Methyl met			-	[1] [1]	
A3	(a)	(i)	symbol	number of	number of	number of		
			<sup>79</sup> 35Br	protons 35	neutrons 44	electrons 35	[1]	
			35DI 81 35Br	35	<u>44</u> 46	36	[1]	
		(ii)			ber of protons (i.e. 3		[1]	
			different nu	umber of neutrons	(i.e. 44 and 146 neu		[4]	
		(iii)		<u>Br (aq) → 2Cl (aq)</u>			[1] [1]	
		(iv)	Astatine, b bromine fro		han bromine, is <u>una</u>	ible to displace	1.1	
	(b)	(i)		less solution turns	dark-brown.		[1]	
	( )	(ii)	The iodine	in pota <u>ssium</u> iodid	e undergoes <u>oxidati</u>		[1]	
					ases from <u>-1 in I<sup>-</sup></u> to		[1]	
				-	bes <u>reduction</u> , as the	e oxidation state of		
	(c)	(i)	oxygen decreases from <u>0 in O<sub>3</sub></u> to <u>-2 in H<sub>2</sub>O</u> . 'Dot-and-cross' diagram [covalent; H <sub>3</sub> C – S – CH <sub>3</sub> ]					
	(-)	(ii)	Dimethyl sulfuide is a simple covalent molecule.				[1]	
					y is required to over			
			intermolecular forces of attraction between molecules.					
A4	(a)	-	er; <u>Ag (s) → A</u>				[1]	
	(h)			$rac{}{} cl_2(g) + 2e^{-1}$			[1] [1]	
	(b)	(i) (ii)						
		(")	lesser.	, is <u>slightly soluble</u>			[1]	
		(iii)	The resulting solution becomes increasingly alkaline.					
			With the removal of H <sup>+</sup> and Cl <sup>-</sup> ions, the remaining Na <sup>+</sup> and OH <sup>-</sup>					
		(iv)	similarity	form the strong al	kali, NaOH.			
		(••)		ode, H+ ions are d	ischarged to form <u>hy</u>	/drogen gas.	[1]	
			difference				[41	
			aqueous se	odium chloride, wh	scharged to form <u>ox</u> ile Cl <sup>-</sup> ions are disch	narged to form	[1]	
			<u>cmonne ga</u>	is for concentrated	aqueous sodium ch			

A5			
_	(a)	(i) energy absorbed for bond-breaking = $610 + 4(414) + 2(463)$ = $3192 \text{ kJ/mol}$	[1]
		energy released for bond-forming = $346 + 5(414) + 358 + 463$ = $3237$ kJ/mol	[1]
		enthalpy change of reaction = $3192 + (-3237) = -45 \text{ kJ/mol}$	[1]
		(ii) <u>More energy is released</u> in bond forming (i.e. C-C, C-H, C-O and O-	[2]
		H) than <u>absorbed</u> in bond breaking (i.e. C=C, C-H and O-H).	
		(iii) on diagram:	[4]
		<ul> <li>the products of the reaction</li> <li>the activation energy for the reaction</li> </ul>	[1] [1]
		<ul> <li>the activation energy for the reaction</li> <li>the enthalpy change of reaction, ΔH</li> </ul>	[1]
	(b)	(i) $C_6H_{12}O_6 \rightarrow 2C_2H_5OH + 2CO_2$	[1]
	(~)	(ii) The fermentation of glucose can be carried out using <u>veast</u> as	[2]
		catalyst. The mixture is kept at a temperature of about <u>37°C</u> ,	
		producing a dilute solution of ethanol, with concentration <u>12-14%</u> .	
		The process is carried out under <u>anaerobic</u> conditions.	[1]
	(0)	Ethanol is purified using <u>fractional distillation</u> .	[1]
	(c)	test 1 : To 5 cm <sup>3</sup> of sample, add 1 g of <u>solid sodium carbonate</u> . observation : With ethanol, there is <u>no visible reaction</u> .	
		With ethanoic acid, <u>effervescence</u> is observed.	
		test 2 : To 5 cm <sup>3</sup> of sample, add an equal volume of <u>acidified</u>	[1]
		potasisum manganate(VII).	
		observation : With ethanol, the <u>purple</u> solution turns <u>colourless</u> .	
		With ethanoic acid, no visible reaction.	
A6	(a)	Choose any one:	[1]
		• The addition polymer contains the <u>C-C chain</u> , while the condensation	
		polymer contains the <u>amide</u> or <u>ester</u> linkages.	
		The formation of addition polymore does not involve any loss of small	
		The formation of addition polymers <u>does not involve any loss of small</u>	
		molecules (i.e. 1 product formed). While the formation of condensation	
		molecules (i.e. 1 product formed). While the formation of condensation polymers involves the loss of small molecules like water (i.e. 2	
		<u>molecules</u> (i.e. 1 product formed). While the formation of condensation polymers <u>involves the loss of small molecules</u> like water (i.e. 2 products formed).	
		<ul> <li><u>molecules</u> (i.e. 1 product formed). While the formation of condensation polymers <u>involves the loss of small molecules</u> like water (i.e. 2 products formed).</li> <li>The formation of addition polymers involves the <u>C=C</u> functional group, while the formation of condensation polymers involves monomers with</li> </ul>	
		<ul> <li><u>molecules</u> (i.e. 1 product formed). While the formation of condensation polymers <u>involves the loss of small molecules</u> like water (i.e. 2 products formed).</li> <li>The formation of addition polymers involves the <u>C=C</u> functional group, while the formation of condensation polymers involves monomers with functional groups like <u>-NH<sub>2</sub>, -COOH and -OH</u>.</li> </ul>	
	(b)	<ul> <li><u>molecules</u> (i.e. 1 product formed). While the formation of condensation polymers <u>involves the loss of small molecules</u> like water (i.e. 2 products formed).</li> <li>The formation of addition polymers involves the <u>C=C</u> functional group, while the formation of condensation polymers involves monomers with</li> </ul>	[1]
	(b)	<ul> <li><u>molecules</u> (i.e. 1 product formed). While the formation of condensation polymers <u>involves the loss of small molecules</u> like water (i.e. 2 products formed).</li> <li>The formation of addition polymers involves the <u>C=C</u> functional group, while the formation of condensation polymers involves monomers with functional groups like <u>-NH<sub>2</sub>, -COOH and -OH</u>.</li> </ul>	[1]
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	(b) (c)	<ul> <li><u>molecules</u> (i.e. 1 product formed). While the formation of condensation polymers <u>involves the loss of small molecules</u> like water (i.e. 2 products formed).</li> <li>The formation of addition polymers involves the <u>C=C</u> functional group, while the formation of condensation polymers involves monomers with functional groups like <u>-NH<sub>2</sub>, -COOH and -OH</u>.</li> <li>H H H H H</li> <li>       </li> <li>-C - C - C - C -  </li> <li>     </li> </ul>	[1]
		<ul> <li><u>molecules</u> (i.e. 1 product formed). While the formation of condensation polymers <u>involves the loss of small molecules</u> like water (i.e. 2 products formed).</li> <li>The formation of addition polymers involves the <u>C=C</u> functional group, while the formation of condensation polymers involves monomers with functional groups like <u>-NH<sub>2</sub>, -COOH and -OH</u>.</li> <li>H H H H</li> <li>       </li> <li>-C - C - C - C -</li> <li>     </li> <li>H CH<sub>3</sub> H CH<sub>3</sub></li> </ul>	
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		$\begin{array}{c} \underline{\text{molecules}} \text{ (i.e. 1 product formed). While the formation of condensation} \\ polymers \underline{\text{involves the loss of small molecules}} \text{ like water (i.e. 2} \\ products formed). \end{array}$ $\begin{array}{c} \text{The formation of addition polymers involves the } \underline{C=C} \text{ functional group,} \\ \text{while the formation of condensation polymers involves monomers with} \\ \text{functional groups like } \underline{-NH_2, -COOH \text{ and } -OH}. \\ \text{H}  \text{H}  \text{H}  \text{H}  \text{H} \\            \\ -C-C-C-C-C-C- \\            \\ \text{H}  CH_3 \text{ H}  CH_3 \\ \text{(i)} & O  O \\        \\ \text{H} - \text{N} - \text{C}_6\text{H}_{12} - \text{N} - \text{H}  \text{H} - \text{O} - \text{C} - \text{C}_4\text{H}_8 - \text{C} - \text{O} - \text{H} \\         &   \end{array}$	[2]
		molecules (i.e. 1 product formed). While the formation of condensation polymers involves the loss of small molecules like water (i.e. 2 products formed). • The formation of addition polymers involves the <u>C=C</u> functional group, while the formation of condensation polymers involves monomers with functional groups like <u>-NH<sub>2</sub>, -COOH and -OH</u> . H H H H H           -C - C - C - C - C -           H CH <sub>3</sub> H CH <sub>3</sub> (i) O O       H - N - C <sub>6</sub> H <sub>12</sub> - N - H H - O - C - C <sub>4</sub> H <sub>8</sub> - C - O - H     H H H H (ii) M <sub>r</sub> of C <sub>12</sub> H <sub>22</sub> N <sub>2</sub> O <sub>2</sub> = 226	[2]
		$\begin{array}{c} \underline{\text{molecules}} \text{ (i.e. 1 product formed). While the formation of condensation} \\ polymers involves the loss of small molecules like water (i.e. 2 products formed). \end{array}$ $\begin{array}{c} \text{ The formation of addition polymers involves the } \underline{C=C} \text{ functional group,} \\ \text{while the formation of condensation polymers involves monomers with} \\ \text{functional groups like } \underline{-NH_2, -COOH \text{ and } -OH.} \\ \text{H}  \text{H}  \text{H}  \text{H}  \text{H} \\            \\ -C-C-C-C-C-L \\            \\ \text{H}  \text{CH}_3  \text{H}  \text{CH}_3 \end{array}$ $\begin{array}{c} \text{O}  \text{O} \\      \\ \text{H}  \text{H} \\ \text{H}  \text{H} \end{array}$	[2]

A	1-1	The minimum elements of the table $\frac{1}{2} = 0.500 + 6000 = 0.000 = 0.00000 = 0.00000 = 0.000000 = 0.00000 = 0.00000000$	[4]					
A7	(a)	The pie charts show that there is $35\%$ of iron in the whole Earth, while	[1]					
		there is only <u>6%</u> in the Earth's crust. This implies that <u>29%</u> of the iron is in						
	<b>/</b> b)	the inner core of the Earth.						
	(b)	(i) Si O						
		percentage 46.7 53.3						
		Ar 28 16						
		no. of moles 1.6679 3.3333						
		$\div 1.6679$ 1 2	[2]					
		empirical formula = $\underline{SiO_2}$ ; name = $\underline{Silcon \ dioxide}$	[2] [1]					
		(ii) Potassium oxide is an i <u>onic</u> compound.	ניז					
		A <u>large</u> amount of heat energy is required to overcome the <u>strong</u>						
		electrostatic forces of attraction between oppositely-charged ions.	[1]					
		Silicon dioxide is a <u>giant covalent</u> molecule.	r.1					
		An even larger amount of heat energy is required to overcome the						
		<u>stronger covalent bonds</u> between atoms.						
		For notassium oxide, the ions are hold rigidly in fixed positions in	[1]					
		For potassium oxide, the ions are held rigidly in fixed positions in the solid state. The <u>absence of mobile charged ions</u> prevents the						
		electric current from being carried.						
		For silicon dioxide, all valence electrons are used for bonding. The	[1]					
		absence of mobile charged electrons prevents the electric current						
	(c)	from being carried. <u>Ca<sup>2+</sup></u> , <u>A/<sup>3+</sup></u> and <u>Si<sub>2</sub>O<sub>8</sub><sup>8-</sup></u>						
	(c) (d)	$M_r$ of phenacite = 110						
	(u)	Mr of prichaence = 110						
		mass of silicon in phenacite = $\frac{28}{20} \times 1000 \text{ g} = 255 \text{ g}$ (to 3 s.f.)						
		mass of silicon in phenacite = $\frac{28}{110}$ x 1000 g = $255$ g (to 3 s.f.) [						
		2(9) $(1000  g - 164  g (to 2  g f))$						
		mass of beryllium in phenacite = $\frac{2(9)}{110}$ x 1000 g = <u>164 g</u> (to 3 s.f.)						
		From the above calculations, 1 kg of phenacite contains a larger mass of						
		silicon than beryllium.						
		no. of mol. of silicon = $\frac{254.545g}{28} = 9.09 \text{ mol}$ (to 3 s.f.)						
		20	[4]					
		163.636g	[1]					
		no. of mol. of beryllium = $\frac{163.636g}{9} = \frac{18.2 \text{ mol}}{1000}$ (to 3 s.f.)						
		From the above calculations, 1 kg of phenacite contains a larger number						
		of moles of beryllium atoms than silicon atoms.						
		·						
<b>A8</b>	(a)	A catalyst provides an alternative pathway with a lower activation energy	[1]					
	()	for the reaction to proceed, leading to a higher frequency of effective	[1]					
		<u>collisions</u> between the particles, resulting in a <u>higher rate</u> of reaction.						
	(b)	Comparing experiments 5, 7, 8 and 9, with the same starting temperature						
	. /	of 19°C, Group 1 and 2 metal compounds are less effective as the time	[1]					
		taken for reaction to finish is longer (i.e. NaNO₃ takes <u>45s</u> , while						
		Mg(NO <sub>3</sub> ) <sub>2</sub> takes <u>46 s</u> ), as compared to the transition metal compounds	[1]					
		which take a shorter time (i.e. $Fe(NO_3)_3$ takes <u>15s</u> , and $Co(NO_3)_2$ takes	_					
		<u>26s</u> .						
	(c)	(i) <u>Fe<sup>2+</sup></u> and <u>Fe<sup>3+</sup></u>	[1]					

	(ii)	<u>Iron(III) ion</u> is the more effective catalyst. Comparing experiments 6 and 7, iron(III) ion took a <u>shorter</u> time (i.e. <u>15s</u> in experiment 7) to complete the reaction, as compared to that of iron(II) ion (i.e. <u>22s</u> in experiment 6), despite the slight change in temperature.	[1]
(d)	(iii) (i)	Transition metals form <u>coloured compounds</u> when hydrated. Yes.	[1]
		Comparing experiments 1 and 5, both NaC <i>l</i> and NaNO <sub>3</sub> took <u>45s</u> to complete the reaction, regardless of the anion. Also, comparing experiments 2 and 6, both FeC <i>l</i> <sub>2</sub> and Fe(NO <sub>3</sub> ) <sub>2</sub> took <u>22s</u> to complete the reaction, regardless of the anion.	[2]
	(ii)	<u>15s</u>	[1]

## Section B [10 m]

	-			
B9	(a)	(i)	From the test with aqueous NaOH, the white precipitate formed is	[1]
			soluble in excess, implying that both $AI^{3+}$ ions and $Zn^{2+}$ ions are	543
			present. If Ca <sup>2+</sup> ion is present, then some white precipitate formed	[1]
			is <u>insoluble</u> in excess.	
			From the test with aqueous NH <sub>4</sub> OH, some white precipitate	[1]
			formed is <u>soluble</u> in excess, confirming that Zn <sup>2+</sup> ions are present,	
			while some white precipitate formed is <u>insoluble</u> in excess,	
			confirming that A <sup>/3+</sup> ions are present.	
		(ii)	No, the ammonia gas produced could be due to the presence of	[1]
			<u>aqueous ammonia</u> .	
			To test for nitrate ions, the scientist should use warm aqueous	[1]
			sodium hydroxide and aluminium foil.	
	(b)	test	1 7 1	[1]
			dilute acid, followed by aqueous silver nitrate.	
		resu		[1]
	(c)		dd excess zinc carbonate [base] powder to warm <u>dilute hydrochloric</u>	[1]
			<u>cid</u> [acid].	
			ilter the mixture to remove the excess zinc chloride.	541
			eat the filtrate till it is saturated.	[1]
			llow the solution to <u>cool</u> and crystals of zinc chloride will form.	
			ilter off the crystals.	[4]
			<u>/ash</u> the crystals with distilled water.	[1]
		<u>Dry</u> th	he pieces of crystals between two pieces of filter paper.	
B10	(a)	(i)	W > Z > Y > X	[1]
			Effervescence is observed.	[1]
			Metal <b>Z</b> , being a <u>reactive</u> metal (i.e. above H in the reactivity	[1]
			series), reacts with acid to displace hydrogen gas.	
			Approach:	[1]
			The student measured and compared the volume of hydrogen gas	
			collected for a metal of fixed mass (i.e. 0.5 g each of metals W, X,	
			<b>Y</b> , <b>Z</b> and <b>M</b> ) to react completely with <u>dilute hydrochloric acid</u> of a	
			fixed concentration (i.e. 1 mol/dm <sup>3</sup> )-and volume (30 cm <sup>3</sup> ) over a 2-	
			minute duration.	

		Procedure:	[1]
		1. Using a 50 cm <sup>3</sup> measuring cylinder, measure 30 cm <sup>3</sup> of HC <i>l</i> and	[,1]
		place it in a 250 cm <sup>3</sup> conical flask.	
		2. Transfer the dilute hydrochloric acid into the conical flask,	
		connect a gas syringe	
		3. Using an electronic mass balance, measure 0.5 g of metal <b>W</b>	
		and add it to the conical flask.	
		Stopper the conical flask and connect a gas syringe to the	
		conical flask using a delivery tube. Start the stopwatch immediately.	
		4. Stop the stopwatch when it reaches 2 minutes.	
		Record the volume of gas produced in a table of suitable format.	
		Conclusion:	[1]
		Metal <b>W</b> will have the highest volume of hydrogen gas collected, followed by metals <b>Z</b> , <b>Y</b> , <u><b>M</b></u> (i.e. 4 <sup>th</sup> most reactive) and <b>X</b> .	
(b)	(i)	There is no visible reaction for experiments 1 and 3.	[1]
(6)	(1)	Calcium and lead, being more reactive than hydrogen, cannot be	[,1]
		displaced from their oxide by hydrogen.	
		In experiment 2, the red copper oxide turns red-brown.	[1]
		Copper, being less reactive than hydrogen, can be displaced from	
	<i></i>	its oxide by hydrogen.	
	(ii)	The tube containing <u>copper oxide</u> .	[1]
		Copper, being the <u>least reactive metal</u> , forms the <u>least stable metal</u> <u>oxide</u> , hence the oxide is <u>more readily reduced</u> by hydrogen.	
		onde, mence the onde is more readily reduced by Hydrogen.	