

**NATIONAL JUNIOR COLLEGE**  
**PRELIMINARY EXAMINATIONS**  
Higher 1

CANDIDATE  
NAME

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SUBJECT  
CLASS

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REGISTRATION  
NUMBER

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**CHEMISTRY**

Paper 2 Structured Questions

**8872/02**

**17 September 2009**

**2 hours**

<p><b>READ THE INSTRUCTION FIRST</b> Write your subject class, registration number and name on all the work you hand in. Write in dark blue or black pen on both sides of the paper. You may use a soft pencil for any diagrams, graphs or rough working. Do not use paper clips, highlighters, glue or correction fluid.</p> <p><b>Section A</b> Answer <b>all</b> questions .</p> <p><b>Section B</b> Answer any <b>two</b> questions.</p> <p>At the end of the examination, fasten all your work securely together. The number of marks is given in brackets [ ] at the end of each question or part question.</p>	<b>For Examiner's Use</b>	
	<b>Section A</b>	
	<b>1</b>	
	<b>2</b>	
	<b>3</b>	
	<b>4</b>	
	<b>5</b>	
	<b>Section B</b>	
	<b>B6</b>	
	<b>B7</b>	
	<b>B8</b>	
	<b>Total</b>	

**Section A**

Answer **all** the questions in this section in the spaces provided.

- 1 5.00 g of impure iron solid was dissolved in an excess of sulfuric acid and the resulting solution containing  $\text{Fe}^{2+}$  was made up to  $250 \text{ cm}^3$ .  $10.0 \text{ cm}^3$  of this solution was titrated against a solution of  $\text{KMnO}_4$  of concentration  $0.01 \text{ mol dm}^{-3}$ . It was found that  $24.00 \text{ cm}^3$  of  $\text{KMnO}_4$  was required.

- (a) Suggest a reason why an indicator was not required for this titration.

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[1]

- (b) Write a balanced redox equation for the reaction between  $\text{Fe}^{2+}$  and  $\text{MnO}_4^-$ .

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[2]

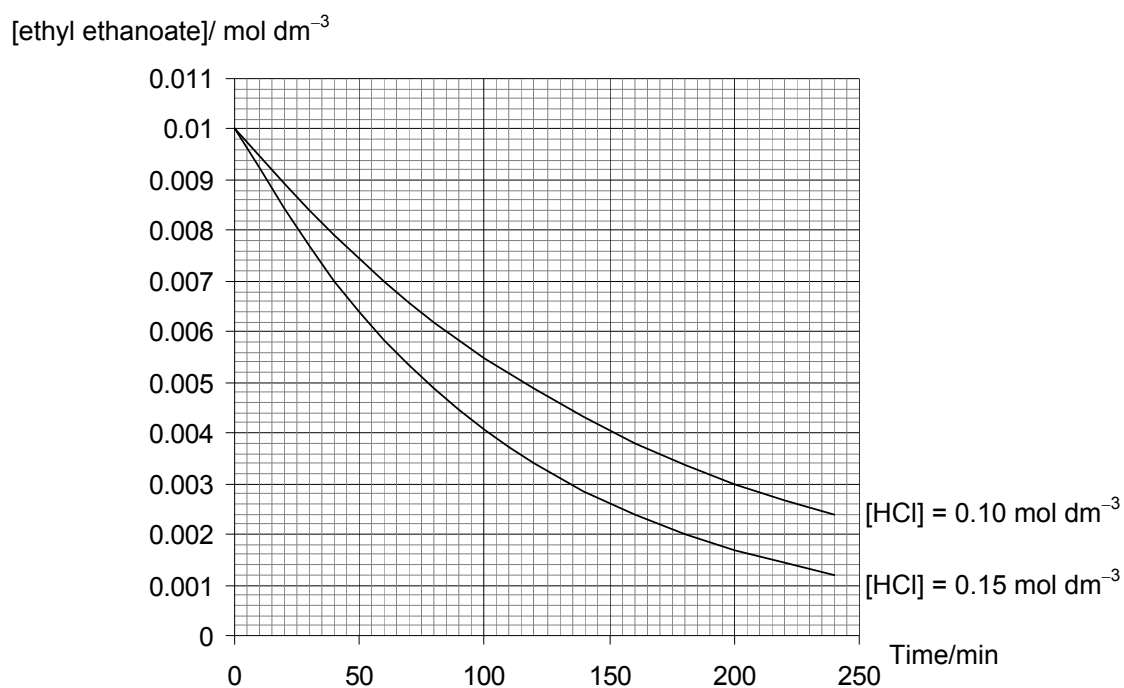
- (c) A student claims that the percentage purity of iron in the sample is 33.5%. Justify if the statement made by the student is true.

[3]

[Total: 6]

**[Turn Over**

- 2 The graphs below shows the rate of hydrolysis of ethyl ethanoate,  $\text{CH}_3\text{COOC}_2\text{H}_5$  in an acidic medium. The reaction was followed twice with different concentrations of HCl and the following results were obtained.



- (a) Using the graphs, determine the order of reaction with respect to both ethylethanoate and HCl.
- (b) Deduce the rate equation and calculate the value of the rate constant, giving its units.
- (c) On the same axis above, sketch the graph when the experiment is repeated using  $0.008 \text{ mol dm}^{-3}$  of ethylethanoate and  $0.15 \text{ mol dm}^{-3}$  HCl.

[Total: 7]

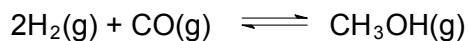
[Turn Over]

3 Methanol is a possible alternative to hydrocarbons as a liquid fuel.

- (a) Given that the enthalpy of combustion of methanol is  $-715 \text{ kJ mol}^{-1}$ , calculate the mass of methanol that should be burnt in order to boil  $1000 \text{ cm}^3$  of water starting from an initial temperature of  $20.0^\circ \text{C}$ . Assume that 50% of the heat obtained from the combustion of methanol is lost to the surroundings and that the specific heat capacity of water is  $4.2 \text{ J K}^{-1} \text{ g}^{-1}$ .

[3]

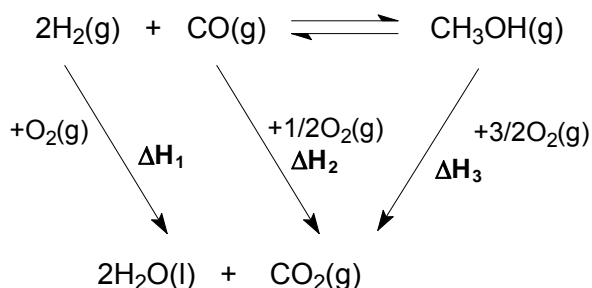
- (b) Methanol can be produced by using a reversible reaction between carbon monoxide and hydrogen.



- (i) Given the following standard enthalpy change of combustion and energy cycle, apply Hess's Law and calculate the enthalpy change for the production of methanol.

$$\Delta H_c^\ominus(\text{H}_2) = -286 \text{ kJ mol}^{-1}$$

$$\Delta H_c^\ominus(\text{CO}) = -283.3 \text{ kJ mol}^{-1}$$



[2]

[Turn Over]

- (ii) When 2.00 mol of hydrogen and 1.00 mol of carbon monoxide were mixed and heated to a high temperature in a container of volume  $1.50 \text{ dm}^3$ , 0.80 mol of methanol was formed at equilibrium.

Calculate a value for the equilibrium constant,  $K_c$ , for this reaction at this temperature and give its units.

- (iii) State and explain the effect on the position of equilibrium for the above reaction when: [3]
- (I) the pressure is increased.
- (II) the temperature is increased.

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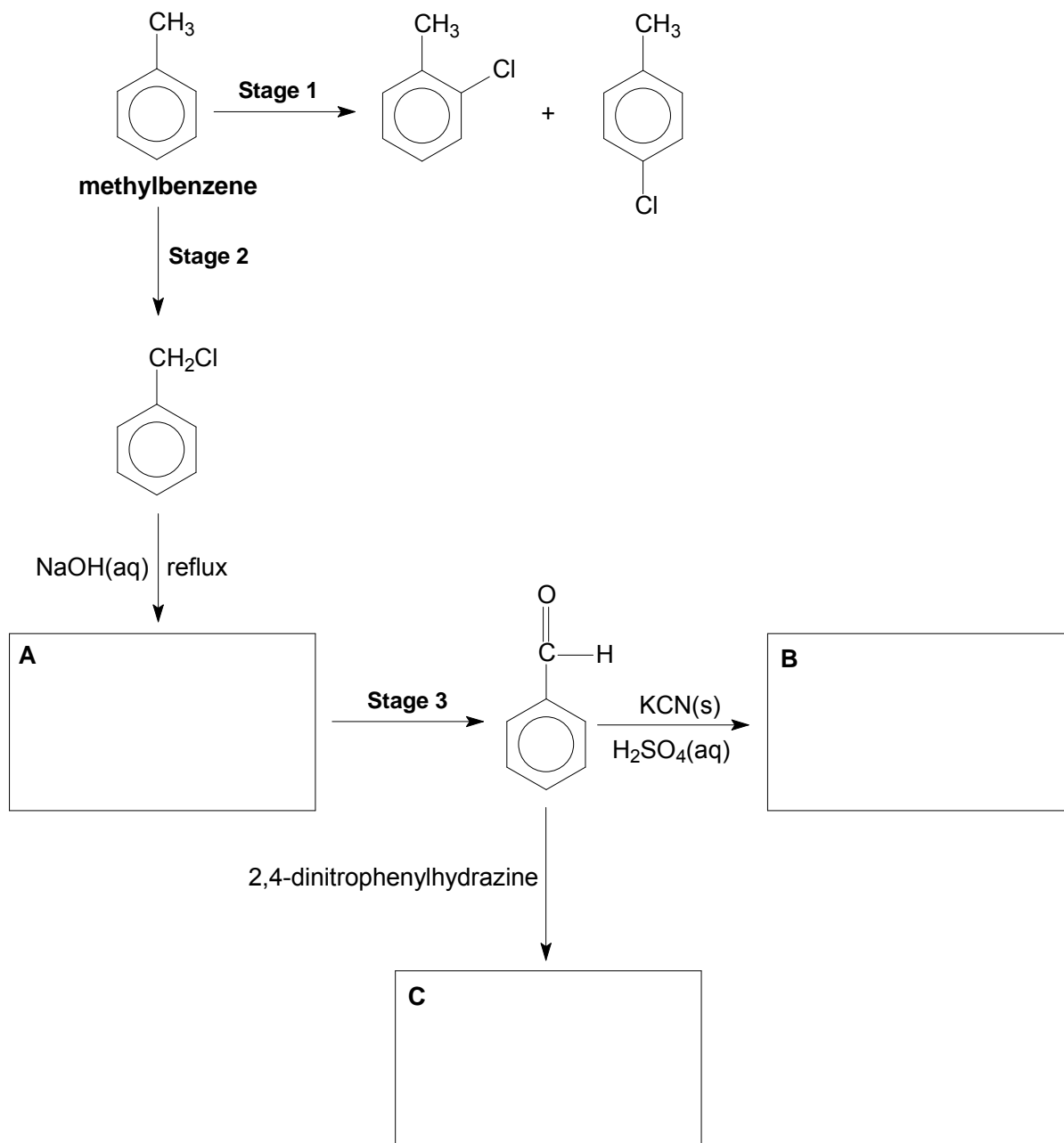
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[3]

[Total: 11]

- 4 The diagram below shows a reaction scheme starting from methylbenzene.



- (a) Draw the structures for Compounds **A–C** in the boxes provided above.

[3]

- (b) Suggest the reagents and conditions required for **Stages 1–3**.

Stage	Reagents and Conditions
1	
2	
3	

[3]

[Total: 6]

- 5 This question is about the extraction of iron from its ore,  $\text{Fe}_2\text{O}_3$  (also known as haematite) using two different methods. The first method of extraction is by using the **Blast Furnace**. The second method is through a process known as the **Thermite Reaction**.

#### Extraction of iron using the Blast Furnace.

Ninety percent of all mining of metallic ores is for the extraction of iron. Industrially, iron is produced starting from iron ores, principally hematite, by reduction with carbon in a blast furnace at temperatures of about 2000 °C.

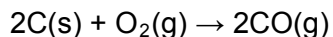
The following table shows the composition of haematite.

Compound	Percentage Composition
$\text{Fe}_2\text{O}_3$	92.0 %
$\text{SiO}_2$	6.0 %
Others	2.0 %

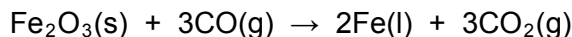
Haematite, carbon in the form of coke, and limestone are continuously fed into the top of the furnace, while a blast of heated air is forced into the furnace at the bottom.

In the blast furnace, the following reactions take place.

**Stage 1:** The coke reacts with oxygen in the air blast to produce carbon monoxide:



**Stage 2:** The carbon monoxide formed then react with the iron ore to form molten iron, becoming carbon dioxide in the process:

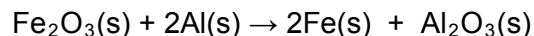


The main impurity in the ore is principally silicon dioxide,  $\text{SiO}_2$ . It is solid at the temperature of the furnace and the furnace would become blocked if it was not removed.

Limestone, which is principally calcium carbonate,  $\text{CaCO}_3$  is decomposed in heat to form calcium oxide and carbon dioxide. Silicon dioxide then reacts with calcium oxide to form calcium silicate called slag which is a liquid. Slag flows to the bottom of the furnace where it floats on the liquid iron and is easily removed. The slag is allowed to cool until it becomes a solid and it is used for road construction.

#### Extraction of iron using the Thermite Reaction

The Thermite reaction is a batch process in which powdered aluminium reacts with iron(III) oxide. The aluminium is able to reduce the iron oxide to elemental iron because aluminium is highly combustible:



The products are aluminium oxide and free elemental iron.



**With reference to the extraction of iron using the Blast Furnace:**

- (a)** State with a reason which compound is the reducing agent in **Stage 2**.

.....  
.....

[2]

- (b)** If 100 kg of haematite is continuously fed into the blast furnace, calculate the mass of iron that can be obtained at the end of the reaction.

[2]

- (c)** Explain with the aid of an equation, how the impurity silicon dioxide is removed from the blast furnace.

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.....  
.....

[2]

**(d) With reference to the extraction of iron using the Thermite Reaction:**

The table below shows the values for the enthalpy of formation of  $\text{Fe}_2\text{O}_3$  and  $\text{Al}_2\text{O}_3$ :

Compound	(Hf / $\text{kJ mol}^{-1}$ )
$\text{Fe}_2\text{O}_3$	-824.2
$\text{Al}_2\text{O}_3$	-1676

Why are the values for the enthalpy of formation of Al and  $\text{O}_2$  not given in the table?

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[1]

**(e)** Using the data above, show by means of calculation whether the Thermite Reaction is an endothermic or exothermic reaction.

[2]

**(f)** Give one advantage of producing iron in the Blast Furnace rather than in the Thermite Reaction.

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[1]

[Total: 10]

## Section B

Answer **two** of the following three questions. Answer these questions on separate answer paper.

- 1 The table below shows some common reagents found in a chemistry laboratory.

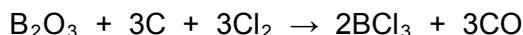
HCl		Ba(OH) <sub>2</sub>
	HNO <sub>3</sub>	
NaCl		NH <sub>3</sub>

- (a) (i) Define the terms Bronsted Acid and Bronsted Base.
- (ii) From the list of compounds given above, choose one Bronsted Acid and one Bronsted Base.
- (iii) Write an equation to show how the Bronsted acid that you have chosen in (a)(ii) function as an acid in presence of water. Indicate which species is the acid and base as well as their respective conjugate acid and base in the equation. [6]
- (b) (i) Calculate the pH of a 25.0 cm<sup>3</sup> Ba(OH)<sub>2</sub> solution of concentration 0.05 mol dm<sup>-3</sup>.
- (ii) What is the final pH when 5 cm<sup>3</sup> of HNO<sub>3</sub> of concentration 0.05 mol dm<sup>-3</sup> is added to the solution in (b)(i)? [4]
- (c) When a weak base NH<sub>3</sub> is partially neutralized by HNO<sub>3</sub>, solution **X** is obtained. Adding a small amount of either acid or base to solution **X** does not change the pH of the solution significantly.
- (i) State the type of solution formed when NH<sub>3</sub> is partially neutralized by HNO<sub>3</sub>.
- (ii) Explain with the aid of equations why the pH of the solution **X** does not change significantly upon addition of a small amount of either acid or base. [3]
- (d) When gaseous HCl is added to 3-methylpent-2-ene, CH<sub>3</sub>CH=C(CH<sub>3</sub>)CH<sub>2</sub>CH<sub>3</sub> a mixture of two products is formed.
- (i) State the type of isomerism exhibited by 3-methylpent-2-ene.
- (ii) Suggest the structural features present in 3-methylpent-2-ene that enables it to exhibit the type of isomerism suggested in (d)(i).
- (iii) Draw the structure of an isomer of 3-methylpent-2-ene which will not exhibit the type of isomerism suggested in (d)(i).
- (iv) Draw the structures of the two products formed when gaseous HCl is added to 3-methylpent-2-ene. Indicate which is the major and which is the minor product. [7]

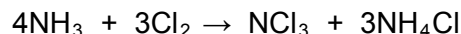
[Total: 20]

[Turn Over

- 2 Boron trichloride,  $\text{BCl}_3$  is produced industrially by direct chlorination of boron oxide and carbon at  $500^\circ\text{C}$ .



Nitrogen trichloride,  $\text{NCl}_3$  on the other hand is formed when ammonia reacts with chlorine gas.



- (a) (i) With the aid of dot-cross diagrams, explain why the shape of both  $\text{BCl}_3$  and  $\text{NCl}_3$  are different from each other. [4]
- (ii) Hydrogen sulphide,  $\text{H}_2\text{S}$ , reacts with boron trichloride,  $\text{BCl}_3$  in a 1:1 ratio to form Compound **D**. [4]
- (l) Draw the structure of Compound **D**. Hence explain why  $\text{BCl}_3$  would react with  $\text{H}_2\text{S}$  in a 1:1 ratio.
- (II) Predict the value of the H-S-H bond angle in Compound **D**. [4]
- (iii) Both nitrogen and phosphorus come from Group V.  $\text{NCl}_3$  does not react with more chlorine gas to form  $\text{NCl}_5$  while  $\text{PCl}_3$  is able to react with more chlorine gas to form  $\text{PCl}_5$ . Explain why this is so. [2]
- (iv) The table below shows the boiling point of three compounds. With reference to the structure and bonding present in the compounds, suggest explanation for the differences in their boiling point.

Compound	Boiling Point / $^\circ\text{C}$
$\text{NH}_3$	- 33
$\text{NCl}_3$	71
$\text{NH}_4\text{Cl}$	520

- (b) Describe, with the aid of equations, what happens when Universal Indicator is added to separate samples of  $\text{NaCl}$  and  $\text{PCl}_5$  in water. [3]
- (c) An organic Compound **E** has the empirical formula  $\text{CH}_2\text{O}$ . Compound **E** undergoes the following reactions. [4]
- It gives brisk effervescence with aqueous sodium carbonate.
  - It forms a yellow precipitate when warmed with aqueous alkaline iodine.
- (i) Draw a structure for Compound **E** which fits the above descriptions.
- (ii) When Compound **E** is warmed with acidified potassium dichromate(VI), a Compound **F** is formed. Suggest a structure for Compound **F** and hence state the expected observation. [3]

[Total: 20]

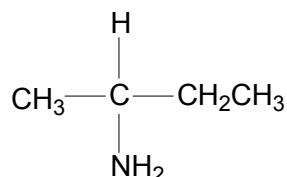
- 3 Butene is a colourless gas present in crude oil as a minor constituent in quantities that are too small for viable extraction.

Butene can be formed from either butan-2-ol,  $\text{CH}_3\text{CH}(\text{OH})\text{CH}_2\text{CH}_3$  or 2-chlorobutane,  $\text{CH}_3\text{CH}(\text{Cl})\text{CH}_2\text{CH}_3$ .

- (a) (i) Illustrate the type of bonding present between butan-2-ol and water with the aid of a suitable diagram.
- (ii) Suggest the reagents and conditions necessary to convert both butan-2-ol and 2-chlorobutane into butene.
- (iii) Give the structures of the three alkenes formed.
- (iv) Suggest the reagents and conditions necessary to distinguish one of the isomers from the other two. In each case, draw the structures of **ALL** the products formed and state the expected observations.

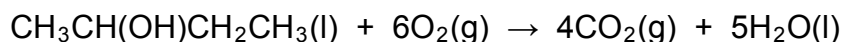
[12]

- (b) Using butan-2-ol as a starting material, suggest a two-step synthesis in which the following compound can be made.



[3]

- (c) Butan-2-ol burns in excess oxygen to give a mixture of carbon dioxide gas and water as shown by the equation below.



- (i) Define the term standard bond energy using oxygen gas as an example.
- (ii) Using the bond energy values from the *Data Booklet*, calculate the enthalpy change for the above reaction.
- (iii) The bond energy values given in the *Data Booklet* are merely average values. Suggest another reason why the value that you have calculated from (c)(ii) differs from the actual value.

[5]

[Total: 20]

[Turn Over]

