



# CHEMISTRY HIGHER LEVEL PAPER 2

Monday 9 May 2011 (afternoon)

2 hours 15 minutes

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#### Examination code

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#### **INSTRUCTIONS TO CANDIDATES**

- Write your session number in the boxes above.
- Do not open this examination paper until instructed to do so.
- Section A: answer all questions.
- Section B: answer two questions.
- Write your answers in the boxes provided.

Please **do not** write on this page.

Answers written on this page will not be marked.



### **SECTION A**

Answer all questions. Write your answers in the boxes provided.

1. Methanol is made in large quantities as it is used in the production of polymers and in fuels. The enthalpy of combustion of methanol can be determined theoretically or experimentally.

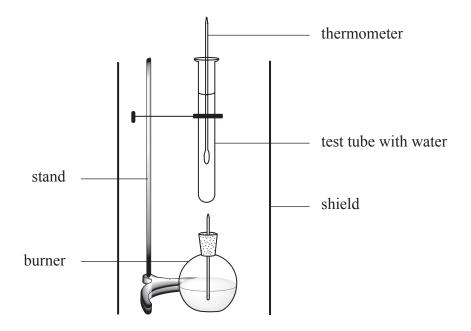
$$\operatorname{CH_3OH}(1) + 1\tfrac{1}{2}\operatorname{O}_2(\mathsf{g}) \to \operatorname{CO}_2(\mathsf{g}) + 2\operatorname{H}_2\operatorname{O}(\mathsf{g})$$

	CH <sub>3</sub> OH(l)	$O_2(g)$	$CO_2(g)$	$H_2O(g)$
Standard enthalpy of formation, $\Delta H_{\rm f}^{\Theta}/{\rm kJ \ mol^{-1}}$	-239	0	-394	-242
Entropy, $S^{\ominus}/J$ $K^{-1}$ mol <sup>-1</sup>	240	205	214	189

(a)	Using the information from the table above, determine the theoretical enthalpy of combustion of methanol.	[2]



(b) The enthalpy of combustion of methanol can also be determined experimentally in a school laboratory. A burner containing methanol was weighed and used to heat water in a test tube as illustrated below.



The following data were collected.

Initial mass of burner and methanol / g	80.557
Final mass of burner and methanol / g	80.034
Mass of water in test tube / g	20.000
Initial temperature of water /°C	21.5
Final temperature of water /°C	26.4


(This question continues on the following page)



*[*2

(ii)	Calculate the heat absorbed, in kJ, by the water.	[3]
(iii)	Determine the enthalpy change, in kJ mol <sup>-1</sup> , for the combustion of methanol.	[2]



(i)	Part (a)
(ii)	Part (b)
Det	ermine the $\Delta S^{\ominus}$ for the combustion of methanol.
	$CH_3OH(1) + 1\frac{1}{2}O_2(g) \rightarrow CO_2(g) + 2H_2O(g)$

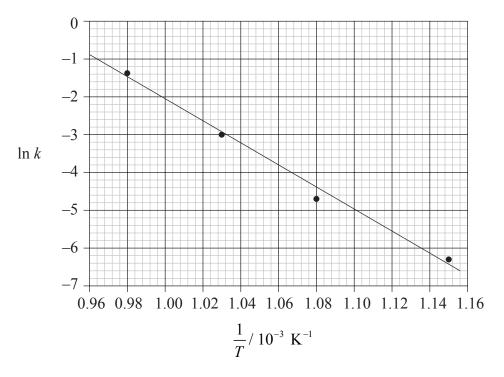


Using the enthalpy of combustion for methanol from Table 12 of the Data Booklet and the $\Delta S^{\ominus}$ determined in part (d), calculate the standard free energy change for the combustion of methanol.	[3]
Explain whether changing the temperature will alter the spontaneity of the reaction.	[1]
	the $\Delta S^{\ominus}$ determined in part (d), calculate the standard free energy change for the combustion of methanol.



2. Consider the following graph of  $\ln k$  against  $\frac{1}{T}$  (temperature in Kelvin) for the second order decomposition of  $N_2O$  into  $N_2$  and O.

$${\rm N_2O} \rightarrow {\rm N_2+O}$$



(a) State how the rate constant, k varies with temperature, T.

[1]


(b) Determine the activation energy,  $E_a$ , for this reaction.

[3]

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[2]

(Question 2 continued)

(c)	The r	ate	expression	for	this	reaction	is	$rate = k [N_2O]^2$	and	the	rate	constant	is
	0.244	$dm^3$	$mol^{-1} s^{-1} at$	750°	°C.			_					

A sample of  $\rm N_2O$  of concentration 0.200 mol  $\rm dm^{-3}$  is allowed to decompose. Calculate the rate when 10% of the  $\rm N_2O$  has reacted.

 	 ٠.

(a)	Explain why the relative atomic mass of cobalt is greater than the relative atomic mass of nickel, even though the atomic number of nickel is greater than the atomic number of cobalt.	[1]
(b)	Deduce the numbers of protons and electrons in the ion Co <sup>2+</sup> .	[1]
(c)	Deduce the electron configuration for the ion Co <sup>2+</sup> .	[1]
(d)	Identify a radioactive isotope of cobalt <b>and</b> state <b>one</b> of its uses.	[1]



4.	Methanol	may	be	produced	by	the	exothermic	reaction	of	carbon	monoxide	gas	and
	hydrogen	gas.											

$$CO(g) + 2H_2(g) \rightleftharpoons CH_3OH(g)$$
  $\Delta H^{\ominus} = -103 \text{ kJ}$ 

State the equilibrium constant expression, $K_c$ , for the production of methanol.	
_	State the equilibrium constant expression, $K_c$ , for the production of methanol.

(b) State and explain the effect of changing the following conditions on the amount of methanol present at equilibrium:

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(ii) increasing the pressure of the reaction at constant temperature. [2]

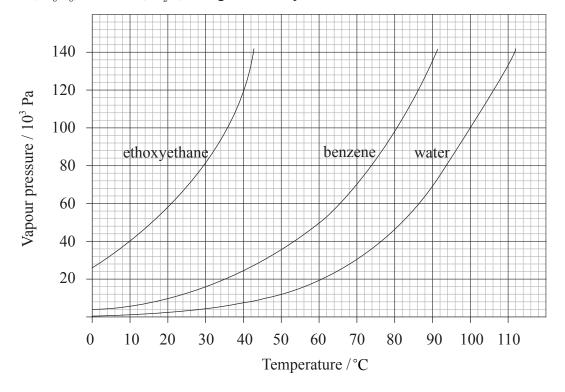
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(c)	The conditions used in industry during the production of methanol are a temperature of 450 °C and pressure of up to 220 atm. Explain why these conditions are used rather than those that could give an even greater amount of methanol.	[2]
(d)	A catalyst of copper mixed with zinc oxide and alumina is used in industry for this production of methanol. Explain the function of the catalyst.	[1]

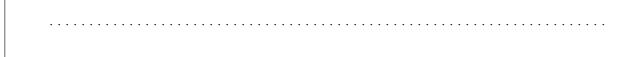


5. The graph below illustrates how the vapour pressures of ethoxyethane, CH<sub>3</sub>CH<sub>2</sub>OCH<sub>2</sub>CH<sub>3</sub>, benzene, C<sub>6</sub>H<sub>6</sub>, and water, H<sub>2</sub>O, change with temperature.



(a) Using data from the graph, explain the difference in vapour pressure of ethoxyethane, benzene and water at  $30\,^{\circ}$ C. [4]


(b) Use the graph to determine the boiling point of benzene at standard pressure. [1]





## **SECTION B**

Answer two questions. Write your answers in the boxes provided.

**6.** 

6.	Carb	on and silicon belong to the same group of the periodic table.
	(a)	Describe and compare <b>three</b> features of the structure and bonding in the three allotropes

	arbon: diamond, graphite and $C_{60}$ fullerene.
Botl	n silicon and carbon form oxides.
Both (i)	n silicon and carbon form oxides. Describe the structure and bonding in $SiO_2$ .



(	ii) Explain why silicon dioxide is a solid and carbon dioxide is a gas at room temperature.	
I	Describe the bonding within the carbon monoxide molecule.	
	Describe the delocalization of pi $(\pi)$ electrons and explain how this can account for the structure and stability of the carbonate ion, $CO_3^{2-}$ .	

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Turn over

e)	Explain the meaning of the term <i>hybridization</i> . State the type of hybridization shown by the carbon atoms in carbon dioxide, diamond, graphite and the carbonate ion.	[5]
•		
f)	(i) Explain the electrical conductivity of molten sodium oxide and liquid sulfur trioxide.	[2]



(ii)	Samples of sodium oxide and solid sulfur trioxide are added to separate beakers of water. Deduce the equation for each reaction and predict the electrical conductivity	
	of each of the solutions formed.	[3]

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[5]

7. (a) Two compounds, **A** and **D**, each have the formula  $C_aH_oCl$ .

Compound **A** is reacted with dilute aqueous sodium hydroxide to produce compound **B** with a formula of  $C_4H_{10}O$ . Compound **B** is then oxidized with acidified potassium manganate(VII) to produce compound **C** with a formula of  $C_4H_8O$ . Compound **C** resists further oxidation by acidified potassium manganate(VII).

Compound **D** is reacted with dilute aqueous sodium hydroxide to produce compound **E** with a formula of  $C_4H_{10}O$ . Compound **E** does not react with acidified potassium manganate(VII).

Deduce the structural formulas for compounds A, B, C, D and E.

A	<b>\:</b>			
В	3 <b>:</b>			
C	<b>:</b>			
D	<b>)</b> :			
E	::			

(This question continues on the following page)



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(b)	Deduce an equation for the reaction between propanoic acid and methanol. Identify the catalyst and state the name of the organic compound, <b>X</b> , formed.	[4]
(c)	In an experiment conducted at 25.0 °C, the initial concentration of propanoic acid and methanol were 1.6 mol dm <sup>-3</sup> and 2.0 mol dm <sup>-3</sup> respectively. Once equilibrium was established, a sample of the mixture was removed and analysed. It was found to contain 0.80 mol dm <sup>-3</sup> of compound <b>X</b> .  (i) Calculate the concentrations of the other three species present at equilibrium.	[3]



(Question 7 continue	ea)	
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	(ii)	State the equilibrium constant expression, $K_c$ , and calculate the equilibrium constant for this reaction at 25.0 °C.	[2]
(d)	(i)	2-chloro-3-methylbutane reacts with sodium hydroxide via an $S_{\rm N}2$ mechanism. Explain the mechanism by using curly arrows to represent the movement of electron pairs.	[4
	(ii)	Explain why the hydroxide ion is a better nucleophile than water.	[2

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(iii) 1-chlorobutane can be converted to a pentylamine via a two stage process.

(Question 7 continued)

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8.

(i)	Define the terms <i>acid</i> and <i>base</i> according to the Brønsted-Lowry theory. Distinguish between a weak base and a strong base. State <b>one</b> example of a weak base.	I
(ii)	Weak acids in the environment may cause damage. Identify a weak acid in the environment <b>and</b> outline <b>one</b> of its effects.	
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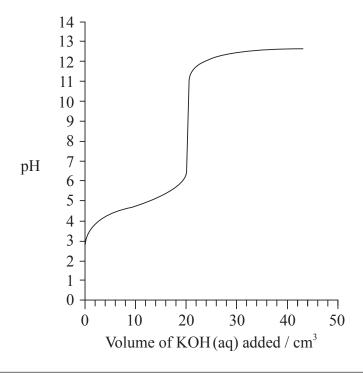
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(iii) The graph below indicates the pH change during the titration of 20.0 cm<sup>3</sup> of 0.100 mol dm<sup>-3</sup> of CH<sub>3</sub>COOH(aq) with 0.100 mol dm<sup>-3</sup> KOH(aq). From the graph, identify the volume of KOH(aq) and the pH at the equivalence point.

[2]




(iv)	Explain how the graph could be used to determine the $pK_a$ of ethanoic acid and	
	determine the $pK_a$ value for these data.	[2]




	the pH at the equivalence point and the final pH reached.
i)	Describe how an indicator works.
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i)	Describe how an indicator works.

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Explain, using an equation, whether a solution of 0.10 mol dm <sup>-3</sup> FeCl <sub>3</sub> (aq) would be acidic, alkaline or neutral.	
Determine the pH of the solution resulting when 100 cm <sup>3</sup> of 0.50 mol dm <sup>-3</sup> HCl(aq) is mixed with 200 cm <sup>3</sup> of 0.10 mol dm <sup>-3</sup> NaOH(aq).	
Determine the pH of the solution resulting when 100 cm <sup>3</sup> of 0.50 mol dm <sup>-3</sup> HCl(aq) is mixed with 200 cm <sup>3</sup> of 0.10 mol dm <sup>-3</sup> NaOH(aq).	
mixed with 200 cm <sup>3</sup> of 0.10 mol dm <sup>-3</sup> NaOH(aq).	
Determine the pH of the solution resulting when 100 cm³ of 0.50 mol dm⁻³ HCl(aq) is mixed with 200 cm³ of 0.10 mol dm⁻³ NaOH(aq).	
mixed with 200 cm <sup>3</sup> of 0.10 mol dm <sup>-3</sup> NaOH(aq).	
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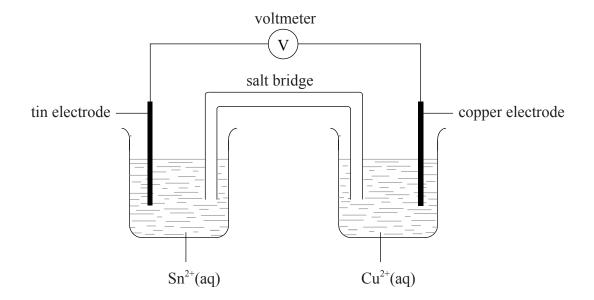


9.

	Draw an electrolytic cell illustrating the electrolysis of molten nickel(II) bromide, NiBr <sub>2</sub> . Include in the diagram the direction of the electron flow, the polarity of	
	electrodes and state the half-equations for the product formed at each electrode.	[5]
(ii)	Deduce the equations for the formation of the major product at the positive electrode (anode) when the following aqueous solutions are electrolysed.	
	<ul><li> dilute sodium chloride</li><li> concentrated sodium chloride</li></ul>	[2



(b) A voltaic cell is constructed from two half-cells as illustrated below.



(1)	reaction occurring in this cell.	[1]
(ii)	Calculate the standard potential for this cell.	[1]
(iii)	State the conditions necessary for the potential of the cell to equal that calculated in part (b) (ii) using the data from Table 14.	[1]



[2]

[3]

(Question 9 continued)

(c)	Using the data below and data from Table 14 of the Data Booklet, predict and explain
	which metal, cadmium or chromium, may be obtained by electrolysis of separate aqueous
	solutions of Cd <sup>2+</sup> (aq) ions and Cr <sup>2+</sup> (aq) ions.

 $E^{\ominus} / V$   $Cd^{2+}(aq) + 2e^{-} \rightarrow Cd(s) \qquad -0.40$   $Cr^{2+}(aq) + 2e^{-} \rightarrow Cr(s) \qquad -0.91$ 


(d) (i) Electrolysis is used in the electroplating of metals. The same amount of current is passed through separate aqueous solutions of  $NiSO_4$ ,  $Sn(SO_4)_2$  and  $Cr_2(SO_4)_3$  in separate electrolytic cells for the same amount of time. State and explain which cell would deposit the greatest amount (in mol) of metal. Identify the electrode at which the metal is deposited.




Nitrogen monoxide may be removed from industrial emissions via a reaction with ammonia as shown by the equation below. $4NH_3(g)+6NO(g) \rightarrow 5N_2(g)+6H_2O(l)$ (i) Deduce the oxidation number of the nitrogen in the reactants and product. [3]	(ii)	For the Sn(SO <sub>4</sub> ) <sub>2</sub> cell, suggest <b>two</b> factors, other than time and current, that would affect the amount of metal deposited during electroplating.	[2]
Nitrogen monoxide may be removed from industrial emissions via a reaction with ammonia as shown by the equation below. $4NH_3(g) + 6NO(g) \rightarrow 5N_2(g) + 6H_2O(l)$			
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	.mm	nonia as shown by the equation below.	
		$4NH_3(g) + 6NO(g) \rightarrow 5N_2(g) + 6H_2O(l)$	[3]
		$4NH_3(g) + 6NO(g) \rightarrow 5N_2(g) + 6H_2O(l)$	[3]
		$4NH_3(g) + 6NO(g) \rightarrow 5N_2(g) + 6H_2O(l)$	[3]
		$4NH_3(g) + 6NO(g) \rightarrow 5N_2(g) + 6H_2O(l)$	[3]
		$4NH_3(g) + 6NO(g) \rightarrow 5N_2(g) + 6H_2O(l)$	[3]
		$4NH_3(g) + 6NO(g) \rightarrow 5N_2(g) + 6H_2O(l)$	[3]
		$4NH_3(g) + 6NO(g) \rightarrow 5N_2(g) + 6H_2O(l)$	[3]
		$4NH_3(g) + 6NO(g) \rightarrow 5N_2(g) + 6H_2O(l)$	[3]



Deduce the oxidation and reduction half-equations and identify the oxidizing agent for the reaction.
30.0 dm³ of ammonia reacts with 30.0 dm³ of nitrogen monoxide at 100 °C.
Identify which gas is in excess and by how much and calculate the volume of nitrogen produced.



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