

## **CHEMISTRY 3.3 Paper 1**

Describe oxidation-reduction processes

Credits: Three

### **INSTRUCTIONS**

Answer **ALL** questions

You are advised to spend 30 minutes answering these questions.

**Question One (Bursary 2003 Question 9: modified)**

The table below gives some standard electrode potentials that may be used in the questions that follow.

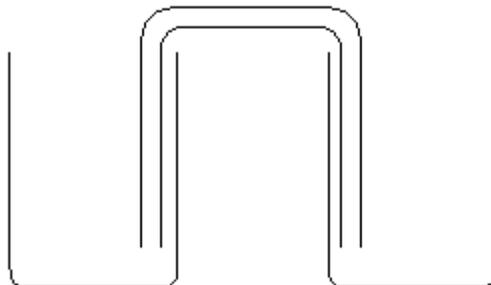
<b>Reduction half equation</b>	<b><math>E^\circ/V</math></b>
$\text{Cu}^{2+}(\text{aq}) + 2\text{e}^- \rightarrow \text{Cu}(\text{s})$	+0.34
$2\text{H}^+(\text{aq}) + 2\text{e}^- \rightarrow \text{H}_2(\text{g})$	0.00
$\text{Mg}^{2+}(\text{aq}) + 2\text{e}^- \rightarrow \text{Mg}(\text{s})$	-2.37
$\text{NO}_3^-(\text{aq}) + 4\text{H}^+(\text{aq}) + 3\text{e}^- \rightarrow \text{NO}(\text{g}) + 2\text{H}_2\text{O}$	+0.96

- a** Identify the species from the table that is the strongest reductant. **A**

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The following electrochemical cell is required:  $\text{Mg}(\text{s})/\text{Mg}^{2+}(\text{aq}) // \text{Cu}^{2+}(\text{aq})/\text{Cu}(\text{s})$

- b** Complete and label the diagram below to show how such a cell could be constructed. **A M**



The salt bridge could be made using a piece of paper soaked in a solution of potassium chloride.

- c** Explain the role of the salt bridge in the cell. **A M**

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**d** Write the net ionic equation for the overall cell reaction. **A**

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**e** Calculate the voltage of the cell in standard conditions. **A M**

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Copper metal reacts with dilute nitric acid but, unlike magnesium, has no reaction with dilute sulfuric acid.

**f** Account for the above observations using the standard reduction potentials in the table. **A M E**

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**g** Write a balanced equation for the reaction of copper metal with dilute nitric acid. **A M**

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**h** What observations could be made when copper reacts with dilute nitric acid? **A M E**

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### Question Two (Bursary 2003 Question 4: modified)

Bromine can be made from a concentrated solution of bromide ions derived from sea water. Chlorine gas is injected into acidified sea water and the bromine is produced by an oxidation-reduction reaction.

- a** Describe what would be observed as the reaction proceeds. **A**

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- b** Explain why this is an oxidation-reduction reaction. **A M**

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Acidification of the sea water is necessary because, at higher pH values, the chlorine would react with water.

- c** Write an equation for the reaction of chlorine gas with water. **A**

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- d** This reaction of chlorine with water is sometimes referred to as an **auto oxidation-reduction** reaction. Discuss this statement with regard to the reaction in **c**. **A M E**

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### Question Three (Bursary 2000 Question 8: modified)

- 1** Oxalic acid is a dicarboxylic acid ( $\text{H}_2\text{C}_2\text{O}_4$ ). On reaction with the  $\text{MnO}_4^-$  ion under acidic conditions, oxalic acid is converted to  $\text{CO}_2$ .

- a** Write the balanced half equation for the conversion of oxalic acid to  $\text{CO}_2$ . **A**

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- b** Circle the words that correctly complete the sentence.

In its reaction with  $\text{MnO}_4^-$ , oxalic acid is acting as

an oxidant.

a reductant.

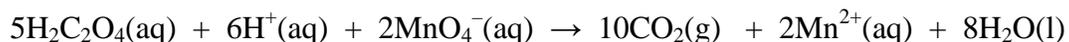
*Justify your answer in terms of oxidation numbers.* **A M**

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- 2** Solutions of potassium permanganate can be standardised by reaction with oxalic acid. The reaction occurring during titration is given below.



In the reaction for the equation given above, how many moles of electrons are transferred?

Justify your answer. **A M**

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- 3** The titration in part **2** above is generally carried out by placing the potassium permanganate in a burette and the oxalic acid in a flask.

**a** What colour is the solution in the flask at the point when exactly enough  $\text{MnO}_4^-$  has been added to react with all of the  $\text{H}_2\text{C}_2\text{O}_4$ ? **A**

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**b** What is the colour of the solution in the flask after the first drop of excess  $\text{MnO}_4^-$  has been added? **A**

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The reaction of  $\text{MnO}_4^-$  with  $\text{H}_2\text{C}_2\text{O}_4$  is catalysed by  $\text{Mn}^{2+}$ . The usual procedure is to heat the oxalic acid solution in a titration flask above  $60\text{ }^\circ\text{C}$  before adding the first portions of permanganate.

**c** Why is heating not required during the remainder of the titration? **A M**

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