WJEC Chemistry A-level

3.2: Redox Reactions

Practice Questions

Wales Specification

 Potassium peroxodisulfate(VI) (persulfate) is a white crystalline compound of formula K₂S₂O₈. It is a powerful oxidising agent and has uses as a food additive, in hair dyes and as a nappy steriliser.

| Temperature / °C | Solubility / g per 100 g H ₂ O | |
|------------------|---|--|
| 0 | 1.75 | |
| 20 | 5.29 | |

(a) Unusually for potassium compounds, it is not very soluble in water.

1 dm³ of a saturated solution of potassium persulfate at 20 °C was cooled to 0 °C. Calculate the mass of solid potassium persulfate that crystallised from the solution. [2]

 (b) (i) A hot solution of potassium persulfate slowly decomposes, giving oxygen as one of the products.

 $2K_2S_2O_8(aq) + 2H_2O(l) \longrightarrow 4KHSO_4(aq) + O_2(g)$

Calculate the maximum volume of oxygen gas that can be produced at 80 °C when a solution containing 0.100 mol of potassium persulfate decomposes as shown above. [2]

[At 80 °C 1 mol of oxygen has a volume of 29.0 dm³]

 Suggest a way that the rate of decomposition of the potassium persulfate solution described in (i) could be measured.

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(c) The diagram below shows a cell that uses persulfate ions in aqueous solution.



(d) The reaction between persulfate ions and iodide ions in aqueous solution is

$$S_2O_8^{2-}$$
 + $2I^- \longrightarrow 2SO_4^{2-}$ + I_2

In an experiment to follow the rate of this reaction, the values below were obtained.

| Experiment | Initial rate / mol dm ⁻³ s ⁻¹ | Initial concentration of $S_2O_8^{2-}$ / mol dm ⁻³ | Initial concentration of I ⁻ / mol dm ⁻³ | |
|------------|--|---|---|--|
| 1 | 8.64×10^{-6} | 0.0400 | 0.0100 | |
| 2 | 3.46×10^{-5} | 0.0800 | 0.0200 | |

(i) The reaction is first order with respect to iodide ions. Use both the initial rate values and the concentrations to show that the order with respect to persulfate ions is also first order.
 [2]



(iii) It is suggested that this reaction occurs in two steps.

| Step 1 | $S_2O_8^{2-}$ + I ⁻ + H ₂ O \longrightarrow $2SO_4^{2-}$ + HOI + H ⁺ | |
|-------------|---|--|
| Step 2 | HOI + H^+ + $I^- \longrightarrow H_2O$ + I_2 | |
| State, usir | ng your answer to (ii), why Step 1 is the rate-determining step. | |
| | | |

Total [14]

[1]

2. The leaves of the rhubarb plant are rich in ethanedioic acid (oxalic acid) which is a poisonous compound. A solution containing ethanedioate ions can be formed by boiling rhubarb leaves with water. It can be separated and samples titrated against acidified potassium manganate(VII) to find the concentration of the ethanedioate solution.

(a) Suggest how the ethanedioate solution could be separated from the rhubarb leaves.

- (b) Write an ion-electron half-equation for the reduction of acidified manganate(VII) ions, MnO₄⁻.
- (c) The ion-electron half-equation for the oxidation of ethanedioate ions is given below.

C₂O₄²⁻(aq) - 2CO₂(g) + 2e⁻

- (i) Give the oxidation states for carbon at the start and end of this reaction. [1]
- Write an equation for the reaction of acidified manganate(VII) ions with ethanedioate ions.

(d) Give a reason why an indicator is not needed in this titration.

(e) Four samples of 25.00 cm³ of the ethanedioate solution were titrated against acidified potassium manganate(VII) solution of concentration 0.0200 moldm⁻³. The volumes of potassium manganate(VII) solution required for complete reaction are listed below.

| | 1 | 2 | 3 | 4 |
|--|-------|-------|-------|-------|
| Volume of KMnO ₄ (aq)/cm ³ | 28.80 | 27.95 | 28.00 | 27.80 |

Use the information given to calculate the concentration of the ethanedioate solution. [4]

(f) Heating ethanedioic acid in glycerol produces methanoic acid, HCOOH.

- (i) Write the expression for the acid dissociation constant, K_a , for methanoic acid. [1]
- (ii) The value of K_a for methanoic acid is 1.8×10^{-4} mol dm⁻³. Calculate the pH of a solution of methanoic acid of concentration 0.2 mol dm⁻³. [3]

(iii) A mixture of methanoic acid and sodium methanoate can be used as a buffer solution. State what is meant by a *buffer solution* and explain how a mixture of methanoic acid and sodium methanoate acts as a buffer.

[3] QWC [1]

(g) Acidified potassium dichromate, K₂Cr₂O₇, is also an oxidising agent.

(i) Give the colour change that occurs when acidified potassium dichromate acts as an oxidising agent

[1]

(ii) When sodium hydroxide is added to a solution of potassium dichromate, a colour change occurs without a redox reaction occurring. Give the formula of the new chromium-containing ion and the colour of the solution formed.

(Total 20)

3. (a) Electrochemical cells are used as power sources in many everyday applications. To decide what to use in a cell, it is necessary to know the standard electrode potential for electrodes. This is measured using a standard hydrogen electrode as a reference standard.

Draw a labelled diagram of the apparatus you would use to measure the standard electrode potential of an Fe³⁺/Fe²⁺ electrode. [5]

- (b) Vanadium is a transition metal that can form compounds with a variety of oxidation states. Zinc however forms compounds with an oxidation state of +2 only.
 - Why can transition elements form compounds with a variety of oxidation states?
 [1]
 - (ii) Give the electronic structure of Zn. [1]
 - (iii) State why zinc forms Zn²⁺. [1]

You will need the standard electrode potentials in the table below to answer part (c).

| Oxidation state of vanadium at start of reaction | nadium at start Reaction | |
|--|--|-------|
| +5 | $\mathrm{VO}_3^-(\mathrm{aq}) + 4\mathrm{H^+}(\mathrm{aq}) + \mathrm{e} \rightleftharpoons \mathrm{VO}^{2+}(\mathrm{aq}) + 2\mathrm{H_2O}(\mathrm{I})$ | +1.00 |
| +4 | $VO^{2+}(aq) + 2H^{+}(aq) + e \rightleftharpoons V^{3+}(aq) + H_2O(l)$ | +0.34 |
| +3 | $V^{3+}(aq) + e \rightleftharpoons V^{2+}(aq)$ | -0.26 |
| +2 | $V^{2+}(aq) + 2e \rightleftharpoons V(s)$ | -1.13 |
| | $Zn^{2+}(aq) + 2e \rightleftharpoons Zn(s)$ | -0.76 |
| F | $Cu^{2+}(aq) + 2e \rightleftharpoons Cu(s)$ | +0.34 |

- (c) Vanadium(V)(aq), as VO₃⁻, is yellow and can be reduced by zinc and aqueous acid producing a series of coloured solutions until the reduction stops with the formation of a violet solution. The reducing agent involves the Zn²⁺(aq)/Zn(s) equilibrium.
 - State the identity of the violet vanadium-containing solution produced in this reduction. Use standard electrode potentials to explain your answer. [3]
 - (ii) What is the standard potential of a cell formed from a standard Zn²⁺(aq)/Zn(s) electrode and a standard Cu²⁺(aq)/Cu(s) electrode? [1]
 - (iii) Write the equilibrium equation for the change occurring at the zinc electrode showing the direction in which the reaction proceeds. [1]
 - Use Le Chatelier's principle to predict the effect on the electrode potential of the zinc electrode of increasing the concentration of Zn²⁺(aq) in the electrode. Explain your answer.
- (d) Halogens can also form compounds with a variety of oxidation states. Some of these including compounds of iodate(V), IO₃⁻, behave as oxidising agents.

A student was investigating the reaction that occurs when iodate(V) oxidises iodide ions to produce iodine. Two possible equations were suggested.

| $IO_3^- + 6H^+ + 5\Gamma$ | $3I_2 + 3H_2O$ | equation 1 |
|-----------------------------|-----------------------------|------------|
| $IO_3^- + 4H^+ + 4\Gamma^-$ | $IO^{-} + 2H_2O + 2I_2$ | equation 2 |

He prepared a solution of potassium iodate(V) by dissolving 0.978 g of KIO₃ in 250 cm³ of solution. He pipetted 25.0 cm³ of this solution into a conical flask, added excess potassium iodide and titrated the iodine produced with 0.100 mol dm⁻³ sodium thiosulfate solution, Na₂S₂O₃. A volume of 27.40 cm³ of this solution was needed to react with the iodate(V).

The equation for the reaction of thiosulfate with iodine is shown below.

$$2S_2O_3^{2-} + I_2 \longrightarrow S_4O_6^{2-} + 2I^{-}$$

- Calculate the number of moles of thiosulfate used to react with the iodine.
- (ii) Deduce the number of moles of iodine present in the 25.0 cm³ sample.
 [1]
- Calculate the number of moles of KIO₃ present in 250 cm³ of the original solution and hence the number of moles present in 25.0 cm³.
- (iv) Use your results from (ii) and (iii) to deduce which of equation 1 and equation 2 suggested above, correctly shows what happens when iodate(V) ions oxidise iodide ions. Show, by calculation, how you came to this conclusion. [2]

Total [20]

Hydrogen Peroxide

If a non-scientist knows only one chemical formula it is most likely to be H₂O for water but how much do you know about another hydrogen oxide, hydrogen peroxide? A molecule of hydrogen peroxide has the molecular formula H₂O₂.

Most chemistry students first meet hydrogen peroxide as a colourless solution that is used to 5 prepare oxygen. Bottles of hydrogen peroxide from a pharmacist are often labelled '20 volume'. This means that one volume of solution decomposes to give 20 volumes of oxygen gas. The equation for the decomposition is:

$$2H_2O_2(aq) \longrightarrow 2H_2O(I) + O_2(g)$$

 $1 dm^3 \qquad 20 dm^3$

This reaction is very slow at room temperature. However the addition of a suitable catalyst 10 increases the rate of decomposition phenomenally. Manganese(IV) oxide, potatoes and blood are all effective. Potatoes and blood both contain the enzyme catalase and one catalase molecule decomposes 50 000 molecules of H₂O₂ per second!

Is hydrogen peroxide an oxidising agent or a reducing agent?

Both in the laboratory and at home hydrogen peroxide is most commonly used as an oxidising agent (so the hydrogen peroxide itself is reduced). The half-equation is:

Reduction $H_2O_2 + 2H^+ + 2e^- \longrightarrow 2H_2O_2$

Since some colouring matter is bleached by oxidation and the product of hydrogen peroxide's reduction is water, it is used as a safe bleaching agent particularly in hair treatment. A peroxide blonde is someone with almost white hair, usually as a result of treatment with hydrogen peroxide.

20 However, if hydrogen peroxide reacts with a more powerful oxidising agent such as potassium manganate(VII), the hydrogen peroxide will act as a reducing agent and will itself be oxidised. The half-equation is:

Oxidation $H_2O_2 \longrightarrow 2H^+ + O_2 + 2e^-$

Therefore hydrogen peroxide can act as both oxidising agent and reducing agent. In fact, it can 25 react with itself so that alternate molecules are oxidised and reduced. The overall equation is obtained by adding the half-equations for the reduction and oxidation, giving

 $2H_2O_2(aq) \longrightarrow 2H_2O(I) + O_2(g)$

which is the standard decomposition equation!

- End of passage -

(a) Using outer electrons only, draw a dot and cross diagram to show the bonding in a hydrogen peroxide molecule(*line 3*).

[1]

(b) Use the equation for the decomposition of hydrogen peroxide *(line 8)* to calculate the concentration, in mol dm^{-3} , of aqueous hydrogen peroxide solution in a bottle of '20 volume hydrogen peroxide' at 25 °C.

[1 mol of oxygen occupies 24 dm³ at 25 °C]

Concentration = mol dm-³

(c) Manganese(IV) oxide *(line 10)* and potassium manganate(VII) *(lines 20-21)* are typical transition metal compounds.

(i) Give two reasons why transition metal compounds can act as catalysts.

[2]

(ii) Explain why transition metal complex ions appear coloured.

[4] QWC [1]

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(d) In an acidic solution, hydrogen peroxide is oxidised to oxygen by potassium manganate (VII) *(lines 20-23).*

(i) Write the half-equation for the reduction of MnO_4^- to Mn^{2+} ions in acidic solution.

(ii) Use your answer to (i) and the half-equation given in *line 23* to deduce the overall equation for this reaction.

[2]

[1]

(iii) 20.0 cm³ of an acidified solution of hydrogen peroxide required 14.80 cm³ of a 0.020 mol dm⁻³ solution of potassium manganate(VII) for complete reaction. Calculate the concentration, in mol dm⁻³, of the hydrogen peroxide solution.

[3]

Concentration = mol dm⁻³

| | | | [2] |
|-----|------|---|------------------------------|
| | | | (Total 18) |
| 6. | Haer | natite is an ore of iron that contains Fe_2O_3 . Iron is extracted from this ore in a | bla <mark>st</mark> furnace. |
| | (a) | Balance the equation for the extraction of iron from Fe_2O_3 . | [1] |
| | | | |
| | (b) | Use oxidation states to show that the reaction in (a) is a redox reaction. | [2] |
| | | | |
| | | | |
| | | | |
| | (C) | A different oxide of iron is iron(II) oxide, FeO. The ions in this compo arrangement similar to that of sodium chloride. | und adopt an |
| | | (i) Give the crystal co-ordination numbers for the ions in FeO. | [1] |
| | | | |
| | | (ii) Draw the arrangement of oxide ions around each iron ion. | [1] |
| (d) | Iro | n can be extracted from FeO according to the equation below. | |
| | | FeO + CO Fe + CO ₂ | |
| | Ca | Iculate the mass of iron that could be extracted from 20.0 kg of iron(II) oxide | e, FeO. <mark>[</mark> 3] |

(e) Explain, using oxidation states, why the decomposition of hydrogen peroxide (line 27) can be

classified as a redox reaction.

(e) Carbon monoxide contains two covalent bonds and one co-ordinate bond. Explain what is meant by the terms *covalent bond* and *co-ordinate bond*, indicating the difference between them.

[2]

(f) Iron is a typical metal. Describe the bonding present in iron. Explain how it can conduct electricity and why it has a high melting temperature.

[4] QWC [1]

(Total 15)