

Boost your performance and confidence with these topic-based exam questions

Practice questions created by actual examiners and assessment experts

Detailed mark scheme

Suitable for all boards

Designed to test your ability and thoroughly prepare you



Time allowed 73 Minutes

Score

/61

%

CHEMISTRY

OCR AS & A LEVEL

Topic Questions

Module 5: Physical chemistry and transiton elements

www.exampaperspractice.co.uk



		s a transition element. Solid compound ions are formed.	s of cobalt are often complexes and in solution,		
(a)	In its complexes, the common oxidation numbers of cobalt are +2 and +3.				
	Cor	Complete the electron configurations of cobalt as the element and in the +3 oxidation state:			
	cob	alt as the element:	1s ² 2s ² 2p ⁶		
	cob	ealt in the +3 oxidation state:	1s ² 2s ² 2p ⁶ [2]		
(b)		te one property of cobalt(II) and cobalt(ch is typical of ions of a transition eleme	III), other than their ability to form complex ions, nt.		
			[1]		
(c)	Cor	mplex ions contain ligands.			
	Sta	te the meaning of the term <i>ligand</i> .			
			[1]		
(d)	Αqι	ueous cobalt(II) sulfate, CoSO ₄ (aq), take	s part in the following reactions.		
	For each reaction, state the formula of the transition element species formed and the type or reaction taking place.				
	(i)	Aqueous cobalt(II) sulfate, CoSO ₄ (aq),	reacts with aqueous sodium hydroxide.		
		transition element species formed:			
		type of reaction:	[2]		
	(ii)	Aqueous cobalt(II) sulfate, CoSO ₄ (aq),	reacts with concentrated hydrochloric acid.		
		transition element species formed:			
		type of reaction:	[2]		

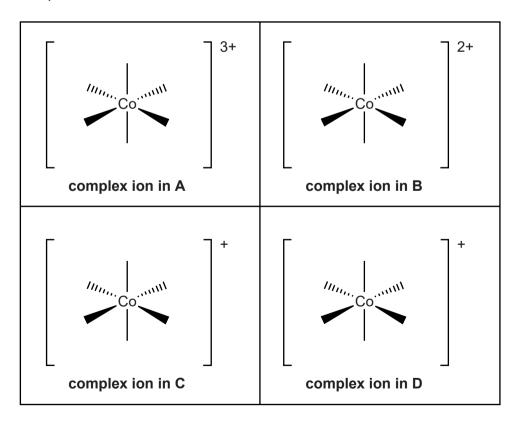


(e) Cobalt(III) chloride, $CoCl_3$, reacts with ammonia to form a range of complexes. These complexes contain different amounts of ammonia. Information about these complexes is summarised below.

The complex ions **C** and **D** are stereoisomers.

complex	formula	formula of complex
Α	$CoCl_3(NH_3)_6$	[Co(NH ₃) ₆] ³⁺ 3C <i>l</i> ⁻
В	$CoCl_3(NH_3)_5$	$[Co(NH_3)_5Cl]^{2+}2Cl^{-}$
С	CoCl ₃ (NH ₃) ₄	[Co(NH ₃) ₄ C <i>l</i> ₂] ⁺ C <i>l</i> ⁻
D	CoCl ₃ (NH ₃) ₄	[Co(NH ₃) ₄ C <i>l</i> ₂] ⁺ C <i>l</i> ⁻

(i) Complete the diagrams below to suggest possible structures for the complex ion in complexes A to D.





(ii) Chemists provided evidence for the formulae of these complexes from their reactions with aqueous silver nitrate. Aqueous silver nitrate reacts with aqueous halide ions in a precipitation reaction.

An excess of silver nitrate solution was reacted with 0.0100 mol of one of the complexes **A** to **D**. 2.868 g of a precipitate was formed.

Determine which complex was reacted.

In your answer you should explain how the result of the experiment would allow the formula of the complex to be identified.
[3]

[Total: 15]



2	2 Iron and platinum are transition elements. They both form ions that combine with ligands complex ions. Some of these complexes are important in biological systems.				orm		
	(a)	Complete the electr	ron structures of:				
		an atom of Fe:	1s ² 2s ² 2p ⁶				
		an ion of Fe ²⁺ :	1s ² 2s ² 2p ⁶				. [2]
	(b)	State one property ion of a transition e		n the ability to fo	rm complex ions,	which is typical o	f an
							. [1]
	(c)	Aqueous iron(II) su	lfate takes part in	redox reactions.			
		Using oxidation nu redox reaction of ac				ve taken place in	the
	6	FeSO ₄ + 7H ₂ SO ₄	+ $Na_2Cr_2O_7 \rightarrow$	3Fe ₂ (SO ₄) ₃ +	Cr ₂ (SO ₄) ₃ + Na ₂	₂ SO ₄ + 7H ₂ O	
							[2]



(d)	Hexaaquairon(II) ions, $[Fe(H_2O)_6]^{2+}$, take part in a ligand substitution reaction with ammonia
	$[Fe(H_2O)_6]^{2+}(aq) + 6NH_3(aq) \rightleftharpoons [Fe(NH_3)_6]^{2+}(aq) + 6H_2O(I)$
	Write an expression for the stability constant, K_{stab} , for this equilibrium.

[2]

.....[2]



(f)	Pla ster	tin, $Pt(NH_3)_2Cl_2$, is a complex of platinum(II) that has two stereoisomers. On reoisomers is used in medicine.	e of these
	(i)	Platin is a neutral complex.	
		Explain why platin is neutral.	
			[1]
	(ii)	Draw diagrams of the two stereoisomers of platin and describe its bonding.	
			[3]
	(iii)	Describe the action of platin in the treatment of cancer patients.	



(g) The use of platin in medicine can cause unpleasant side effects for patients.

In the search for alternatives, chemists often start with the current drug and modify its properties by chemically changing some of the groups.

A recent discovery is a drug called carboplatin. The structure of carboplatin is similar to platin except that a single 1,1-cyclobutanedicarboxylate ion replaces the two chloride ligands in the structure of platin.

Draw the structures of,

- the 1,1-cyclobutanedicarboxylate ion
- carboplatin

1,1-cyclobutanedicarboxylate ion

carboplatin

[2]

[Total: 18]



3 The Dissolved Oxygen Concentration (DOC) in rivers and lakes is important for aquatic life. If the DOC falls below 5 mg dm⁻³, most species of fish cannot survive.

Environmental chemists can determine the DOC in water using the procedure below.

A sample of river water is shaken with aqueous Mn²⁺ and aqueous alkali.
 The dissolved oxygen oxidises the Mn²⁺ to Mn³⁺, forming a pale brown precipitate of Mn(OH)₃.

$$O_2(aq) + 4Mn^{2+}(aq) + 8OH^{-}(aq) + 2H_2O(I) \rightarrow 4Mn(OH)_3(s)$$

 The Mn(OH)₃ precipitate is then reacted with an excess of aqueous potassium iodide, which is oxidised to iodine, I₂.

$$2\mathsf{Mn}(\mathsf{OH})_3(\mathsf{s}) \ + \ 2\mathsf{I}^-(\mathsf{aq}) \ \longrightarrow \ \mathsf{I}_2(\mathsf{aq}) \ + \ 2\mathsf{Mn}(\mathsf{OH})_2(\mathsf{s}) \ + \ 2\mathsf{OH}^-(\mathsf{aq})$$

 The iodine formed is then determined by titration with aqueous sodium thiosulfate, Na₂S₂O₃(aq).

$$2S_2O_3^{2-}(aq) + I_2(aq) \rightarrow S_4O_6^{2-}(aq) + 2I^{-}(aq)$$

A 25.0 cm³ sample of river water was analysed using the procedure above.

The titration required 24.6 $\rm cm^3$ of 0.00100 mol $\rm dm^{-3}~Na_2S_2O_3(aq)$.

(a) (i) Calculate the DOC of the sample of river water, in mg dm⁻³.

$$DOC =mg dm^{-3}$$
 [4]



ter for fish to
[1]
NO₂⁻ ions can
−1 .
[1]
trate(III) ions.
[2]
[Total: 8]



Potassium manganate(VII) can be prepared in the laboratory by a two-step synthesis starting from manganese(IV) oxide.

Step 1

In this step, manganese(IV) oxide is heated strongly with potassium hydroxide and potassium chlorate(V), a powerful oxidising agent.

Manganese(IV) oxide, MnO₂, is oxidised to manganate(VI) ions.

Step 2

Potassium manganate(VI) is separated from the alkaline mixture from step 1 as a green solid.

In this step, potassium manganate(VI) is heated in water. Manganate(VI) ions disproportionate forming manganate(VII) ions and a precipitate of manganese(IV) oxide.

(a) In step 1, a redox reaction takes place.

Add the correct number of electrons to the correct sides of the incomplete oxidation and reduction half-equations shown below.

$$MnO_2 + 4OH^- \rightarrow MnO_4^{2-} + 2H_2O$$

 $3H_2O + ClO_3^- \rightarrow 6OH^- + Cl^-$

$$3H_2O + ClO_3^- \rightarrow 6OH^- + Cl^-$$

[2]

(b) In step 2, an equilibrium is set up.

$$3MnO_4^{2-}(aq) + 2H_2O(I) \rightleftharpoons 2MnO_4^{-}(aq) + MnO_2(s) + 4OH^{-}(aq)$$

Suggest, with the aid of an equation, how the equilibrium position shifts.

The equilibrium position can be shifted by bubbling carbon dioxide gas through the mixture.



(c) Aqueous potassium manganate(VII), KMnO₄, in acidic conditions can be used in analysis.

A student analyses a sample of sodium sulfite, Na₂SO₃, using the following method.

- The student dissolves 0.720 g of impure sodium sulfite in water.
- The solution is made up to 100.0 cm³.
- The student titrates 25.0 cm³ of this solution with 0.0200 mol dm⁻³ KMnO₄ under acidic conditions. The volume of KMnO₄(aq) required to reach the end-point is 26.2 cm³.

The equation for the reaction is shown below.

$$2 \text{MnO}_4^{\;-} \; + \; 6 \text{H}^+ \; + \; 5 \text{SO}_3^{\;2-} \; \longrightarrow \; 2 \text{Mn}^{2+} \; + \; 5 \text{SO}_4^{\;2-} \; + \; 3 \text{H}_2 \text{O}$$

Determine the percentage purity of the sample of sodium sulfite.

[Total: 10]



5 Haematite is the main ore of iron. The percentage of iron in a sample of haematite can be determined using the method below.

Method

- **Stage 1.** An excess of concentrated hydrochloric acid is added to a 3.25 g sample of haematite. The iron(III) oxide in the haematite reacts to form a solution containing Fe³⁺ ions.
- **Stage 2.** An excess of aqueous tin(II) chloride is added. Sn²⁺ reduces the Fe³⁺ present to Fe²⁺. Excess Sn²⁺ is removed.
- **Stage 3.** The solution is diluted and made up to 250.0 cm³ in a volumetric flask.
- **Stage 4.** A 25.0 cm³ sample of this solution is pipetted into a conical flask.
- **Stage 5.** The solution in the conical flask is titrated with $0.0200\,\mathrm{mol\,dm^{-3}}$ aqueous potassium dichromate(VI), $\mathrm{K_2Cr_2O_7}$. The Fe²⁺ ions are oxidised to Fe³⁺ ions.
- Stage 6. Stages 4 and 5 are repeated to obtain an average titre of 26.5 cm³.

You are provided with the following electrode potentials.

You may need to use this information throughout this question.

(a)	Write an equation for the reaction between iron(III) oxide and concentrated hydrochloric acid
	occurring in Stage 1.

_____[1]

(b) Write equations for the reactions involving iron ions in Stages 2 and 5.

Stage 2



	percentage iron = % [5]
(d)	Aqueous potassium manganate(VII), $KMnO_4(aq)$, is not suitable for titrating the solution in this method. Aqueous potassium dichromate(VI), $K_2Cr_2O_7(aq)$, is used instead.
	Suggest and explain why potassium dichromate(VI), $K_2Cr_2O_7$, is suitable for this titration whereas potassium manganate(VII), $KMnO_4$, is not suitable.
	·
	[2]
	[Total: 10]

(c) Calculate the percentage by mass of iron in the haematite ore.