# **Transition Elements**

1. \* B and C are compounds of two different transition elements.

A student carries out test tube reactions on aqueous solutions of  ${\bf B}$  and  ${\bf C}$ . The observations of the student's tests are shown below.

	Test	B(aq)	C(aq)
	NH <sub>3</sub> (aq) added dropwise	green precipitate <b>D</b>	grey-green precipitate <b>E</b>
1	excess NH₃(aq) added	no further change	purple solution <b>F</b>
	HNO₃(aq)	no change	no change
2	followed by Ba(NO <sub>3</sub> ) <sub>2</sub> (aq)	white precipitate <b>G</b>	no change
	HNO₃(aq)	no change	no change
3	followed by AgNO₃(aq)	no change	white precipitate <b>H</b>

Analyse the results to identify ${\bf B}$ to ${\bf H}$ , and construct ionic equations for the formation of products ${\bf D}$ to ${\bf H}$ .				

Transition metal ions can bond to ligands to form complex ions with different shapes.  Explain what is meant by the terms ligand, coordination number and ligand substitution, using suitable examples of complex ions with different shapes, limited to monodentate ligands. Your answer should include diagrams and equations where appropriate.	
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3.

	<u>[6]</u>
Iron(II) g	lluconate, C <sub>12</sub> H <sub>22</sub> FeO <sub>14</sub> , is the active ingredient in some brands of iron supplements.
	nt carries out an experiment to determine the mass of iron(II) gluconate in one tablet of an
iron sup	plement, using the method below.
	The student crushes two tablets and dissolves the powdered tablets in dilute
Stage 1	sulfuric acid
Stage 2	The student makes up the solution from <b>Stage 1</b> to 250.0 cm <sup>3</sup> in a volumetric flask.
_	The student then titrates 25.0 cm <sup>3</sup> portions of the solution obtained in <b>Stage 2</b> with
	0.00200 mol dm <sup>-3</sup> potassium manganate(VII).
Stage 3	TI 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1
	The student obtains a mean titre of 13.50 cm <sup>3</sup> .
	In this titration, 1 mol of manganate(VII) ions reacts with 5 mol of iron(II) ions.
i.	Explain why the student used 0.00200 mol dm <sup>-3</sup> potassium manganate(VII) solution for
••	this titration, rather than the more usual concentration of 0.0200 mol dm <sup>-3</sup> used in
	manganate(VII) titrations.
	[1]

ii. Use the student's results to determine the mass, in mg, of iron(II) gluconate in <b>one</b> tablet.					
	Give your answer to 3 s	ignificant figures.			
	mass of iron(II) glu	uconate in one tablet =	mg <b>[5</b> ]		
iii.	Some iron supplements	contain iron(II) sulfate or iron(II	) fumarate.		
		, ,	of two iron supplements, <b>A</b> and <b>B</b> .		
г					
	Iron supplement	Iron compound	Mass of iron compound in one tablet / mg		
	Iron supplement	Iron compound iron(II) sulfate, FeSO <sub>4</sub>	Mass of iron compound in one tablet / mg		
		·	in one tablet / mg		
	A	iron(II) sulfate, FeSO <sub>4</sub>	in one tablet / mg		
	A B	iron(II) sulfate, FeSO <sub>4</sub> iron(II) fumarate, C <sub>4</sub> H <sub>2</sub> FeO <sub>4</sub>	180 210		
	A B	iron(II) sulfate, FeSO <sub>4</sub> iron(II) fumarate, C <sub>4</sub> H <sub>2</sub> FeO <sub>4</sub> <b>Table 18.3</b>	180 210		
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4.	Red blo	od cells contain haemoglobin.
	Explain	using ligand substitutions:
		ow haemoglobin transports oxygen around the body hy carbon monoxide is toxic.
5(a).	A stude  Experir The stu	estion is about reactions of ions and compounds of transition elements. In transition elements on a solution containing $[Cr(H_2O)_6]^{3+}(aq)$ . The solution are solution containing $[Cr(H_2O)_6]^{3+}(aq)$ until a solution is formed.
	<b>Experir</b> The stu	ment 2 dent carries out the following reaction sequence.
	Step 1	NaOH(aq) is added slowly to a solution containing $[Cr(H_2O)_6]^{3+}(aq)$ in a boiling tube. A grey–green precipitate forms. An excess of NaOH(aq) is added to the boiling tube.
	Step 2 Step 3 Step 4	The precipitate dissolves and a green solution forms containing a 6 coordinate complex ion. $H_2O_2$ is added to the mixture and the boiling tube is heated. A yellow solution forms. The solution in the boiling tube is acidified. The solution now contains $Cr_2O_7^{2-}$ (aq).
	i.	What is the formula of the complex ion in the purple solution that forms <b>in Experiment</b> 1?
	ii.	Suggest an equation for the reaction in <b>Experiment 2, Step 1</b> . Include state symbols.
		<u></u>

(b).

ii. Draw a 3-D diagram for the <b>2</b> .		• ,	, i.e.
Include the charge of the i	on.		
v. What is the formula of the	ion that causes the yellow col	our in Experiment 2, Step	<b>3</b> ?
. State the colour of the colo			
v. State the colour of the solu	ution that forms in <b>Experiment</b>	. 2, Step 4.	
	oxidation states. <b>Table 18.1</b> s		
ueous solution.			
ueous solution.	oxidation states. <b>Table 18.1</b> s	hows the colours of the ions	
Decide the second secon	oxidation states. <b>Table 18.1</b> s <b>Vanadium ion</b>	hows the colours of the ions	
Dxidation state of vanadium +5	oxidation states. <b>Table 18.1</b> s <b>Vanadium ion</b> VO <sub>2</sub> *(aq)	hows the colours of the ions  Colour  yellow	
+4	Vanadium ion  VO2 <sup>+</sup> (aq)  VO3 <sup>+</sup> (aq)	Colour yellow blue green	- -
Dxidation state of vanadium +5 +4	oxidation states. <b>Table 18.1</b> since $Vanadium\ ion$ $VO_2^{+}(aq)$ $VO^{2+}(aq)$ $V^{3+}(aq)$ $V^{2+}(aq)$	Colour yellow blue	
Dxidation state of vanadium +5 +4 +3	Vanadium ion  VO2 <sup>+</sup> (aq)  VO3 <sup>+</sup> (aq)	Colour yellow blue green	
Dxidation state of vanadium +5 +4 +3	Vanadium ion  VO2 <sup>+</sup> (aq)  V3 <sup>+</sup> (aq)  V <sup>3+</sup> (aq)  V <sup>2+</sup> (aq)	Colour yellow blue green	
Dxidation state of vanadium  +5  +4  +3  +2	Vanadium ion  VO2 <sup>+</sup> (aq)  V3 <sup>+</sup> (aq)  V <sup>3+</sup> (aq)  V <sup>2+</sup> (aq)	Colour yellow blue green	
Dxidation state of vanadium  +5  +4  +3  +2	Vanadium ion  VO2 <sup>+</sup> (aq)  V3 <sup>+</sup> (aq)  V <sup>3+</sup> (aq)  V <sup>2+</sup> (aq)	Colour yellow blue green	s in

ii. The student adds excess iron to a solution containing VO<sup>2+</sup>(aq) ions, and observes that the colour of the solution changes from blue to green and then to violet.

Use the relevant standard electrode potentials shown in **Table 18.2** to explain these observations.

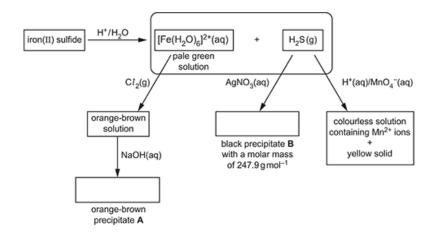
	Redox system	E°/V
1	$V^{2+}(aq) + 2e^- \rightleftharpoons V(s)$	-1.18
2	$Fe^{2+}(aq) + 2e^- \rightleftharpoons Fe(s)$	-0.44
3	$V^{3+}(aq) + e^- \rightleftharpoons V^{2+}(aq)$	-0.26
4	$VO^{2+}(aq) + 2H^+ + e^- \rightleftharpoons V^{3+}(aq) + H_2O(I)$	+0.34
5	$Fe^{3+}(aq) + e^- \rightleftharpoons Fe^{2+}(aq)$	+0.77
6	$VO_{2+}(aq) + 2H^+ + e^- \rightleftharpoons VO^{2+}(aq) + H_2O(I)$	+1.00

**Table 18.2** 

		[3]
iii.	Construct an equation for the <b>first</b> colour change from blue to green.	
		[1]

**6(a).** This question is about reactions of iron compounds.

A student carries out the reactions in the flowchart, starting with iron(II) sulfide.



i. In the boxes, write the formulae of **A** and **B**.

iv.

ii. The student thinks that the reaction of iron(II) sulfide with H<sup>+</sup> / H<sub>2</sub>O is a redox reaction.

Explain, with reasons, whether the student is correct.

[1]

iii. Write the equation for the reaction of [Fe(H<sub>2</sub>O)<sub>6</sub>]<sup>2+</sup>(aq) with C/<sub>2</sub>(g).

Construct an equation for the reaction of  $H_2S(g)$  with  $H^+(aq)/MnO_4^-(aq)$ .

[2]

(b). \*Compound C is a hydrated ionic compound with the empirical formula: FeH<sub>18</sub>N<sub>3</sub>O<sub>18</sub>.

A student investigates the thermal decomposition of compound C as outlined below.

## Stage 1

The student gently heats 0.00300 mol of compound  ${\bf C}$  to remove the water of crystallisation. 0.486 g of water is collected, leaving 0.00300 mol of the anhydrous compound  ${\bf D}$ .

#### Stage 2

The student strongly heats 0.00300 mol of compound  $\bf D$ , which decomposes to form a solid oxide E (molar mass of 159.6 g mol<sup>-1</sup>) and 270 cm<sup>3</sup> of a gas mixture, measured at RTP, containing gases  $\bf F$  and  $\bf G$ .

### Stage 3

The student cools the 270 cm<sup>3</sup> gas mixture of **F** and **G**.

- Gas **F** is a compound that condenses to form 0.414 g of a liquid.
- Gas **G** remains and has a volume of 54 cm<sup>3</sup>, measured at RTP.
  - Gas **G** is tested and it relights a glowing splint.

Determine the formulae of <b>C</b> , <b>D</b> , <b>E</b> , <b>F</b> and <b>G</b> .				
Show all your working and equations for the	e reactions.			

7.

	 6]
Cobalt(II) forms complex ions with water ligands and with chloride ligands.	
<ul> <li>With water ligands, cobalt(II) forms a pink octahedral complex ion, [Co(H<sub>2</sub>O)<sub>6</sub>]<sup>2+</sup>.</li> <li>With chloride ligands, cobalt(II) forms a blue tetrahedral complex ion.</li> <li>A student dissolves cobalt(II) sulfate in water in a boiling tube. A pink solution forms.</li> </ul>	
Experiment 1 The student places the boiling tube in a water bath at 100 °C. Concentrated hydrochloric acid is added dropwise. The colour of the solution changes from pink to blue.	
Experiment 2 The student places the boiling tube from experiment 1 in an ice/water bath at 0 °C. The colour of the solution changes from blue to pink.	
<ul> <li>Write the equilibrium equation for the reaction that takes place when the colour of the solution changes.</li> </ul>	
[1	1
ii. Explain the observations and predict whether the formation of the blue colour is exothermic or endothermic.	
[2	2]

8. This question is about ethanedioic acid, (COOH)2, and ethanedioate ions, (COO<sup>-</sup>)2.

The ethanedioate ion, shown below, can act as a bidentate ligand.



Fe<sup>3+</sup> forms a complex ion with three ethanedioate ions. The complex ion has two optical isomers.

Draw the 3D shapes of the optical isomers.

In your diagrams, show the structure of the ethanedioate ligands and any overall charge.

[3]

**9(a).** This question is about some reactions of d block elements and their ions.

**Table 21.1** shows standard electrode potentials which will be needed within this question.

Zn <sup>2+</sup> (aq) + 2e <sup>-</sup>	$\leftarrow$	Zn(s)	E <sup>⊕</sup> = −0.76 V
Cr <sup>3+</sup> (aq) + e <sup>-</sup>	$\rightleftharpoons$	Cr <sup>2+</sup> (aq)	E <sup>e</sup> = −0.42 V
Ni <sup>2+</sup> (aq) + 2e <sup>-</sup>	$\rightleftharpoons$	Ni(s)	E <sup>⊕</sup> = −0.25 V
$I_2(aq) + 2e^-$	$\rightleftharpoons$	2I <sup>-</sup> (aq)	$E^{\Theta} = +0.54 \text{V}$
Fe <sup>3+</sup> (aq) + e <sup>-</sup>	$\rightleftharpoons$	Fe <sup>2+</sup> (aq)	E <sup>⊕</sup> = +0.77 V
$Cr_2O_7^{2-}(aq) + 14H^+(aq) + 6e^-$	$\rightleftharpoons$	$2Cr^{3+}(aq) + 7H_2O(I)$	$E^{\Theta} = +1.33 \text{V}$
$H_2O_2(aq) + 2H^+(aq) + 2e^-$	$\rightleftharpoons$	2H <sub>2</sub> O(I)	E <sup>⊕</sup> = +1.78 V

**Table 21.1** 

Complete the electron configuration of

a Ni atom: 1s²		
a Ni <sup>2+</sup> ion: 1s <sup>2</sup>	2	[2]

Reaction 1:	Aqueous copper(II) sulfate reacts with excess aqueous ammonia in a ligar substitution reaction. A deep-blue solution is formed, containing an octah complex ion, <b>C</b> , which is a trans isomer.	
Reaction 2:	Copper(I) oxide reacts with hot dilute sulfuric acid in a disproportionation reaction. A blue solution, $\mathbf{D}$ , and a brown solid, $\mathbf{E}$ are formed.	
Reaction 3:	Copper(II) oxide reacts with warm dilute nitric acid in a neutralisation react to form a blue solution. Unreacted copper(II) oxide is filtered off, and the solution is left overnight in an evaporating basin.  A hydrated salt, <b>F</b> , crystallises, with the percentage composition by mass	
	Cu, 26.29%; H, 2.48%; N, 11.59%; O, 59.63%.	
Identify <b>C-F</b> by for	mulae or structures, as appropriate.	
Include equations,	any changes in oxidation number, and working.	[6]

(b). \* Three different reactions of copper compounds are described below.

10(a).

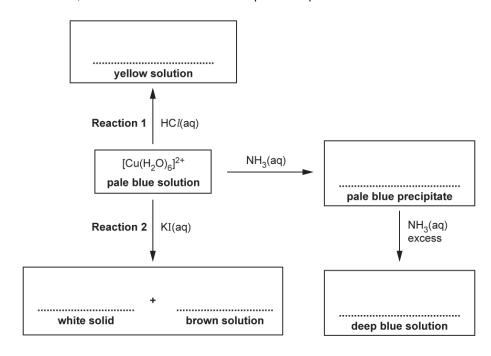
A hydrated nickel(II) complex, <b>A</b> , is heated in a	a crucible to remove the water of
crystallisation.	
The anhydrous complex <b>B</b> is formed. The results are	shown below.
Mass of crucible + hydrated complex A	- 50 551 a
	= 59.554 g
Mass of crucible + anhydrous complex <b>B</b> Mass of crucible	= 58.690 g
Mass of crucible	= 51.257 g
The anhydrous complex <b>B</b> is analysed and found to I	nave a molar mass of 309.7 g mol <sup>-1</sup> and to
contain the following percentage composition by mas	
Ni, 18.95%; C, 23.25%; N, 27.12	%; H, 7.75%; C <i>I</i> , 22.93%.
The enhydrous complex B contains a setion C comp	riging Ni. C. N. and H. anly
The anhydrous complex <b>B</b> contains a cation <b>C</b> comp	nsing Ni, C, N and H only.
Cation <b>C</b> is six-coordinate, contains three molecules	of the bidentate ligand <b>D</b> , and exists as
optical isomers.	
·	
Determine the formula of <b>A</b> , <b>B</b> , <b>C</b> and <b>D</b> and show the	e 3D structures for the optical isomers of <b>C</b> .
Show all your working	
Show <b>all</b> your working.	

 [6]

(b). This question is about reactions of ions and compounds of transition elements.

The flowchart shows reactions of the complex ion  $[Cu(H_2O)_6]^{2+}.$ 

i. In the boxes, write down the formulae of the species responsible for the observations.



[5]

ii. Name the type of reaction for **Reaction 1** and **Reaction 2**.

Reaction 1	
Reaction 2	

**11.** Chromium (III) picolinate, shown below, is a neutral complex that can be prepared from the weak acid, picolinic acid.

Chromium(III) picolinate is used in tablets as a nutritional supplement for chromium.

i. Draw the structure of the ligand in chromium(III) picolinate.

ii. A typical tablet of chromium(III) picolinate contains 200 µg of chromium.

Calculate the mass, in g, of chromium (III) picolinate in a typical tablet. 1  $\mu g$  =  $10^{-6}$  g.

Give your answer to **three** significant figures.

mass	
=	

**12.** Acid rain is caused by the reaction of acid gases with water and oxygen in the air.

Coal often contains traces of iron(II) disulfide, FeS2.

 $FeS_2$  is an ionic compound of  $Fe^{2+}$  ions and  $S_2^{2-}$  ions.

i. Write the electron configuration, in terms of sub-shells, of an Fe<sup>2+</sup> ion.

[1]

	ii.	Draw a 'dot-and-cross' diagram for FeS <sub>2</sub> .	
		Show outer electrons only.	
			[2]
13(a).	This qu	uestion is about chemicals used by gardeners.	
	A garde (NH <sub>4</sub> ) <sub>2</sub> F	en product contains hydrated ammonium iron(II) sulfate, (NH <sub>4</sub> ) <sub>2</sub> Fe(SO <sub>4</sub> ) <sub>2</sub> •xH <sub>2</sub> O. Fe(SO <sub>4</sub> ) <sub>2</sub> •xH <sub>2</sub> O contains 27.55% by mass of water of crystallisation.	
	Calcula	ate the value of x in the formula $(NH_4)_2Fe(SO_4)_2•xH_2O$ .	
	Show y	our working.	
		x =	[3]
(b).	The ga	orden product in the previous question part is a solid mixture of the following ingredier	nts:
	•	Hydrated ammonium iron(II) sulfate, $(NH_4)_2Fe(SO_4)_2 \cdot xH_2O$ , which is soluble in water Crushed limestone (calcium carbonate) Sand.	er
	i.	Suggest why crushed limestone has been included in this garden product.	
			[1]

- ii. \*Plan a procedure on a test tube scale to show that the solid mixture contains the following ions:
  - NH<sub>4</sub>+, Fe<sup>2+</sup> and SO<sub>4</sub><sup>2-</sup> present in (NH<sub>4</sub>)<sub>2</sub>Fe(SO<sub>4</sub>)<sub>2</sub>•xH<sub>2</sub>O
  - CO<sub>3</sub><sup>2-</sup> present in crushed limestone.

		Show your reasoning, including relevant equations.
(c ).	Some g	ardeners spray crops with 'Bordeaux mixture' which contains several compounds.
,-		the compounds in Bordeaux mixture is prepared by a student. The student added aqueous sodium de to aqueous copper(II) sulfate.
	i.	Write an ionic equation, including state symbols, for the reaction in this preparation.
		What would be observed?
		Equation
		Observation
	ii.	<b>[2]</b> It is believed that Cu <sup>2+</sup> ions in the Bordeaux mixture interact with protein molecules in fungi, preventing growth. Proteins are polyamides.
		Suggest the interactions which occur between Cu <sup>2+</sup> ions and protein molecules in fungi.

	[2]
<b>14</b> A stud	dent investigates reactions of cobalt ions, as outlined below.
•	A student dissolves cobalt(II) chloride in water. A pink solution forms containing the hexaaqua complex ion <b>B</b> .
•	The student adds an excess of concentrated ammonia solution to the pink solution until there is no further change.
•	A pale brown solution forms which contains the complex ion [Co(NH <sub>3</sub> ) <sub>6</sub> ] <sup>2+</sup> .
i.	Write the equation for the formation of $[Co(NH_3)_6]^{2+}$ from complex ion <b>B</b> .
	State the type of reaction.
	Equation
	Type of reaction
	ro1
ii.	<b>[2]</b> Draw a 3-D diagram of the [Co(NH₃)₅] <sup>2+</sup> ion.

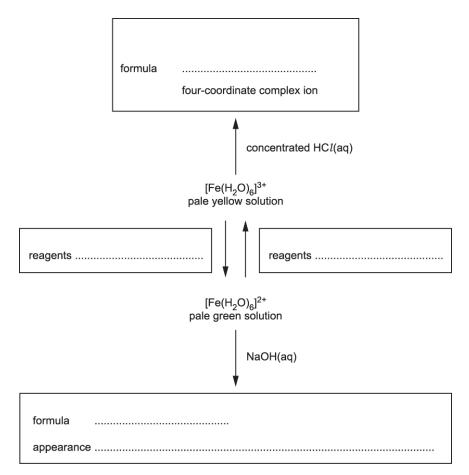
III.	o o	The solution is warmed with aqueous hydrogen peroxide, H <sub>2</sub> O <sub>2</sub> (aq).  The H <sub>2</sub> O <sub>2</sub> oxidises cobalt(II) to cobalt(III), to form a red-brown solution containing coordinate complex ion <b>C</b> .  Concentrated hydrochloric acid is added to the red-brown solution.  Yellow crystals of a complex <b>D</b> are formed.	j a six-
	Complex	x <b>D</b> has the percentage composition by mass:	
		Co, 22.03%; N, 31.41%, H, 6.73%; CI, 39.83%.	
	Determi	ne the formulae of <b>C</b> and <b>D</b> , showing clearly the ligands and any charges.	
	Show al	I your working.	
			[4]
iv.	Write ha	alf equations and an overall equation for the oxidation of $[Co(NH_3)_6]^{2+}$ to ${f C}$ by hydre in (iii).	ogen
	Half equ	ations	
	Overall	equation	
			[3]
15.	When	heated with dilute acid, $MnO_4^{2-}$ (aq) ions disproportionate into $MnO_4^-$ and $MnO_2$ .	
	i.	Balance the equation for this disproportionation reaction.	
		$MnO_4^{2-}(aq) + H^+(aq) \rightleftharpoons MnO_4^-(aq) + MnO_2(s) + H_2O(I)$	
			[1]
	ii.	Although $MnO_4^{2-}$ (aq) ions disproportionate in acidic conditions, $MnO_4^{2-}$ (aq) ions stable under alkaline solutions.	s are
		Explain this difference in stability, in terms of equilibrium.	
			[2]

- **16** This question is about reactions and properties of d-block elements.
- Iron forms many compounds and ions in the +2 and +3 oxidation states.
  - i. Complete the electron configuration of iron in its +2 oxidation state.

1s <sup>2</sup>	 	
<b>[41</b> ]		

ii. The flowchart below shows reactions of iron in its +2 and +3 oxidation states.

Complete the flowchart using formulae for reagents and iron-containing products.



[4]

**17.** Hydroxide ions, OH<sup>-</sup>, and cyanide ions, CN<sup>-</sup>, can react with some aqueous solutions of transition metal compounds.

When nickel(II) sulfate is dissolved in water, a pale green solution forms containing a six-coordinate complex ion  ${\bf C}$ .

- $\circ \quad \text{Aqueous potassium hydroxide is added to aqueous nickel(II) sulfate}.$ A green solid **D** forms.
- An excess of aqueous potassium cyanide is added to aqueous nickel(II) sulfate.

  A yellow solution forms containing a four-coordinate complex ion **E** that contains **only** nickel, carbon and nitrogen.

  i. In **C**, **D** and **E**, nickel has the +2 oxidation state. Suggest the formulae of **C**, **D** and **E**.

	Complex ion C:	
	Solid D:	
	Complex ion E:	
ii.	Write equations, and name the types o	f reaction, for the formation of <b>D</b> and <b>E</b> .
	Formation of solid <b>D</b> from aqueous nicl	kel(II) sulfate.
	Equation:	
	Type of reaction:	
	Formation of complex ion <b>E</b> from comp	lex ion <b>C</b> .
	Equation:	
	Type of reaction:	

**18(a).** Hydrated copper(II) methanoate, Cu(HCOO)<sub>2</sub>•xH<sub>2</sub>O, is a copper salt.

A student carries out the procedure below to prepare  $Cu(HCOO)_2 \cdot xH_2O$  and to determine the value of x in its formula.

#### Step 1

The student prepares Cu(HCOO)<sub>2</sub>•xH<sub>2</sub>O by reacting a copper compound with aqueous methanoic acid to form Cu(HCOO)<sub>2</sub>(aq) and allowing the solvent to evaporate.

#### Step 2

The student dissolves 2.226 g of Cu(HCOO)<sub>2</sub>•xH<sub>2</sub>O in water and makes up the solution to 250.0 cm<sup>3</sup>.

### Step 3

Using a pipette, the student adds 25.0 cm<sup>3</sup> of this solution to a conical flask followed by an excess of KI(aq).

The  $Cu^{2+}(aq)$  ions react to form a precipitate of copper(I) iodide and  $I_2(aq)$ . In this reaction, 2 mol  $Cu^{2+}$  form 1 mol  $I_2$ .

#### Step 4

The student titrates the iodine in the resulting mixture with 0.0420 mol dm<sup>-3</sup> Na<sub>2</sub>S<sub>2</sub>O<sub>3</sub> (aq).

Cu(HCOO)<sub>2</sub>•xH<sub>2</sub>O: 1s<sup>2</sup> .....

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

23.5 cm<sup>3</sup> 0.0420 mol dm<sup>-3</sup> Na<sub>2</sub>S<sub>2</sub>O<sub>3</sub> (aq) is required to reach the end point.

Complete the electron configuration of copper in

copper(I) iodide:	1s <sup>2</sup> [2]	
		at
State symbols are	not required.	
		[1]
Write an ionic equa	ation, including state symbols, for the reaction in <b>step 3</b> .	
	Choose a suitable would take place to State symbols are	CODDELLI TOUTOE.

[1]

. `	In <b>step</b>	4, the student adds a solution to observe the end point accurately.			
).	Name the solution and state the colour change at the end point.				
	Solution added:	n 			
	Colour	change:			
		[2]			
	(e).	Determine the value of $x$ in $Cu(HCOO)_2 \cdot xH_2O$ .			
		Show your working.			

19(a). This question is about the chemistry of copper compounds and complex ions.

The flowchart shows two reactions of aqueous copper(II) sulfate.

In the boxes, write the formulae and colours of the complex ions formed.

NH <sub>3</sub> (aq)		
complex ion:	complex ion:	
colour:	colour:	
		[3]

**(b).** Cu<sup>2+</sup> ions form a complex ion **A** with two ethanedioate ions and two water molecules. The ethanedioate ion is a bidentate ligand.

The skeletal formula of the ethanedioate ion is shown in Fig. 1.1 below.

i. What is meant by the term bidentate ligand?

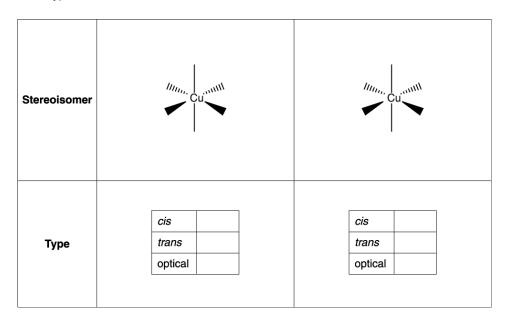
ii. The complex ion **A** exists as three stereoisomers.

The shape of one of the stereoisomers is shown below. The charge has been omitted.

## Complex A

Complete the 3D diagrams of the other two stereoisomers of  $\bf A$ . You do **not** need to include any charges.

Indicate with ticks whether the stereoisomers are *cis*, *trans*, optical or a combination of these types.



[3]

iii. What is the empirical formula, including the charge, of the complex ion A?

[2]

**20(a).** Some electrode potentials for ions are shown below.

i. Complete the electron configurations for Fe<sup>2+</sup> and Br<sup>-</sup>.

Fe<sup>2+</sup>:
1s<sup>2</sup>
Br<sup>-</sup>: 1s<sup>2</sup>

ii.	Predict the products of reacting Fe(s) separately with $I_2(aq)$ , $Br_2(aq)$ and $Cl_2(aq)$ .					
	Explain your predictions using the electrode potential data above.					
	[3]					
In this to addition	Fe <sup>2+</sup> ions can be used to test for NO <sub>3</sub> <sup>-</sup> ions. In this test, aqueous iron(II) sulfate is added to a solution containing NO <sub>3</sub> <sup>-</sup> ions, followed by slow addition of concentrated sulfuric acid. The sulfuric acid forms a layer below the aqueous solution. In the presence of NO <sub>3</sub> <sup>-</sup> ions, a brown ring forms between the two layers.					
Two rea	actions take place.					
Reactio	In the acid conditions Fe <sup>2+</sup> ions reduce NO <sub>3</sub> <sup>-</sup> ions to NO.  n 1: Fe <sup>2+</sup> ions are oxidised to Fe <sup>3+</sup> ions.  Water also forms.					
Reactio	A ligand substitution reaction of [Fe(H <sub>2</sub> O) <sub>6</sub> ] <sup>2+</sup> takes place in which one NO ligand exchanges with one water ligand. A deep brown complex ion forms as the brown ring.					
Constru	ct equations for these two reactions.					
Reactio	n 1:					
Reactio	n 2:					
	[3]					

An octanedral complex ion <b>A</b> , C9H30N6NI <sup>ST</sup> , exists as two optical isomers.	
In complex ion <b>A</b> , Ni <sup>3+</sup> is bonded to three molecules of a bidentate ligand <b>B</b> .	
i. State what is meant by a bidentate ligand.	
	[1]
ii. What is the molecular formula of the bidentate ligand <b>B</b> ?	<b>L_4</b>
	[1]
iii. Draw a possible structure for <b>B</b> and explain how <b>B</b> is able to act as a bidentate ligand.	
	[2]
iv. What is the coordination number of complex ion <b>A</b> ?	
	[1]
	r:1
v. Complete the 3-D diagrams of the shapes of the optical isomers of complex ion <b>A</b> .	
You can show the bidentate ligand simply as	
Ni Ni Ni	

•	aqueous sodium hydroxide excess aqueous ammonia hydrochloric acid.
	In your answer you should link observations with equations.

(b). Describe the reactions of EITHER aqueous copper(II) ions OR aqueous cobalt(II) ions with:

**22(a).** A student carries out an investigation to prepare and analyse a sample of barium ferrate(VI), BaFeO<sub>4</sub>. The steps in the investigation are shown below.

#### Step 1

The student adds solid iron(III) oxide to a hot aqueous solution containing an excess of hydroxide ions. The student bubbles chlorine gas through the mixture.

A solution forms containing aqueous ferrate(VI) ions, FeO<sub>4</sub><sup>2-</sup>(aq), and aqueous chloride ions.

#### Step 2

The student adds aqueous barium chloride to the resulting solution.

A precipitate of impure barium ferrate(VI) forms.

The precipitate is filtered, washed with distilled water and dried.

The student obtains 0.437 g of impure solid barium ferrate(VI).

#### Step 3

An excess of acidified aqueous potassium iodide is added to the solid from step 2.

The BaFeO<sub>4</sub> reacts as shown below, and the impurity does not react. A solution forms containing aqueous iodine,  $I_2$ (aq).

$$BaFeO_4(s) + 8H^+(aq) + 3I^-(aq) \rightarrow Fe^{3+}(aq) + Ba^{2+}(aq) + 11/2I_2(aq) + 4H_2O(l)$$

#### Step 4

(b).

The student determines the amount of  $I_2$  formed by carrying out a titration with aqueous sodium thiosulfate,  $Na_2S_2O_3(aq)$ .

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

26.4 cm<sup>3</sup> of 0.100 mol dm<sup>-3</sup> Na<sub>2</sub>S<sub>2</sub>O<sub>3</sub>(aq) are required to reach the end point.

Construct an equation for the oxidation of iron(III) oxide (step 1).

Write an <b>ionic</b> equation for the	e formation of barium ferrate(VI) ( <b>step 2</b> ).	
Include state symbols.		
		[1]

[2]

(c).	In <b>step</b>	<b>3</b> , w	hat i	s the	reducing	agent?
------	----------------	--------------	-------	-------	----------	--------

Explain your answer in terms of electrons.

reducing agent
explanation

When the solution is not alkaline, ferrate(VI) ions react with water. The reaction forms a gas with a density of 1.333 × 10 <sup>-3</sup> g cm <sup>-3</sup> , measured at room temperature and pressure, and an orange-brown precipitate.  Determine the formulae of the gas and the precipitate.		
Determine the percentage, by mass, of barium ferrate(VI) in the 0.437 g of solid formed in <b>ste</b> Give your answer to <b>one</b> decimal place.  percentage of barium ferrate(VI) =		
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Give your answer to <b>one</b> decimal place.  percentage of barium ferrate(VI) =	(d). The solid sample of barium ferrate(VI) obtained in step 2 is impure.	
percentage of barium ferrate(VI) =	Determine the percentage, by mass, of barium ferrate(VI) in the 0.437 g of solid formed in s	te
When the solution is not alkaline, ferrate(VI) ions react with water. The reaction forms a gas with a density of 1.333 × 10 <sup>-3</sup> g cm <sup>-3</sup> , measured at room temperature and pressure, and an orange-brown precipitate.  Determine the formulae of the gas and the precipitate.  Write an equation for the reaction that takes place.	Give your answer to <b>one</b> decimal place.	
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The reaction forms a gas with a density of 1.333 × 10 <sup>-3</sup> g cm <sup>-3</sup> , measured at room temperature and pressure, and an orange-brown precipitate.  Determine the formulae of the gas and the precipitate.  Write an equation for the reaction that takes place.	percentage of barium terrate(v1) =	. %
The reaction forms a gas with a density of 1.333 × 10 <sup>-3</sup> g cm <sup>-3</sup> , measured at room temperature and pressure, and an orange-brown precipitate.  Determine the formulae of the gas and the precipitate.  Write an equation for the reaction that takes place.		
write an equation for the reaction that takes place.  gas  precipitate	When the solution is not alkaline, ferrate(VI) ions react with water. The reaction forms a gas with a density of $1.333 \times 10^{-3}$ g cm <sup>-3</sup> , measured at room temperature and pressure, and an orange-brown precipitate.	
precipitate	<ul> <li>Determine the formulae of the gas and the precipitate.</li> <li>Write an equation for the reaction that takes place.</li> </ul>	
precipitate		
orecipitate	gas	
	· · · · · · · · · · · · · · · · · · ·	
εquation		
	εquation	

[2]

23(a).		nts in the d-block of the Periodic Table form ions that combine with ligands to form compl lost d-block elements are also classified as transition elements.	ex
	Explair	n why two of the Period 4 d-block elements (Sc—Zn) are <b>not</b> also transition elements.	
	In your	answer you should link full electron configurations to your explanations.	
			[6]
(b).		balt(III) ion, $Co^{3+}$ , forms a complex ion <b>A</b> with two chloride ligands and two ethanediamin $I_2CH_2NH_2$ , ligands.	e,
	The str	ructure of ethanediamine is shown below.	
		$H_2N$ $NH_2$	
	i.	Explain how ethanediamine is able to act as a bidentate ligand.	
			[2]
	ii.	Write the formula of complex ion <b>A</b> .	
			[1]
	iii.	What is the coordination number of cobalt in complex ion <b>A</b> ?	
			[1]

İ۷.

	Draw 3-D diagrams to show the three stereoisomers.	
		[3]
The ed	quilibrium reaction for the transport of oxygen by haemoglobin (Hb) in blood can be ented as <b>equation 5.1</b> .	
	$Hb(aq) + O_2(aq) \rightleftharpoons HbO_2(aq)$ equation 5.1	
i.	Explain how ligand substitution reactions allow haemoglobin to transport oxygen in blood.	1
		[2]
ii.	Write an expression for the stability constant, $K_{\text{stab}}$ , for the equilibrium involved in the transport of oxygen by haemoglobin.	he
	Use the simplified species in <b>equation 5.1</b> .	
		[1]
iii.	In the presence of carbon monoxide, less oxygen is transported in the blood.	• •
ш.	Suggest why, in terms of bond strength and stability constants.	
	Suggest why, in terms of bond strength and stability constants.	
		[2]

Complex ion **A** has *cis* and *trans* stereoisomers. One of these stereoisomers also has an optical isomer.

24(a).	<ol> <li>A redox reaction takes place when copper metal is heated with concentrated sulfuric acid. A solution forms and 95.0 cm<sup>3</sup> of a colourless gas is collected, measured at RTP. The gas has mass of 254 mg.</li> </ol>						
	i.	Write the electron configuration, in terms of sub-shells, for a copper atom.					
		<u>[1]</u>					
	ii.	Suggest the identity of the colourless gas and write an equation for the reaction taking place.					
		State symbols are <b>not</b> required in the equation.					
		Show your working for calculations.					
		gas:					
		equation:					

(b). A student carries out two experiments based on redox reactions of iron and chromium.

Use the standard electrode potentials below to help you answer the questions that follow.

[4]

Fe <sup>2+</sup> (aq) + 2e <sup>-</sup>	$\rightleftharpoons$	Fe(s)	<i>E</i> ⊖ = −0.44 V
2H+ (aq) + 2e-	$\rightleftharpoons$	H2(g)	<i>E</i> ⊖ = 0.00 V
Fe <sup>3+</sup> (aq) + e <sup>-</sup>	$\rightleftharpoons$	Fe <sup>2+</sup> (aq)	<i>E</i> ⊖ = +0.77 V
O <sub>2</sub> (g) + 4H <sup>+</sup> (aq) + 4e <sup>-</sup>	$\rightleftharpoons$	$2H_2O(I)$	<i>E</i> ⊖ = +1.23 V
$Cr_2O_7^{2-}(aq) + 14H^+(aq) + 6e^-$	$\rightleftharpoons$	2Cr <sup>3+</sup> (aq) + 7H <sub>2</sub> O(I)	<i>E</i> ⊖ = +1.33 V
$CI_2(g) + 2e^-$	$\rightleftharpoons$	2C <i>l</i> ⁻(aq)	<i>E</i> ⊖ = +1.36 V
$H_2O_2(aq) + 2H^+(aq) + 2e^-$	$\rightleftharpoons$	2H <sub>2</sub> O(I)	<i>E</i> ⊖ = +1.78V

For each experiment, identify the species causing the observations shown in bold text and write overall equations for any reactions taking place.

State symbols are **not** required in the equations.

- i. Experiment 1
  - The student adds iron filings to dilute hydrochloric acid.
     A green solution forms and gas bubbles are seen.
  - 2. The student bubbles air through the green solution. The solution turns an **orange-brown colour**.

	1:	
	2:	
		[3]
ii.	Experiment 2  The student heats a green solution of chromium(III) sulfate with dilute acid and hydrogen peroxide, H <sub>2</sub> O <sub>2</sub> .  The solution turns an orange colour.	
		[3]

25(a).	This question is about the chemistry of transition elements.	
	Many chromium compounds contain chromium in the +3 oxidation state.	
	Complete the electron configurations of chromium as the element and in the +3 oxidation st	tate.
	Chromium as the element:	
	1s <sup>2</sup> 2s <sup>2</sup> 2p <sup>6</sup>	
	Chromium in the +3 oxidation state: 1s <sup>2</sup> 2s <sup>2</sup> 2p <sup>6</sup>	
		[2]

(b). Compound I is a complex with the empirical formula CoN<sub>4</sub>H<sub>12</sub>Cl<sub>3</sub>.

The formula of compound  ${\bf I}$  contains one chloride ion and a complex ion  ${\bf J}$ , which has two stereoisomers.

Draw and label the three-dimensional structures of the two stereoisomers of complex ion  ${\bf J}$ . Include the charge of the complex ion in your diagrams.

[3]

(c). A and B are compounds of two different transition elements.

Two students carry out test-tube tests on aqueous solutions of  ${\bf A}$  and  ${\bf B}$ . They then analyse the results to identity  ${\bf A}$  and  ${\bf B}$ .

The observations of **Student 1's** tests are shown below.

	Test	A(aq)	B(aq)
1	NH <sub>3</sub> (aq) added dropwise	green precipitate <b>C</b>	pale-brown precipitate <b>E</b>
	Excess NH <sub>3</sub> (aq) added	violet solution <b>D</b>	no further change
2	HNO₃(aq)	no change	no change
	followed by Ba(NO <sub>3</sub> ) <sub>2</sub> (aq)	white precipitate <b>F</b>	no change
	HNO₃(aq)	no change	no change
3	followed by AgNO <sub>3</sub> (aq)	no change	yellow precipitate, <b>G</b>

I.	Analyse the results and answer the following.
	Identify <b>A</b> and <b>B</b> and write their formulae.
	Construct ionic equations for the formation of the products <b>C–G</b> from the relevant ions present in <b>A(aq)</b> and <b>B(aq)</b> .
ii.	What was the purpose of adding HNO <sub>3</sub> (aq) first in <b>Test 2</b> and <b>Test 3</b> ?
	[ <sup>*</sup>
iii.	Student 2 accidentally used HC/(aq) instead of HNO <sub>3</sub> (aq) for both Test 2 and Test 3.
	What different observations would <b>Student 2</b> obtain? Explain your reasoning.
	[(

	[1	<u>1]</u>
iv.	How could the procedure be modified to be more certain of the conclusions from <b>Test 3</b>	?

**END OF QUESTION PAPER**