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Transition Metal Basics

Question Paper

Level	International A Level
Subject	Chemistry
Exam Board	Edexcel
Topic	Transition Metals & Organic Nitrogen Chemistry
Sub Topic	Transition Metal Basics
Booklet	Question Paper

Time Allowed: 86 minutes

Score: /71

Percentage: /100

Grade Boundaries:

A*	Α	В	С	D	E	U
>85%	'77.5%	70%	62.5%	57.5%	45%	<45%

1	Which of the following elements is both a d-block element and a transition element?						
	■ A Calcium						
	⊠ B	Copper					
	⊠ C	Scandiu	um				
		Zinc					
					(Total for Question 1 = 1 mark)		
2 T				ation energies (in kJ mol ⁻ ansition element?	⁻¹) of four different elements.		
2 T		ould be		_	⁻¹) of four different elements.		
2 T	Which c	ould be	e those of a tra	ansition element?	⁻¹) of four different elements.		
2 T	Which c	ould be	e those of a tra	ansition element?	⁻¹) of four different elements.		
2 T	Which c	658 578	e those of a tra 1310 18	ansition element?	⁻¹) of four different elements.		
2 ⊤	Which c	658 578 738	e those of a tra 1310 18 14	ansition element?	(Total for Question 2 = 1 mark)		

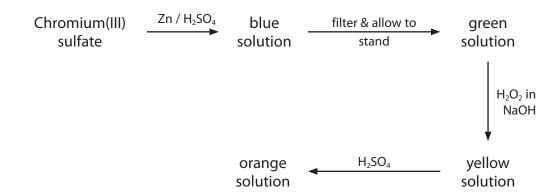
3 The electronic configuration of the element vanadium is

			3d	4s
	⊠ A	[Ar]	↑↓ ↑↓ ↑	
	ВВ	[Ar]	↑ ↑	
	⋈ C	[Ar]	↑	<u> </u>
	⊠ D	[Ar]	<u> </u>	11
				(Total for Question 3 = 1 mark)
4			als typically form a number of pest explanation for this prope	ions which are stable in aqueous erty is that
	⊠ A		erences in the successive ioniz	zation energies are similar to the f the ions.
	⊠ B	all the i	ons are formed by the remova	al of electrons from the d subshell.
	⊠ C	the ioni	ization energies of transition r	metals are low.
	■ D	-	Iration enthalpies of transition ose of ions of s and p block m	metal ions are always more exothermic etals.
				(Total for Question 4 = 1 mark)
5	The ele	ectronic c	configuration of the iron(II) ion	n, Fe ²⁺ , is
			3d	4s
	⋈ A	[Ar]	$\uparrow\downarrow$ \uparrow \uparrow	$\uparrow\downarrow$
	⋈ B	[Ar]	<u> </u>	$\uparrow\downarrow$
	⊠ C	[Ar]	<u> </u>	\uparrow
	☑ D	[Ar]	↑↓ ↑ ↑ 	
				(Total for Question 5 = 1 mark)

5			ion metal compounds often have catalytic properties. The best explanation is that						
	X	Α	transition metal compounds usually have a much larger surface area than other metal compounds.						
	×	В	transition metal ions readily promote electrons to higher energy levels by absorbing electromagnetic radiation in the visible region.						
	X	C	relatively small amounts of ene of a transition metal.	rgy are required to change t	he oxidation state				
	X	D	the ionization energies of trans other metals.	ition metals are much lower	than those of				
				(Total for	Question 6 = 1 mark)				
7 \			the shapes of the dichlorocupra llorochromate(III) ion, CrCl ₄ ?	ate(I) ion, CuCl ₂ , and the					
			CuCl ₂	$CrCl_4^-$					
	X	A	V shaped	tetrahedral					
	X	В	linear	tetrahedral					
	×	C	V shaped	square planar					
	×	D	linear	square planar					
				(Total for	Question 7 = 1 mark)				
	The iron(II) ion forms complexes with monodentate ethanoate ions and bidentate ethanedioate ions. The complexes with ethanedioate ions are more stable. What is the best explanation for this?								
	X	A	Ethanedioate ions form stronge	er bonds than ethanoate ion	s with iron(II) ions.				
	X	В	Ethanedioic acid is a stronger a	cid than ethanoic acid.					
	X	C	The formation of the ethanedic	oate complex produces more	particles in solution.				
	X	D	Ethanedioic acid forms stronge	r hydrogen bonds than etha	noic acid.				
				(Total for	Question 8 = 1 mark)				

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9 The diagram below summarises a sequence of reactions involving chromium compounds.

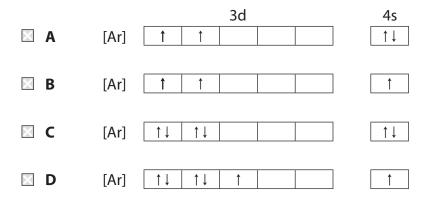


How many different oxidation states of chromium are involved in this sequence?

- **B** 3
- **◯ C** 4
- **D** 5

(Total for Question 9 = 1 mark)

10 The electronic configuration of the element chromium is

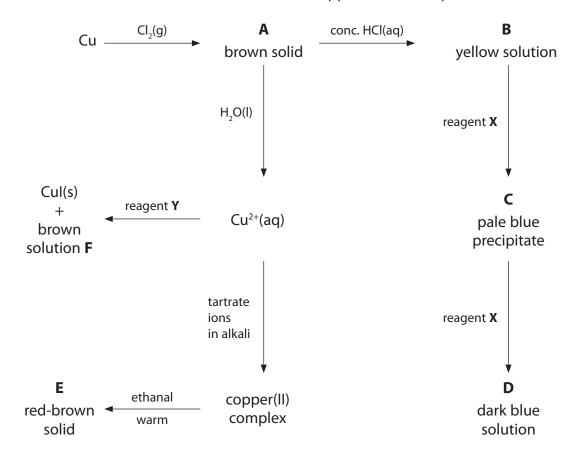


(Total for Question 10 = 1 mark)

11	In wh	nich of th	ne following compounds does iro	on have the highest oxidation number?
		Fe ₃ O ₄		
	⊠ В	K ₂ FeO ₄		
		Na ₄ Fe(C	CN) ₆	
	⊠ D	Na ₃ Fe(C	CN) ₆	
				(Total for Question 11 = 1 mark)
12	What	t is the e	lectronic configuration of the Fe	i+ ion?
			3d	4s
	⊠ A	[Ar]	↑ ↑	
	⊠ B	[Ar]	$\begin{array}{ c c c c c c c c c c c c c c c c c c c$	
	⊠ C	[Ar]	<u> </u>	<u> </u>
	⊠ D	[Ar]	<u> </u>	$\uparrow\downarrow$
				(Total for Question 12 = 1 mark)
13	polluta	ant wher		biomass fuels and is a significant mosphere. One method of removing nium(IV) oxide catalyst.
	The b e	est expla	nation for the use of titanium(IV) oxide is that the titanium
	⊠ A	contain	s many active sites on which the	reaction can occur.
	⊠ B		ly oxidized to a higher oxidation ation state +4.	state which can then be reduced back
	⊠ C		ly reduced to a lower oxidation s ation state +4.	tate which can then be oxidized back
	⊠ D	has par	tially filled d orbitals in its +4 oxi	dation state.
				(Total for Question 13 = 1 mark)

	A transition metal ion, M, forms a complex with a bidentate ligand, B. The formula of the complex is MB_3 so the shape of the complex is most likely to be							
×	Α	trigonal planar.						
X	В	pyramidal.						
×	C	trigonal bipyramidal.						
X	D	octahedral.						
		(Total for Question 14 = 1 mark)						
		,						
		h of the following lists all the types of bond that are present in a crystalline of the compound tetraamminecopper(II) sulfate?						
		h of the following lists all the types of bond that are present in a crystalline						
	mple	h of the following lists all the types of bond that are present in a crystalline e of the compound tetraamminecopper(II) sulfate?						
saı	mple A	h of the following lists all the types of bond that are present in a crystalline of the compound tetraamminecopper(II) sulfate? Ionic, covalent and dative covalent						
sai	mple A B	h of the following lists all the types of bond that are present in a crystalline of the compound tetraamminecopper(II) sulfate? Ionic, covalent and dative covalent Ionic and dative covalent						

16 The scheme below summarises some reactions of copper and its compounds.



(a) (i) Identify the copper containing species **A** to **E** either by name, including the oxidation number, or by formula. Also, identify the brown solution, **F**.

A B C C D E (ii) Identify the reagents X and Y. (2)

Υ

(6)

(iii) Identify the organic product of the reaction between the copper(II) complex and ethanal. Hence explain the role of ethanal in the reaction.	(2)
(iv) The reaction between Cu ²⁺ (aq) and reagent Y forms Cul(s) and a brown solution, F . This reaction is the first stage in a method for the determination of the concentration of Cu ²⁺ (aq).	
Outline briefly how this method is used. Practical details are not required.	(1)
*(b) (i) Explain why B is coloured.	(4)
(ii) Explain why B and D have different colours.	(2)

(c)	Aq	ueous copper(I) ions undergo a disproportionation reaction.	
	(i)	Write the ionic equation for this reaction. Include state symbols in your answer.	(1)
	(ii)	Explain, stating the relevant oxidation numbers, why the reaction in (c)(i) is classified as a disproportionation.	(1)
	(iii)	se the standard reduction potentials on page 17 of the Data Booklet to calculate $E_{\rm cell}^{\ominus}$ for this disproportionation. Hence show that this reaction is thermodynamically feasible.	(2)
		(Total for Question 16 = 21 ma	rks)

17	This qu	uestion is about the element chromium and some of its compounds.	
	(a) (i)	Complete the electronic configuration of the chromium atom, using the s, p, d notation.	(1)
		[Ar]	
	*(ii)	State how this electronic configuration of the chromium atom is unusual compared with most other transition metals.	
		Give two reasons why chromium has this electronic configuration.	(2)

(1	b) (i)	A solution containing chromium(II) ions can be produced in the laboratory by reducing a solution of potassium dichromate(VI) using zinc in 50% hydrochloric acid. This reaction takes place in two steps. Dichromate(VI) ions are reduced to chromium(III) ions and the chromium(III) ions are then reduced to chromium(II) ions.	
		Use the relevant standard reduction potentials from page 17 or standard electrode potentials on pages 14 and 16 of the Data Booklet to calculate $E_{\rm cell}^{\ominus}$ for each step.	
		Use your $E_{\text{cell}}^{\ominus}$ values to explain why both steps are spontaneous.	(3)
	(ii)	State the colour changes you would expect to see during this reaction.	(1)
	(iii)	A student suggests that the hydrogen produced by the reaction of the zinc with hydrochloric acid in this experiment is a serious risk. Evaluate the student's suggestion.	(1)

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(c) Chromium(II) ions in aqueous solution are quickly oxidized by the oxygen in air. One method of stabilising chromium(II) ions is by adding a solution of sodium ethanoate, forming a complex, $[Cr_2(CH_3CO_2)_4(H_2O)_2]$. This complex may be represented by the structure below.

(i)	What type of ligand is the ethanoate ion in this complex?	(1)
(ii)	State the type of bonding which occurs between the ligands and the chromium(II) ions.	(1)
(iii)	Suggest two unusual features in the structure and bonding of this complex.	(2)
(iv)) This complex is red. Explain why the colour of chromium(II) ethanoate is different from that of $Cr(H_2O)_6^{2+}$ (aq).	(2)

(v) Predict the number and relative areas of the peaks in the \boldsymbol{low} resolution

proton nmr spectrum of $Cr_2(CH_3CO_2)_4(H_2O)_2$.

Justify your answers.	
	(2)

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(d) An experiment is carried out to check the oxidation number of chromium in chromium(II) ethanoate $Cr_2(CH_3CO_2)_4(H_2O)_2$.

 $1.00 \text{ g} (2.66 \times 10^{-3} \text{ mol}) \text{ of chromium(II)}$ ethanoate is dissolved in 25 cm³ of 1.00 mol dm^{-3} sulfuric acid.

The solution is diluted with distilled water until the volume is 250 cm³.

25.0 cm³ portions of the diluted solution are titrated with 0.00750 mol dm⁻³ potassium manganate(VII).

Calculate the volume of potassium manganate(VII) needed to oxidize the chromium(II) ions present in each 25.0 cm 3 portion to the +6 oxidation state. The manganese is reduced to the +2 oxidation state.

Comment on your answer and suggest how the experiment could be improved to give a more suitable titre.

(5)

Comment	
	(Total for Question 17 = 21 marks)

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(3)

- **18** Crystals of copper(II) sulfate dissolve in water to form a blue solution, **A**. When dilute aqueous ammonia is added to this solution, a pale blue precipitate, **B**, forms which dissolves in excess aqueous ammonia to form a dark blue solution, **C**.
 - (a) (i) Give the **formula** of the copper species in **A**, **B** and **C**. You should include all of the ligands present in each species.

(**) -	
(ii) Explain why solution A is coloured.	(4)
	(-)
(iii) Explain why solution A is a different colour to solution C .	
(iii) Explain why solution A is a different colour to solution C .	(2)

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(b) A more concentrated solution of **C** may be prepared by using concentrated aqueous ammonia in place of dilute aqueous ammonia. The crystalline sulfate of **C** may be obtained by cooling the mixture in an ice bath and adding ethanol. The filtered crystals may be recrystallized using ethanol as the solvent.

The steps of the recrystallization are summarised below. In the spaces provided, explain the purpose of each step, referring particularly to any words in **bold** type.

(5)

	(Total for Question 18 = 14 marks)
Step 4	The mixture was filtered using suction filtration .
Step 3	The filtrate was cooled in an ice bath .
Step 2	The hot solution was filtered .
Step 1	The solid was dissolved in the minimum amount of hot ethanol.