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Surname	Other n	ames
Pearson Edexcel International Advanced Level	Centre Number	Candidate Number
Chemistry	/	
Advanced Unit 5: General Principle	es of Chemistry II Inic Nitrogen Chei	
Advanced Unit 5: General Principle Metals and Orga	es of Chemistry II nnic Nitrogen Che otic assessment)	Paper Reference
Advanced Unit 5: General Principle Metals and Orga (including synop	es of Chemistry II nnic Nitrogen Che otic assessment)	mistry

Instructions

- Use **black** ink or **black** ball-point pen.
- **Fill in the boxes** at the top of this page with your name, centre number and candidate number.
- Answer **all** questions.
- Answer the questions in the spaces provided
 - there may be more space than you need.

Information

- The total mark for this paper is 90.
- The marks for **each** question are shown in brackets
 - use this as a guide as to how much time to spend on each question.
- Questions labelled with an asterisk (*) are ones where the quality of your written communication will be assessed
 - you should take particular care with your spelling, punctuation and grammar, as well as the clarity of expression, on these questions.
- A Periodic Table is printed on the back cover of this paper.

Advice

- Read each question carefully before you start to answer it.
- Try to answer every question.
- Check your answers if you have time at the end.
- Show all your working in calculations and include units where appropriate.

Turn over ▶





SECTION A

Answer ALL the questions in this section. You should aim to spend no more than 20 minutes on this section. For each question, select one answer from A to D and put a cross in the box ⊠. If you change your mind, put a line through the box ₩ and then mark your new answer with a cross ⋈.

1 Which electrode and solution would be used to measure the standard electrode potential of the manganese(III)/manganese(II) half-cell?

$$Mn^{3+}(aq) + e^{-} \rightleftharpoons Mn^{2+}(aq)$$

		Electrode	Solution
×	Α	manganese	1 mol dm ⁻³ Mn ³⁺ (aq)
×	В	manganese	1 mol dm ⁻³ with respect to Mn ³⁺ (aq) and Mn ²⁺ (aq)
×	C	platinum	1 mol dm ⁻³ Mn ³⁺ (aq)
X	D	platinum	1 mol dm ⁻³ with respect to Mn ³⁺ (aq) and Mn ²⁺ (aq)

(Total for Question 1 = 1 mark)

- 2 In which pair of species are the oxidation numbers of the d-block elements the same?
 - \triangle **A** $[Cr(NH_3)_4Cl_2]^+$ and $[Mn(H_2O)_6]^{2+}$
 - B CrO₄²⁻ and TiCl₃
 - \square **C** Cr_2O_3 and $[Fe(CN)_6]^{3-}$
 - \square **D** $\operatorname{Cr}_2\operatorname{O}_7^{2-}$ and MnO_4^-

(Total for Question 2 = 1 mark)

3 One mole of metal ions reacted in solution with one mole of sulfur dioxide. The half-equation for the sulfur dioxide reaction is

$$SO_2(aq) + 2H_2O(l) \rightarrow SO_4^{2-}(aq) + 4H^+(aq) + 2e^-$$

The original oxidation number of the metal was +3. What was the oxidation number of the metal after the reaction?

- \blacksquare **B** +2
- **◯ C** +4
- **D** +5

(Total for Question 3 = 1 mark)

- 4 This question concerns four complexes.
 - (a) Which complex has a tetrahedral structure?

(1)

- \triangle A [CrCl₄]
- \boxtimes **B** $[CuCl_2]^-$
- \square **C** [Pt(NH₃)₂Cl₂]
- \square **D** $[TiCl_6]^{2-}$
- (b) Which complex contains a metal in the +1 oxidation state?

(1)

- \triangle **A** $[CrCl_4]^-$
- \boxtimes **B** [CuCl₂]⁻
- \square **C** [Pt(NH₃)₂Cl₂]
- \square **D** $[TiCl_6]^{2-}$

(Total for Question 4 = 2 marks)



- **5** An ion of metal M has a charge of +n. It forms a complex ion with a charged bidentate ligand, L.
 - (a) The formula of the complex ion formed between the metal ion and the bidentate ligand is $[ML_2]^{(+n-4)}$. What is the charge on ligand L?

(1)

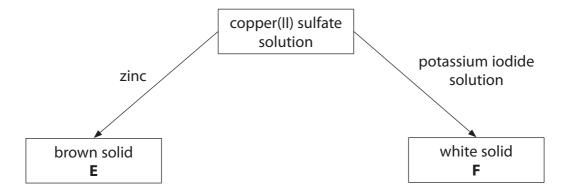
- \triangle A +2
- B 0
- **区** −2
- D -4
- (b) Another complex ion can be formed in which both of the bidentate ligands L, in $[ML_2]^{(+n-4)}$, are replaced by the neutral monodentate ligand Z. What is the formula of the complex ion?

(1)

- \blacksquare **A** $[MZ_2]^{n+}$
- \boxtimes **B** $[MZ_2]^{2n+}$
- \square **C** $[MZ_4]^{n+}$
- \square **D** $[MZ_4]^{2n+}$

(Total for Question 5 = 2 marks)

6 Two reactions of copper(II) sulfate solution are shown.



(a) What is the insoluble brown solid **E**?

(1)

- ☑ B Copper(I) oxide
- ☑ D Zinc sulfate
- (b) What is the insoluble white solid **F**?

(1)

- **A** Copper
- ☑ B Copper(I) iodide
- **D** Potassium sulfate

(Total for Question 6 = 2 marks)

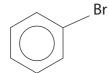
- **7** When benzene reacts with a mixture of concentrated nitric and sulfuric acids, the reaction is
 - ☑ A electrophilic addition.
 - **B** electrophilic substitution.
 - **C** nucleophilic addition.
 - **D** nucleophilic substitution.

(Total for Question 7 = 1 mark)

- **8** Benzene and phenol react with bromine.
 - (a) What is the organic product when benzene reacts with excess bromine in the presence of ultraviolet light?

(1)

A



⊠ B

Br

⊠ C

□ D

Br

electrophile.

(b) Bromine reacts more readily with phenol than with benzene. This is because the (1)
A benzene ring in phenol is more susceptible to nucleophilic attack.
B benzene ring in phenol is deactivated because the oxygen of the OH group is very electronegative.
C lone pair of electrons on the oxygen atom in phenol overlap with the delocalised electrons in the benzene ring.
D lone pair of electrons on the oxygen atom enable phenol to act as an

(Total for Question 8 = 2 marks)

9 The structure of compound **G** is

(a) What is the systematic name of compound **G**?

(1)

- ☑ A 4-methylbenzene-1,3-diamine
- B 4-methylbenzene-1,5-diamine
- **D** 4-methylphenyldiamine
- (b) What is the organic species formed in the reaction between compound **G** and **excess** ethanoyl chloride?

(1)

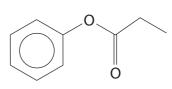
 CH_3

CH₃

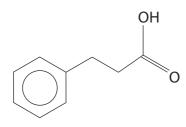
(Total for Question 9 = 2 marks)

10 Four different compounds, **P**, **Q**, **R** and **S**, are structural isomers with molecular formula $C_9H_{10}O_2$.

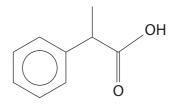
compound P



compound Q



compound **R**



compound S

(a) Which compound does **not** exhibit optical isomerism but does react with sodium hydrogencarbonate to give a colourless gas?

(1)

- A Compound P
- B Compound Q
- $oxed{oxed}$ Compound R
- D Compound S
- (b) Which compound reacts with sodium hydroxide solution to give sodium benzoate as one of the products?

(1)

- A Compound P
- ☑ B Compound Q
- C Compound R
- ☑ D Compound S

(Total for Question 10 = 2 marks)

0

11 A reaction sequence is shown.

compound
$$\mathbf{W}$$

$$\begin{array}{c}
\text{Step 1} \\
\text{Cr}_2O_7^{2-}/\text{H}^+
\end{array}$$

$$\begin{array}{c}
\text{Step 2} \\
\text{O}
\end{array}$$

$$\begin{array}{c}
\text{Step 3}
\end{array}$$

- (a) What is compound **W**?
 - (1)
- A Butan-1-ol
- B Butan-2-ol
- C 2-methylpropan-1-ol
- D 2-methylpropan-2-ol
- (b) Which substances are required for Step 2?

(1)

- A Acidified potassium dichromate(VI)
- B lodine in alkali, followed by hydrochloric acid
- Sodium hydroxide solution followed by hydrochloric acid
- D Ammoniacal silver nitrate (Tollens' reagent)
- (c) Which is the reagent for Step **3**?

- **A** Aqueous chlorine
- **B** Chlorine gas
- C Hydrochloric acid
- D Phosphorus(V) chloride



(d) Which is the reagent for Step 4?

(1)

- A CH₃CH₂CH₂CH₂NH₂
- B CH₃CH(NH₂)CH₂CH₃
- C CH₃CH₂CH₂CONH₂
- ☑ D CH₃CH(CONH₂)CH₃

(Total for Question 11 = 4 marks)

TOTAL FOR SECTION A = 20 MARKS

SECTION B

Answer ALL the questions. Write your answers in the spaces provided.

- **12** Aluminium and iron are both metallic elements. There are similarities and differences in the properties of their compounds.
 - (a) Both elements form compounds in which their oxidation number is +3.
 - (i) Complete the electronic configuration of the ${\rm Al}^{3+}$ and ${\rm Fe}^{3+}$ ions, using the s, p, d notation.

(2)

(ii) Aluminium only forms compounds in which its oxidation number is +3, whereas iron has compounds with a variety of oxidation numbers.

Suggest a reason why iron forms stable compounds with more than one oxidation number.

(1)

*(b) Most aluminium compounds are colourless but iron(III) compounds a	are coloured

Explain why $[Fe(H_2O)_6]^{3+}$ ions are coloured.

(4)



- (c) Aluminium ions and iron(III) ions form complexes in solution. These solutions are acidic.
 - (i) Draw the structure of the $[Fe(H_2O)_6]^{3+}$ complex ion, showing clearly the shape around the Fe^{3+} ion, and which atoms in the ligands are attached to the Fe^{3+} ion.

(2)

*(ii) The following equilibrium occurs in aqueous solution

$$[Fe(H_2O)_6]^{3+} + H_2O \implies [Fe(OH)(H_2O)_5]^{2+} + H_3O^+$$

Suggest why one of the water ligands loses a proton.

(2)



(d) Aluminium chloride, AlCl₃, and iron(III) chloride, FeCl₃, can both be used as catalysts in Friedel-Crafts reactions.

The reaction between benzene and chloromethane, using an aluminium chloride catalyst, can be summarised as

$$C_6H_6 + CH_3Cl + AlCl_3 \rightarrow C_6H_6CH_3^+ + AlCl_4^-$$

 $C_6H_6CH_3^+ + AlCl_4^- \rightarrow C_6H_5CH_3 + HCl + AlCl_3$

Suggest, by reference to the electronic structure of AlCl₃, how the AlCl₄ ion forms.

(2)

- (e) Thiocyanate ions, SCN $^-$, are used to test for the presence of Fe $^{3+}$ ions in aqueous solution. A blood red colour, caused by the complex ion $[Fe(SCN)(H_2O)_5]^{2+}$, is seen.
 - (i) State the type of reaction taking place.

(1)

(ii) Draw a dot-and-cross diagram of the thiocyanate ion, [SCN]. Hence suggest a structure of the ion, showing all the bonds and which atom has the negative charge.

(2)

(iii) Suggest **two** ways in which the thiocyanate ion could bond to the Fe³⁺ in the complex.

(1)

(f) Aluminium hydroxide, Al(OH)₃, is amphoteric.

Write **ionic** equations for the reactions of aluminium hydroxide with hydrochloric acid and with sodium hydroxide solution.

State symbols are not required.

(2)

lonic equation with hydrochloric acid

(Total for Question 12 = 19 marks)



- **13** This question is about carboxylic acids.
 - (a) An organic compound, **T**, contains the elements carbon, hydrogen and oxygen only. **T** contains a carboxylic acid group and one other functional group.
 - *(i) A sample of compound **T** of mass 2.25 g was burned completely, producing 3.30 g of carbon dioxide and 1.35 g of water. In the mass spectrum of compound **T**, the molecular ion peak is at m/e = 90.

Use all the data to calculate the molecular formula of compound **T**. You **must** show your working.

(6)

(ii)	The mass spectrum of compound T has a peak at $m/e = 45$. Give the displayed formulae of two species that could produce this peak.	(2)
(iii)	The low resolution proton nmr spectrum of compound T has four peaks with areas in the ratio 1:2:2:1. Draw the structure of compound T and explain how your structure is consistent with the proton nmr data.	(3)



(b) The structure of 2-hydroxy-2-phenylpropanoic acid is shown.

- *(i) Outline how 2-hydroxy-2-phenylpropanoic acid can be synthesised in **three** steps starting from benzene and an acyl chloride.
 - Include the reagents for each step in the synthesis and draw the structures of the two organic intermediates.

(5)

(ii) Poly(2-hydroxy-2-phenylpropanoic acid) is a possible biodegradable polymer.

Draw a section of this polymer, showing **two** repeat units.

(2)

(Total for Question 13 = 18 marks)

- **14** This question is about redox reactions.
 - (a) Vanadium exists in different oxidation states which can be interconverted using suitable oxidising and reducing agents.

Some relevant standard electrode potentials are shown in the table.

Half-equation	E [⊕] /V
$Zn^{2+}(aq) + 2e^- \rightleftharpoons Zn(s)$	-0.76
$V^{3+}(aq) + e^- \rightleftharpoons V^{2+}(aq)$	-0.26
$SO_4^{2-}(aq) + 4H^+(aq) + 2e^- \Longrightarrow SO_2(aq) + 2H_2O(l)$	+0.17
$VO^{2+}(aq) + 2H^{+}(aq) + e^{-} \rightleftharpoons V^{3+}(aq) + H_2O(l)$	+0.34
$Fe^{3+}(aq) + e^{-} \rightleftharpoons Fe^{2+}(aq)$	+0.77
$VO_{2}^{+}(aq) + 2H^{+}(aq) + e^{-} \rightleftharpoons VO^{2+}(aq) + H_{2}O(l)$	+1.00

(i) Write the overall half-equation for the oxidation of vanadium(II) ions to the vanadium(V) ions, VO_2^+ . State symbols are not required.

(1)

(ii) Select, from the table, a reducing agent which should reduce vanadium(V) to vanadium(IV) but not to vanadium(III). Justify your answer.

(3)

(iii) Write the ba	alanced equation	for the disp	proportionati	ion of V ³⁺	into V ²⁺	and '	VO ²⁺
•	whether this rea		sible under st	tandard c	condition	ns.	
State symbo	ols are not require	ed.					

(3)

(b) Excess potassium iodide solution was added to 25.0 cm³ of an aqueous solution of bromine.

$$Br_2 + 2I^- \rightarrow 2Br^- + I_2$$

The iodine produced was titrated with sodium thiosulfate solution. All of the iodine required 24.20 cm³ of 0.100 mol dm⁻³ sodium thiosulfate solution for reaction.

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

(i) Calculate the concentration of the aqueous solution of bromine.

(2)

(ii) In another experiment, 100.0 cm³ of the same aqueous solution of bromine was treated directly with 0.00100 moles of sodium thiosulfate.

Excess potassium iodide was then added to the unreacted bromine.

The iodine produced required 16.80 cm³ of the 0.100 mol dm⁻³ sodium thiosulfate solution for reaction.

Deduce the mole ratio of $S_2O_3^{2-}$: Br_2 and hence write a balanced equation for the reaction between thiosulfate ions and bromine. State symbols are not required.

(5)

(Total for Question 14 = 14 marks)

TOTAL FOR SECTION B = 51 MARKS



SECTION C

Answer ALL the questions. Write your answers in the spaces provided.

15

Sweeteners

Table sugar (sucrose) contributes to tooth decay and obesity, so some manufacturers add other natural or artificial sweeteners to food.

Honey is a natural sweetener and has about the same relative sweetness as sucrose. The sweetness arises from the simple sugars glucose and fructose. Honey also contains pinocembrin which is an antioxidant.

Saccharin is an artificial sweetener, which was first produced in 1879. It is over 300 times sweeter than sucrose.

It is normally used as the sodium salt, which is very soluble in water.

Aspartame is an ester of the dipeptide formed from the amino acids phenylalanine and aspartic acid. It was first produced in 1965 and is about 200 times sweeter than sucrose.

(a) (i) Give the molecular formula for pinocembrin.

(1)

(ii) Label the chiral carbon atom in pinocembrin with an asterisk (*).

(1)

(b) One route for the synthesis of saccharin starts with the reaction between methylbenzene and chlorosulfonic acid, HSO₃Cl.

Draw the mechanism for this reaction.

The electrophile is *SO₂Cl.

(3)

(c) Another route for the synthesis of saccharin starts with the reaction between methyl 2-aminobenzoate and nitrous acid (prepared from sodium nitrite and hydrochloric acid).

SO₂NH₂

saccharin

(i) Give the structure of the Intermediate ${\bf A}.$

(1)

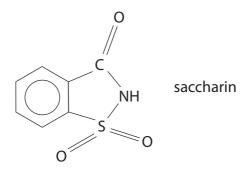
SO₂Cl

(ii) Suggest a substance for Reagent **B** in Step **4**.

(1)

(iii) Complete the equation for the formation of saccharin in Step 5.

(iv) The sodium salt of saccharin is much more soluble in water than saccharin.



The acidic hydrogen is the one that is attached to the nitrogen atom.

Complete the structure of the sodium salt of saccharin, showing the charges on both ions.



(d) Phenylalanine and aspartic acid are amino acids.

$$\begin{array}{c} O \\ O \\ NH_2 \end{array}$$

(i) Draw the structure of the organic compound formed when hydrochloric acid is added to phenylalanine.

(1)

(ii) Draw the structure of the organic product formed when **excess** sodium hydroxide solution is added to aspartic acid.

(iii) The structure of aspartame is

aspartame

Draw the structure of the dipeptide formed between phenylalanine and aspartic acid that is required to form aspartame.

(2)

(iv) Identify, by name or formula, the alcohol needed to form aspartame from the dipeptide in (d)(iii).



(e) Sugar alcohols, such as sorbitol, are also used as artificial sweeteners. Sorbitol is made from glucose.

(i) State the reagent needed to convert glucose into sorbitol.

(1)

(ii) Describe a chemical test which will distinguish between glucose and sorbitol. Give the result for each substance.

(3)

(iii) Sorbitol is dehydrated to form sorbitan. Sorbitan contains a ring with four carbon atoms and one oxygen atom. Complete the formula of sorbitan.

(1)



(Total for Question 15 = 19 marks)

TOTAL FOR SECTION C = 19 MARKS TOTAL FOR PAPER = 90 MARKS



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5		1.0 H hydrogen					6)	(8)	55.8	ē	iron 26	101.1	Ru	ruthenium 44	190.2	o O	osmium 76	[277]		hassium r		150		
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•			nass	loc	umber		193	(0)	52.0	ხ	chromium manganese	95.9	Wo	nolybdenum 42	183.8		tungsten 74	[392]	Sg	seaborgium 106		144	PN	neodymium p
		Key	relative atomic mass	atomic symbol	name atomic (proton) number		(5)	(c)	50.9		vanadium 23	92.9		niobium 41	180.9	Та	tantalum 73	[292]		dubnium 105		141	P	жазеодутіцт 59
			relati	ato	atomic		5	(4)	47.9		titanium 22	91.2	Zr	zirconium 40	178.5	Ŧ	hafnium 72	[261]	₹	rutherfordium 104		140		cerium 58
							ę	(5)	45.0	S	scandium 21	88.9		yttrium 39	138.9	La*	lanthanum 57	[227]		actinium 89			Ş	
	7	(2)	9.0	Be	beryllium 4	24.3	Mg magnesium	12	40.1	g	calcium 20	97.8	Ş	strontium 38	137.3		barium 56	[326]	Ra	radium 88			* Lanthanide series	* Actinide series
	-	(£)	6.9	=	lithium 3	23.0	_ E	=	39.1	×	potassium 19	85.5		rubidium 37	132.9	S	caesium 55	[223]	Ŀ	francium 87			* Lanth	* Actinic
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175	Ľ	lutetium	71	[257]	ב	lawrencium	103
173	Х	ytterbium	70	[254]	å	nobelium	102
169	T	thulium	69	[526]	PW	mendelevium	101
167	ᆸ	erbium	89	[253]	F	fermium	100
165	유	holmium	29	[254]	E	einsteinium	66
163	D	dysprosium	99	[251]	ຽ	californium	98
159	ΤP	terbium	65	[245]	æ	berkelium	97
157	PS	gadolinium	64	[247]	Ę	anium	96
152	Eu	europium	63	[243]	Am	americium	95
150	Sm	samarium	62	[242]	Pu	plutonium	94
[147]	Pm	promethium	61	[237]	å	neptunium	93
144	PN	neodymium	9	238	_	uranium	92
141	Pr	praseodymium	29	[231]	Pa	protactinium	91
140	Ç	cerium	28	232	£	thorium	06
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