Please check the examination details below	v before entering your candidate information
Candidate surname	Other names
Pearson Edexcel International Advanced Level	Candidate Number
Monday 18 Janu	uary 2021
Morning (Time: 1 hour 45 minutes)	Paper Reference WCH15/01
Chemistry	
International Advanced Lev Unit 5: Transition Metals and	vel d Organic Nitrogen Chemistry
You must have: Scientific calculator, Data booklet	Total Marks

Instructions

- Use **black** ink or 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.
- Show all your working in calculations and include units where appropriate.

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.
- In the question marked with an **asterisk** (*), marks will be awarded for your ability to structure your answer logically showing how the points that you make are related or follow on from each other where appropriate.
- 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.

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 \boxtimes . If you change your mind, put a line through the box \boxtimes and then mark your new answer with a cross \boxtimes .

1 When an alkene is added to a solution of potassium manganate(VII), the purple solution turns colourless.

In terms of electron transfer and oxidation number, how does the manganese change in this reaction?

		Liection transfer	Oxidation number
X	A	gains electrons	increases
X	В	gains electrons	decreases
×	C	loses electrons	increases
X	D	loses electrons	decreases

(Total for Question 1 = 1 mark)

2 The standard hydrogen electrode uses an electrode of platinum coated in a finely divided form of the metal called platinum black.

What is the purpose of this coating?

- A to increase the rate of the equilibrium between the hydrogen gas and the hydrogen ions
- **B** to provide an inert protective coating for the electrode
- C to increase the electrical conductivity of the electrode
- **D** to ensure that the conditions remain standard

(Total for Question 2 = 1 mark)



3 An electrochemical cell is set up to measure $E_{\text{cell}}^{\ominus}$ for the reaction

$$Fe(s) + Sn^{2+}(aq) \rightarrow Fe^{2+}(aq) + Sn(s)$$

(a) What is the cell diagram for this cell?

(1)

- **A** Fe(s) | Fe²⁺(aq) || Sn(s) | Sn²⁺(aq)
- **B** $Fe^{2+}(aq) | Fe(s) | | Sn^{2+}(aq) | Sn(s)$
- \square **C** Fe(s) | Fe²⁺(aq) || Sn²⁺(aq) | Sn(s)
- \square **D** Fe²⁺(aq) | Fe(s) || Sn(s) | Sn²⁺(aq)
- (b) The standard electrode potential for the Fe / Fe²⁺ electrode system is -0.44 V and $E_{\text{cell}}^{\ominus}$ for the reaction is +0.30 V.

What is the standard electrode potential for the Sn / Sn²⁺ electrode system?

(1)

- **■ B** -0.14 V
- C +0.14 V
- \square **D** +0.74 V

(Total for Question 3 = 2 marks)

Use this space for any rough working. Anything you write in this space will gain no credit.

4 Hydrogen-oxygen fuel cells can operate in acidic or alkaline conditions.

What is the reaction at the anode in an alkaline hydrogen-oxygen fuel cell?

- \square **A** $O_2(g) + 2H_2O(I) + 4e^- \rightarrow 4OH^-(aq)$
- **B** $4OH^{-}(aq) \rightarrow O_2(g) + 2H_2O(I) + 4e^{-}$
- \square **C** $2H_2O(I) + 2e^- \rightarrow H_2(g) + 2OH^-(aq)$
- \square **D** $H_2(g) + 2OH^-(aq) \rightarrow 2H_2O(1) + 2e^-$

(Total for Question 4 = 1 mark)

- **5** Which of these has the greatest number of unpaired electrons in each of its atoms?
 - A chromium
 - **B** iron

 - **D** vanadium

(Total for Question 5 = 1 mark)

- 6 Nickel is classified as a transition metal. This is because nickel
 - A is a d block element
 - **B** has partially filled d orbitals
 - C forms stable ions with partially filled d orbitals
 - **D** forms stable compounds in which it has different oxidation states

(Total for Question 6 = 1 mark)

Use this space for any rough working. Anything you write in this space will gain no credit.

ω, .	****	c u.	e the shapes of these complexes?	(1)
	<	A	both complexes are square planar	
	<	В	both complexes are tetrahedral	
	<	C	Pt(NH ₃) ₂ Cl ₂ is tetrahedral and CrCl ₄ is square planar	
	<	D	Pt(NH ₃) ₂ Cl ₂ is square planar and CrCl ₄ is tetrahedral	
			the bonding between the ligands and the central atom in omplexes?	
				(1)
2	K	Α	the bonding in both complexes is ionic	
	<	В	the bonding in both complexes is dative covalent	
2	K	C	the bonding in Pt(NH ₃) ₂ Cl ₂ is dative covalent and in CrCl ₄ is ionic	
	K	D	the bonding in Pt(NH ₃) ₂ Cl ₂ is ionic and in CrCl ₄ is dative covalent	
			(Total for Question 7 = 2	marks)
Coba	alt c	hlo	ride is used as a test for the presence of water.	
Γhis	test	de	pends on the fact that	
X	Α	ar	hydrous cobalt(II) chloride is blue and hydrated cobalt(II) chloride is pi	nk
X	В	ar	hydrous cobalt(II) chloride is pink and hydrated cobalt(II) chloride is bl	ue
X	C	cc	obalt(II) chloride is blue and cobalt(III) chloride is pink	
X	D	cc	obalt(II) chloride is pink and cobalt(III) chloride is blue	
			(Total for Question 8 = 1	l mark)
his	spa	ace	for any rough working. Anything you write in this space will gain n	



9 Iodide ions are oxidised by peroxodisulfate ions in aqueous solution.

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

This reaction is catalysed by adding Fe²⁺ ions to the solution.

This catalysis is effective because

- \blacksquare **A** Fe²⁺ reacts with iodide ions and with peroxodisulfate ions
- **B** Fe²⁺ has many electrons in its outermost subshells
- C Fe²⁺ has many active sites on which the reaction can occur
- \square **D** Fe²⁺ is readily oxidised to Fe³⁺ which is then reduced to Fe²⁺

(Total for Question 9 = 1 mark)

- 10 The delocalised electrons in benzene result from the overlap of
 - \square **A** s orbitals to form σ bonds
 - \square **B** s orbitals to form π bonds
 - \square **C** p orbitals to form σ bonds
 - \square **D** p orbitals to form π bonds

(Total for Question 10 = 1 mark)

11 The reaction of ethene with bromine occurs under normal laboratory conditions but the reaction of benzene with bromine to form bromobenzene requires heat and the presence of a catalyst.

The best explanation for the difference in reactivity is that the delocalised electrons in benzene

- A repel electrophiles
- **B** result in a kinetic barrier to intermediate formation
- C result in benzene having an endothermic enthalpy of formation
- D make benzene thermodynamically stable with respect to the formation of bromobenzene

(Total for Question 11 = 1 mark)

12 What is the name of the compound shown?

- A 1-methylpropanamide
- **B** 3-methylpropanamide
- **D** 3-methylbutanamide

(Total for Question 12 = 1 mark)

13 Separate 0.1 mol dm⁻³ aqueous solutions of ammonia, butylamine and phenylamine were prepared.

Which of the following sequences shows the solutions in order of **increasing** pH?

- A butylamine, phenylamine, ammonia
- B ammonia, butylamine, phenylamine
- C phenylamine, ammonia, butylamine
- **D** ammonia, phenylamine, butylamine

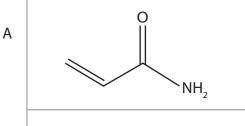
(Total for Question 13 = 1 mark)

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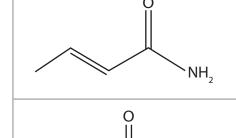
14 The repeat unit of a polymer is shown.

What is the structure of the monomer?

 \times

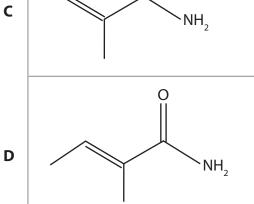


⊠ B



X

X



(Total for Question 14 = 1 mark)

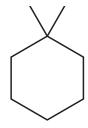
15 The repeat unit of a polymer is shown.

This polymer could be

- A both a polypeptide and a polyamide
- **B** neither a polypeptide nor a polyamide
- C a polypeptide but not a polyamide
- **D** a polyamide but not a polypeptide

(Total for Question 15 = 1 mark)

16 The structure of a hydrocarbon is shown.



How many peaks will there be in the ${}^{13}\text{C}$ NMR spectrum of this compound?

- A four
- **B** five
- C seven
- **D** eight

(Total for Question 16 = 1 mark)

Use this space for any rough working. Anything you write in this space will gain no credit.

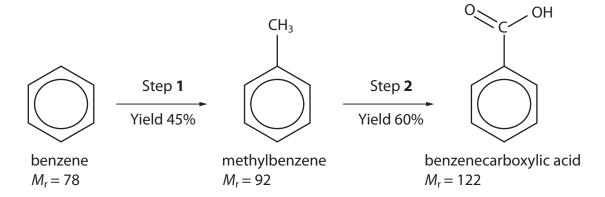
17 When a sample of a hydrocarbon is burned completely in oxygen, 2.64 g of carbon dioxide and 0.81 g of water are formed.

Which of these could be the **molecular** formula of the hydrocarbon?

- \square A C_2H_3
- \square **B** C_4H_3
- \square **C** C_4H_6
- \square **D** $C_{12}H_9$

(Total for Question 17 = 1 mark)

18 Benzenecarboxylic acid may be produced from benzene in a two-step synthesis.



8.24 g of benzenecarboxylic acid was formed in this synthesis.

What mass of benzene was used?

- B 5.27 g

(Total for Question 18 = 1 mark)

TOTAL FOR SECTION A = 20 MARKS

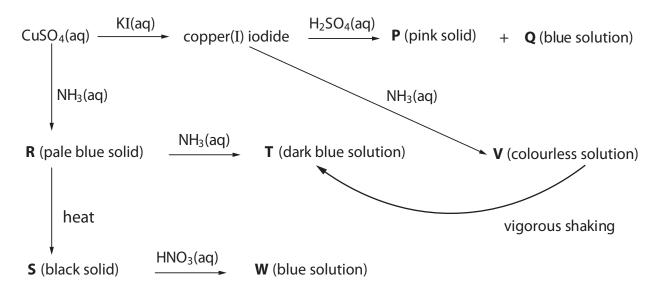
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SECTION B

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

19 The diagram summarises some reactions of copper compounds.



(a) Identify, by name (including the oxidation state) or formula, the species in the sequence that contain copper.

(7)

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- (b) **T** and **V** are the same type of chemical species.
 - (i) Name this type of chemical species.

(1)



A detailed explanation of the fact that T is coloured is not required.	(3)
(iii) Suggest an explanation for the change of $oldsymbol{V}$ into $oldsymbol{T}$ on shaking.	(2)
The reaction between copper(I) iodide and sulfuric acid is a disproportionation.(i) Write the ionic equation for this disproportionation reaction.	
State symbols are not required.	(1)
(ii) Show that the reaction in (c)(i) is thermodynamically feasible. Use the standard electrode potentials of the relevant half-cells from the Data Booklet.	
Hom the Data Booklet.	(2)



(d) The rare mineral mitscherlichite has the chemical formula K₂CuCl₄•nH₂O.

4.26 g of mitscherlichite was dissolved in distilled water and the solution made up to 250.0 cm³. Excess potassium iodide solution was added to a 25.0 cm³ portion of this solution and the iodine formed was titrated against a solution of sodium thiosulfate with a concentration of 0.0500 mol dm⁻³.

This procedure was repeated until concordant results were obtained. The mean accurate titre was 26.65 cm³.

The equations for the reactions are

$$2Cu^{2+} + 4I^- \rightarrow 2CuI + I_2$$

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

Calculate the value of n, the number of moles of water of crystallisation per mole of mitscherlichite.

(6)

(Total for Question 19 = 22 marks)

20 Ketones are useful starting compounds in organic synthesis.

This question is about butanone.

- (a) The mass spectrum of butanone has significant peaks at m/z = 43 and at m/z = 57.
 - (i) Give the structures of the species responsible for these two peaks.

(2)

(ii) Give the structure of **one** other species that you would expect to produce a peak at a different m/z value in the mass spectrum of butanone.

(1)

(b)	Devise a reaction scheme to prepare propan-1-ol from butanone, using no more than four steps.	
	Identify the reagents and essential conditions for each step and give the name or structure of each of the intermediate compounds.	
		(4)

(c)	Devise a reaction scheme to prepare 2-methylbut-2-ene from butanone, using no more than four steps.	
	Identify the reagents and essential conditions for each step and give the name or structure of each of the intermediate compounds.	(5)
	(Total for Question 20 = 12 ma	rks)



21 A yellow crystalline solid **E** dissolved in distilled water to give a yellow solution. Addition of dilute sulfuric acid to this solution produced an orange solution **F**. Warming **F** with ethanol resulted in a green solution **G**, and the formation of ethanal.

A standard cell was set up using solutions of **F** and **G** for the right-hand electrode and ethanol and ethanal for the left-hand electrode.

$$E_{\text{cell}}^{\ominus}$$
 was found to be +1.94 V.

$$CH_3CHO(aq) + 2H^+(aq) + 2e^- \rightleftharpoons C_2H_5OH(aq)$$
 $E^{\ominus} = -0.61 \text{ V}$

(a) Deduce the formulae of the ions responsible for the colours of **F** and **G**, using the standard electrode potential and E^{\ominus} given, and the values in the Data Booklet.

(2)

(b) Write the overall equation for the reaction in the cell. State symbols are not required.

(2)

(c) Write the ionic equation for the reaction of the aqueous solution of **E** with dilute sulfuric acid. State symbols are not required.

(1)

(Total for Question 21 = 5 marks)



22 Using excess oxygen, 25 cm³ of a gaseous hydrocarbon C_xH_v was burned completely.

After cooling to room temperature the total gas volume was measured and found to be 75 cm³ less than the total gas volume before the mixture was ignited.

When the product gases were shaken with potassium hydroxide solution, the total gas volume decreased by a further 100 cm³.

All gas volumes were measured at room temperature and pressure.

A general equation for the combustion of a hydrocarbon is

$$C_xH_y + \left[x + \frac{y}{4}\right]O_2 \rightarrow xCO_2 + \left[\frac{y}{2}\right]H_2O$$

(a) Determine the molecular formula of C_xH_y. You **must** show your working.

(3)

(b) When C_xH_y was added to a little bromine water and the mixture shaken, the bromine water remained yellow.

Suggest **two** possible structures for C_xH_y .

(2)

(Total for Question 22 = 5 marks)



23	Compare and contrast the mechanism of the action of platinum as a catalyst in the removal of pollutants from car engine exhaust fumes with that of vanadium(V) oxide as a catalyst in the Contact Process for the manufacture of sulfuric acid.	
	General definitions of catalysts are not required.	
		(6)

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	TOTAL FOR SECTION B = 50 MARKS
	TOTAL FOR SECTION B = 50 MARKS



SECTION C

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

Coffee Chemistry

24 There are over a thousand chemical compounds in coffee and their physiological effects are the subject of considerable speculation and research. The verdict on coffee is contradictory: some of the compounds have been identified as toxic and even carcinogenic but others are antioxidants associated with cancer prevention. Recent research has identified compounds in coffee that might be used in the treatment of prostate cancer.

By far the best known compound in coffee is caffeine, the most widely consumed psychoactive drug in the world. In small amounts it is a stimulant but doses in excess of 10 g per day are toxic. Caffeine contains amide and amine groups.

Chlorogenic acid is responsible for the acidic taste of coffee. It is an antioxidant and has also been shown to slightly decrease blood pressure.

Caffeic acid, quinic acid and acetoin are also present in coffee.

(a) Another way of drawing the structure of caffeine is shown.

(i) The bonding represented by this diagram of caffeine differs from that given in the passage.

Explain what this diagram indicates about the bonding in caffeine, stating the effect on the structure of caffeine.

(ii) Su etl	ggest why caffeine is a much weaker nylamine, even though the right-hanc	base than a primary amine suc I ring has two amine groups.	h as (2)
(ii) Su etl	ggest why caffeine is a much weaker nylamine, even though the right-hand	base than a primary amine suc I ring has two amine groups.	
(ii) Su	ggest why caffeine is a much weaker nylamine, even though the right-hand	base than a primary amine suc	

(3)

- (b) A 200 cm³ cup of coffee contains approximately 85 mg of caffeine.
 - (i) Calculate the concentration, in mol dm⁻³, of caffeine in this cup of coffee. Give your answer to an appropriate number of significant figures.

(4)

(ii) The removal of caffeine from the body is a first order reaction with a half-life of between three and seven hours for an adult.

An adult drinks coffee containing a total of 160 mg of caffeine.

Calculate to the nearest hour the **minimum** time needed for the amount of caffeine in their body to drop to 20 mg.

(2)

(c) Chlorogenic acid is an ester of caffeic acid, a compound that is present in all plants.

(i) A student suggested that caffeic acid could be synthesised by an electrophilic substitution of 1,2-dihydroxybenzene.

Draw the mechanism of this electrophilic substitution, including the formation of a suitable electrophile.

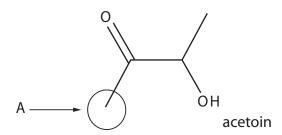
(5)

(ii) Deduce the structure of quinic acid which combines with caffeic acid to form chlorogenic acid.

(1)

quinic acid

(d) The structure of acetoin is shown with one of the proton environments labelled.



(i) Identify the other proton environments of acetoin on the structure and label them B, C etc.

(1)

(ii) Complete the table to show the splitting pattern in the high resolution proton NMR spectrum of acetoin.

(2)

Proton environment	Splitting pattern
А	

(Total for Question 24 = 20 marks)

TOTAL FOR SECTION C = 20 MARKS
TOTAL FOR PAPER = 90 MARKS

ytterbium lutetium

175

Ξ

Tm thulium

167 Er erbium 68

Holmium F

165

lawrencium

nobelium

mendelevium

fermium

californium einsteinium

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orted	Rn radon 86	[222]	Xenon xenon 54	131.3	Kr krypton 36	0 0	Ar argon	10	Ne se	20.2	4.0 He hettum 2	0 (8)	
peen repo	At astatine 85	[210]	I iodine 53	126.9	Br bromine 35	-	35.5 CL chlorine	6	- Linerine	19.0	(77)	1	
116 have l	Po polonium 84	[509]	Te tellurium 52	127.6	Se selenium 34	0	32.1 Sulfur	8	0	16.0	(16)	9	
Elements with atomic numbers 112-116 have been reported but not fully authenticated	Bi bismuth 83	209.0	Sb antimony 51	121.8	As arsenic 33	C1	31.0 P	7	Z	14.0	(15)	'n	
atomic nul but not f	Pb lead 82	207.2	S # 05	118.7	Ge germanium 32	4	28.1 Si	6	٥	12.0	(14)	4	
ents with	T1 thallium 81	204.4	In indium	114.8	Ga gallium 31	2	27.0 Al	5	a	10.8	(13)	m	
Elem	Hg mercury 80	200.6	Cd cadmium 48	112.4	Zn zinc 30	(77)							ents
[272] Rg roentgenium	Au gold 79	197.0	Ag silver 47	107.9	Cu copper 29	(11)							The Periodic Table of Elements
[268] [271] [272] Mt Ds Rg meitnerium damstadtium roentgenium	Pt platinum 78	195.1	Pd palladium 46	106.4	N icket 28	(0.)							le of
[268] Mt neitnerium	Ir iridium 77	192.2	Rh rhodium 45	102.9	Co cobalt 27	(2)							lab
HS hassium r	Os osmium 76	190.2	Ru ruthenium 44	101.1	Fe iron 26	6					1.0 Hydrogen		DIBOL
[264] Bh	Re rhenium 75	186.2	Tc technetium 43	1861	Mn nanganese 25	()							e Pe
[266] Sg	W tungsten 74	183.8	Mo Tc molybdenum technetium 42 43	95.9	Cr Mn chromium manganese 24 25	6		ımper	lo	nass			
[262] Db dubnium	Ta tantalum 73	180.9	níobium r	92.9	V vanadium 23	(6)		atomic (proton) number	atomic symbol	relative atomic mass	Key		
[261] Rf	Hf hafnium 72	178.5	Zr zirconium 40	91.2	Ti titanium 22	(2)		atomic	ator	relativ			
[227] Ac* actinium	La* lanthanum 57	138.9	¥ yttrium 39	88.9	Sc scandfum 21	(2)							
[226] Ra radium	Ba barium L 56	137.3	Sr strontium 38	87.6	Ca calcium 20	71	Ag	4	Be	9.0	(2)	7	
[223] Fr franctum	Cs caesium 55	132.9	Rb rubidium 37	85.5	K potassium 19	Ε,	23.0 Na		5	6.9	(1)	÷.	

* Lanthanide series

140