Surname	Othe	er names
Pearson Edexcel GCE	Centre Number	Candidate Number
Chemistr	'y	
	ples of Chemistry ganic Nitrogen Ch optic assessment)	emistry
Unit 5: General Princip	ganic Nitrogen Ćh optic assessment)	Paper Reference
Unit 5: General Princip Metals and Org (including syn	ganic Nitrogen Ćh optic assessment) 6 – Morning	emistry

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.

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.
- Keep an eye on the time.
- Try to answer every question.
- Check your answers if you have time at the end.

P 4 6 6 6 1 A 0 1 3 2

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 of the following gives the oxidation states of manganese in the ions shown?

		MnO ₄ ²⁻	MnO ₃
×	A	+7	+6
X	В	+6	+5
×	c	+7	+5
X	D	+6	+6

(Total for Question 1 = 1 mark)

2 Which of the following gives the electrodes and electrolyte that are used in an alkaline hydrogen fuel cell?

Electrodes		Electrodes	Electrolyte	
×	A	graphite	potassium hydroxide solution	
×	В	graphite	water with a little salt	
X	С	platinum	potassium hydroxide solution	
X	D	platinum	water with a little salt	

(Total for Question 2 = 1 mark)

- **3** Which of the following **cannot** be used to detect alcohol in a breathalyser test?
 - ☑ A Fractional distillation
 - ☑ B Fuel cell
 - ☑ C Infrared spectroscopy
 - ☑ D Reduction of dichromate(VI) ions

(Total for Question 3 = 1 mark)

- **4** A titration using potassium manganate(VII) in dilute sulfuric acid can be used to determine the percentage of
 - ☑ A aspirin in aspirin tablets.
 - **B** chlorine in bleach.
 - **C** copper in an alloy.
 - ☑ D iron(II) sulfate in iron tablets.

(Total for Question 4 = 1 mark)

Which of the following gives the electronic configurations for a chromium atom and a chromium(II) ion?

Cr		Cr	Cr ²⁺	
X	A	[Ar]3d ⁴ 4s ²	[Ar]3d ⁴	
X	В	[Ar]3d ⁵ 4s ¹	[Ar]3d⁴	
X	c	[Ar]3d ⁴ 4s ²	[Ar]3d²4s²	
X	D	[Ar]3d ⁵ 4s ¹	[Ar]3d³4s¹	

(Total for Question 5 = 1 mark)

6 Aqueous sodium hydroxide and aqueous ammonia are added to separate solutions of the same metal ion. The observations are shown in the table below.

Reagent added	A few drops	Excess
NaOH(aq)	green precipitate	green precipitate remains
NH₃(aq)	green precipitate	green precipitate dissolves to form a blue solution

The metal ion is

- \triangle A $Cr^{3+}(aq)$.
- \square **B** Fe²⁺(aq).
- \square **C** Fe³⁺(aq).
- \square **D** Ni²⁺(aq).

(Total for Question 6 = 1 mark)

7 The reaction between cerium(IV) ions and thallium(I) ions is very slow.

$$2Ce^{4+}(aq) + Tl^{+}(aq) \rightarrow 2Ce^{3+}(aq) + Tl^{3+}(aq)$$

Which of these ions could catalyse this reaction?

- A Al³+
- B Fe³⁺
- C Na⁺

(Total for Question 7 = 1 mark)

- **8** Which of these hydroxides is amphoteric?
 - \triangle A Cu(OH)₂
 - \square **B** Mg(OH)₂
 - \square C Ni(OH)₂
 - \square **D** $Zn(OH)_2$

(Total for Question 8 = 1 mark)

9	During a titration between acidified manganate(VII) ions and sulfate(IV) ions, the
	manganate(VII) ions are reduced to manganese(II) ions and the sulfate(IV) ions are
	oxidized to sulfate(VI) ions.

The mole ratio of manganate(VII) ions to sulfate(IV) ions in this reaction is

- **■ B** 7:4
- **∠ C** 2:5
- **D** 4:7

(Total for Question 9 = 1 mark)

- **10** The total number of compounds with the structural formula C₆H₃CH₃(NO₂)₂, which contain a benzene ring, is
 - **A** four.
 - **B** five.
 - C six.
 - **D** seven.

(Total for Question 10 = 1 mark)

11 This question is about the ester shown below.

(a) The number of peaks seen in the **low** resolution proton nmr spectrum of this ester is

(1)

- A two.
- **B** three.
- C four.
- **D** five.
- (b) The peak in the **high** resolution proton nmr spectrum corresponding to the proton in **bold** on the structure above will

(1)

- A not be split.
- **B** be split into three peaks.
- □ C be split into four peaks.
- **D** be split into seven peaks.

(Total for Question 11 = 2 marks)

12 Safranal is one of the substances that contributes to the aroma of saffron.



Separate samples of safranal were tested with bromine water, 2,4-dinitrophenylhydrazine and Fehling's solution.

What are the final observations when safranal is tested with each of those reagents?

		Bromine water	2,4-dinitrophenylhydrazine	Fehling's solution
×	A	orange solution	orange solution	red precipitate
×	В	colourless solution	orange precipitate	red precipitate
×	c	orange solution	orange solution	blue solution
×	D	colourless solution	orange precipitate	blue solution

(Total for Question 12 = 1 mark)

13 The structure of the organic product of the reaction between phenol and excess bromine water is

⊠ A



В

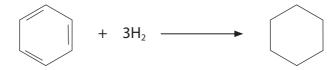
⊠ C

ОН

⊠ D

(Total for Question 13 = 1 mark)

14 If it is assumed that the structure of benzene has three localised double bonds (structure **X**), the calculated standard enthalpy change of hydrogenation is –360 kJ mol⁻¹.



structure X

The actual standard enthalpy change of hydrogenation of benzene is -208 kJ mol⁻¹.

From these data, it can be deduced that the

- A actual benzene structure is kinetically more stable than structure **X** as it requires a high activation energy to react.
- B actual benzene structure is thermodynamically more stable than structure X as it has a lower enthalpy content.
- structure **X** is kinetically unstable as it undergoes addition reactions at room temperature.
- structure **X** is thermodynamically more stable than the actual benzene structure as the standard enthalpy change of hydrogenation is more exothermic.

(Total for Question 14 = 1 mark)



15 The repeat unit for poly(propenamide) is

(Total for Question 15 = 1 mark)

16 The structures of three amino acids are shown in the table.

Amino acid	Structure
cysteine	HSCH ₂ CH(NH ₂)COOH
glycine	H ₂ NCH ₂ COOH
threonine	CH₃CH(OH)CH(NH₂)COOH

The tripeptide glycine-cysteine-threonine is

- A H₂NCH₂CONHCH(CH(OH)CH₃)CONHCH(CH₂SH)COOH
- **■ B** H₂NCH₂CONHCH(CH₂SH)CONHCH(CH(OH)CH₃)COOH
- C H₂NCH(CH(OH)CH₃)CONHCH(CH₃SH)CONHCH₂COOH
- D H₂NCH(CH₂SH)CONHCH₂ CONHCH(CH(OH)CH₃)COOH

(Total for Question 16 = 1 mark)

17 The amino acid alanine, H₂NCH(CH₃)COOH, exists as a solid at room temperature.

The most important reason for this is that it

- **A** exists as a zwitterion.
- **B** forms hydrogen bonds.
- **C** is amphoteric.
- **D** has strong London forces.

(Total for Question 17 = 1 mark)

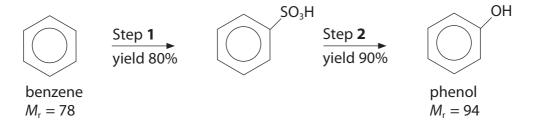
18 Complete combustion of a hydrocarbon produced 0.66 g of carbon dioxide and 0.225 g of water.

Which of the following molecular formulae is consistent with these data?

- \square A C_3H_6 .
- \square **B** C₃H₈.
- \square **C** C₆H₆.
- \square **D** C₆H₁₀.

(Total for Question 18 = 1 mark)

19 Phenol can be produced from benzene as shown in the reaction sequence below.



The mass of phenol, to 2 decimal places, produced from 3.90 g of benzene is

- **■ B** 3.76 g.
- ☑ D 4.70 g.

(Total for Question 19 = 1 mark)

TOTAL FOR SECTION A = 20 MARKS

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SECTION B

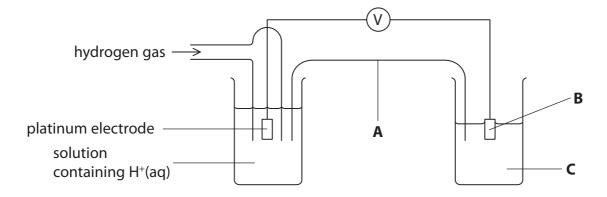
Answer ALL the questions. Write your answers in the spaces provided.

- **20** Vanadium exists in different oxidation states which can be interconverted using suitable oxidising and reducing agents.
 - (a) Use relevant standard electrode potential values, on page 14 of the Data Booklet, to complete the table below in which two E^{\ominus} values are missing.

(1)

Half-equation	E [⊕] /V
$V^{2+}(aq) + 2e^- \rightleftharpoons V(s)$	
$V^{3+}(aq) + e^- \rightleftharpoons V^{2+}(aq)$	
$VO^{2+}(aq) + 2H^{+}(aq) + e^{-} \implies V^{3+}(aq) + H_2O(I)$	+0.34
$I_2(aq) + 2e^- \rightleftharpoons 2I^-(aq)$	+0.54
$VO_2^+(aq) + 2H^+(aq) + e^- \implies VO^{2+}(aq) + H_2O(I)$	+1.00

(b) The standard electrode potential of $V^{3+}(aq) + e^- \rightleftharpoons V^{2+}(aq)$ is measured using the apparatus below.



(i) Identify, by name or formula, the substances needed in the salt bridge and the right-hand half-cell to measure the standard electrode potential.

(3)

- A Salt bridge containing a solution of
- **B** Electrode made of
- **C** Solution containing
 - (ii) State the **three** standard conditions needed for this measurement.

(2)

1

2.....

3

*(c)	A solution containing iodide ions, I^- , was added to an acidified solution containing vanadium(V) ions, VO_2^+ .	
	Predict the oxidation state of the vanadium ions left at the end of the reaction. Justify your prediction by calculating the $E_{\text{cell}}^{\ominus}$ for any relevant reaction(s).	
	Write the ionic equation for any reaction(s) occurring. State symbols are not required.	
	(5)	
	(Total for Question 20 = 11 marks)	
	(10 tal. 101 Quantum 20 11 mana)	

21 (a) The structures of 2-aminopropanoic acid and 3-aminopropanoic acid are shown.

2-aminopropanoic acid

3-aminopropanoic acid

(i) Explain how the **low** resolution proton nmr spectra of these two amino acids differ.

(2)

(ii) Explain whether or not 3-aminopropanoic acid is chiral.

(1)

(iii) Write ionic equations for the reaction of 3-aminopropanoic acid with

(2)

H⁺ ions

OH-ions



(iv) Draw two repeat units of the polymer formed when 3-aminopropanoic acid polymerizes.

(1)

(b) The food colouring E110 is also known as Sunset Yellow.

It can be synthesised as shown below.

(i) Give the reagents and condition for **Step 1**.

(2)



(ii) Draw the structure of the reagent needed for Step 2 .	(1)
(iii) Explain why Sunset Yellow can exist as geometric isomers.	(1)
*(iv) Describe the essential steps of the method that you would use to prepare a pure, dry sample of the solid Sunset Yellow from an impure sample of the food colouring. You may assume that ethanol is a suitable solvent for this method.	(4)



(v) Suggest how you could check that a sample of Sunset Yellow is pure.

(1)

(c) Explain how a chemist could use phenylmethanol to synthesise a sample of benzamide in three steps.

C NH₂

phenylmethanol

benzamide

Include the reagents for the steps in the synthesis and draw the structures of **all** the intermediates.

(5)

(Total for Question 21 = 20 marks)

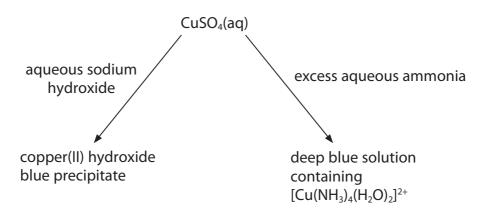
co	opper and zinc are both in the d-block of the Periodic Table. Copper forms ompounds that contain Cu ⁺ and Cu ²⁺ ions but zinc only forms compounds that ontain Zn ²⁺ ions. Complete the electronic configurations of the Cu ²⁺ ions and Zn ²⁺ ions and hence explain why copper is classified as a transition metal but zinc is not.	(2)
Cu ²⁺	[Ar]	
Zn ²⁺	[Ar]	
(b) Some photochromic glasses contain silver(I) and copper(I) chlorides.	
(.2	Explain, with the aid of an equation, why these photochromic glasses go darker in sunlight.	
	sumgnt.	(2)

(c)	Copper forms a complex ion with the formula $[CuCl_4]^{2-}$. This has the same shape as $[Pt(NH_3)_2Cl_2]$.	
	Draw the shape of the $[CuCl_4]^{2-}$ ion and state the type of bonding between the ligands and the metal ion.	(2)
	Shape	(2)
	Bonding	
(d) The [CuCl ₂] ⁻ ion is formed by boiling a solution of copper(II) chloride with copper	
	turnings and concentrated hydrochloric acid.	
	(i) Write an equation for this reaction. State symbols are not required.	(1)
	(ii) State the meaning of the term disproportionation and explain whether or not this reaction to form the $[CuCl_2]^-$ ion is a disproportionation reaction.	
		(2)

(iii) Explain why the $[CuCl_2]^-$ ions are colourless.

(2)

(e) Copper(II) sulfate solution reacts with aqueous sodium hydroxide and with aqueous ammonia.



(i) Write the **ionic** equation for the reaction of copper(II) sulfate solution with aqueous sodium hydroxide. Include state symbols.

(1)

(ii) State the type of reaction occurring overall when excess aqueous ammonia is added to copper(II) sulfate solution.

(1)

(f)	1,2-diaminoethane is a bidentate ligand. It reacts with copper(II) ions in aqueous solution.	
	$[Cu(H_2O)_6]^{2+} + 3H_2NCH_2CH_2NH_2 \implies [Cu(H_2NCH_2CH_2NH_2)_3]^{2+} + 6H_2O$	
	(i) State what is meant by the term bidentate .	(1)
	(ii) Explain, in terms of entropy, why the reaction takes place.	(2)
	(Total for Question 22 = 16 ma	rks)

TOTAL FOR SECTION B = 47 MARKS

SECTION C

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

23

Analgesics

Analgesics are taken to relieve the symptoms of pain.

Paracetamol

This has been synthesised in a three-step process since the 1950s.

Ibuprofen

This was first synthesised in the 1960s from propanoic acid. That process involved six steps and produced more waste than the required drug.

It is now manufactured in a three-step process. Some data about these two analgesics are given in the table below.

Analgesic	Molecular formula	Molar mass / g mol ⁻¹
paracetamol	C ₈ H ₉ NO ₂	151
ibuprofen	C ₁₃ H ₁₈ O ₂	206

(a) **Name** the functional group in paracetamol, other than the phenol group.

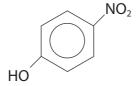
(1)

(b) Paracetamol is made from phenol in a three-step process.

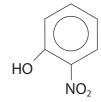


dilute H₂SO₄ NaNO₃

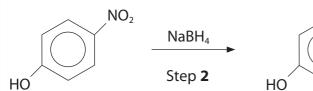
Step 1

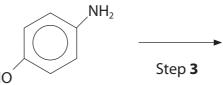


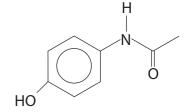
4-nitrophenol



2-nitrophenol







(i) In a typical nitration of an arene, the electrophile is formed as shown below.

HNO₃

H₂SO₄

 $H_2NO_3^+$

HSO₄

Equation 1

 $H_2NO_3^+ \rightarrow H_2O + NO_2^+$

Equation 2

Identify the acid-base conjugate pairs in **Equation 1**. Write your answers on the dotted lines under the equation.

(1)

(ii) Give a mechanism for the nitration of phenol by NO_2^+ to form 4-nitrophenol.	(3)
(iii) Explain why phenol is nitrated much more readily than benzene.	(2)
(iv) State the type of reaction taking place in Step 2 .	(1)
(v) Suggest a reagent for Step 3 .	(1)

(vi) 2-nitrophenol has a melting temperature of 46 $^{\circ}$ C and 4-nitrophenol has a melting temperature of 114 $^{\circ}$ C.

Suggest, in terms of intermolecular forces, why these two compounds have different melting temperatures.

(2)

(c) Paracetamol can be hydrolysed to form 4-aminophenol and ethanoic acid.

$$H_{0}$$
 $+ H_{2}O$ $+ CH_{3}COOH$

The amount of 4-aminophenol produced can be determined using a redox titration. The half-equation for the oxidation of 4-aminophenol is given below.

The oxidizing agent is ammonium cerium(IV) sulfate and ferroin indicator is used to detect the end-point of the titration. During the reaction, the Ce⁴⁺ ions are reduced to Ce³⁺ ions.

(i) Write the overall equation for the reaction between Ce⁴⁺ ions and 4-aminophenol.

(1)



(ii) In an experiment, 0.500 g of a tablet containing paracetamol was hydrolysed and the solution was made up to 100 cm³.

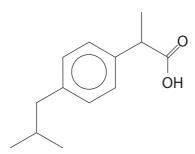
 $20.0~cm^3$ portions of the resulting solution were titrated with $0.100~mol~dm^{-3}$ ammonium cerium(IV) sulfate solution.

The mean titre was 12.60 cm³.

Calculate the percentage, by mass, of paracetamol in the tablet.

(5)

(d) (i) Identify the chiral carbon atom in ibuprofen with an asterisk (*).



(ii) Suggest a problem in the manufacture of a single isomer of a chiral drug and describe a way that the pharmaceutical industry might overcome this problem.

(2)

(1)

(e) Ibuprofen was originally made in a six-step process but is now made in a three-step process.

Suggest a specific environmental reason why the manufacturing process was changed.

(1)



(f) Ibuprofen is not very soluble in water. It can be made into an ionic, soluble salt by reacting it with lysine.

Draw the structures of **both** the cation and the anion in the soluble salt formed when ibuprofen reacts with lysine.

(2)

(Total for Question 23 = 23 marks)

TOTAL FOR SECTION C = 23 MARKS
TOTAL FOR PAPER = 90 MARKS



The Periodic Table of Elements

0 (8)	(18)	4.0	1	helium	(
7					1
9					
2					
4					1877
3					
		1.0	I	hydrogen	
	_				
					1/2:
2					

4.0 He helium	20.2 Ne	neon 10	39.9	Ar argon 18	83.8	고	krypton 36	131.3	Xe	xenon 54	[222]	Ru	radon 86				
(77)	19.0 Z	e.		CI chlorine a	8 6.67		bromine kr 35	126.9 1		iodine x	[210]	_	e.		reported		
		_					(2) (1) (1) (1) (1) (1) (1) (1) (1) (1) (1	127.6 12		200			Ē		nave been	pa	
(16)	16.0	oxygen 8	32.1	Sulfur 16	79.0	Se	selenium 34	127		tellurium 52	[209]	P	polonii 84		2-116 h	enticat	
(15)	14.0 N	nitrogen 7	31.0	P phosphorus 15	74.9	As	arsenic 33	121.8	Sb	antimony 51	209.0	Bi	bismuth 83		Elements with atomic numbers 112-116 have been reported but not fully authenticated		
(14)	12.0 C	carbon 6	28.1	Si silicon 14	72.6	Ge	germanium 32	118.7	Sn	tiu 20	207.2	Ъ	lead 82				
(13)	10.8 B	boron 5	27.0	AI aluminium 13	2.69	Ga	gallium 31	114.8	п	indium 49	204.4	F	thallium 81		ents with		
				(12)	65.4	Zu	zinc 30	112.4	В	cadmium 48	200.6	H	mercury 80		Elen		
				(11)	63.5	J	copper 29	107.9	Ag	silver 47	197.0	Αn	gold 79	[272]	Rg	roentgenium 111	
				(10)	58.7	ź	nickel 28	106.4	Pq	palladium 46	195.1	¥	platinum 78	[271]	Ds	darmstadtium 110	
				(6)	58.9	ပိ	cobalt 27	102.9	R	rhodium 45	192.2	ļ	iridium 77	[368]	Mt	meitnerium damstadtium 109 110	
1.0 Hydrogen				(8)	55.8	Fe	iron 26	101.1	Ru	ruthenium 44	190.2	os	osmium 76	[277]		hassium 108	
				0	54.9	Wn	manganese 25	[86]	7	technetium 43	186.2	Re	rhenium 75	[264]	Bh	bohrium 107	
A	mass bol	umber		(9)	52.0	င်	vanadium chromium manganese 23 24 25	95.9	Wo	molybdenum 42	183.8	>	tungsten 74	[366]	Sg	m seaborgium bo	
Key	relative atomic mass atomic symbol	name atomic (proton) number		(5)	50.9	>	vanadium 23	92.9		niobium 41	180.9	Ta	E		В	dubniu 105	
	relati ato	atomic		(4)	47.9	ï	titanium 22	91.2	Zr	zirconium 40	178.5	Ŧ	hafnium 72	[261]	Rf	rutherfordium 104	
				(3)	45.0	Sc	scandium 21	88.9	>	yttrium 39	138.9	La*	lanthanum 57	[227]		actinium 89	
(2)	9.0 Be	beryllium 4	24.3	Mg magnesium 12	40.1	Ca	calcium 20	87.6	Sr	strontium 38	137.3	Ba	barium 56	[326]	Ra	radium 88	
(1)	6.9 Li	lithium 3	702200	Na sodium 11	39.1	¥	potassium 19	85.5	ВЪ	rubidium 37	132.9	S	caesium 55	[223]	노	francium 87	

^{*} Lanthanide series

^{*} Actinide series

Ce Pr Nd Pm Sm Eu Gd Tb Dy Ho Er Tm Yb Lu cerium præcedymium necedymium promeethium samarium gadolinium terbium dysprosium holmium erbium thutlium ytterbium thutlium t		
Nd Pm Sm Eu Gd Tb Dy Ho Er Tm	175 Lu lutetium 71	[257] Lr lawrencium 103
Nd Pm Sm Eu Gd Tb Dy Ho Er	Yb ytterbium 70	No nobelium 102
Nd Pm Sm Eu Gd Tb Dy Ho	169 Tm thulium 69	[256] Md mendelevium 101
Nd Pm Sm Eu Gd Tb Dy Nocodymium promethium samarium europium gadolinium terbium dysprosium d	167 Er erbium 68	[253] Fm fermium 100
144 [147] 150 152 157 159 159 150 15	Ho hotmium 67	Es einsteinium 99
Nd Pm Sm Eu Gd	Dy dysprosium 66	Cf Cf californium 98
144 [147] 150 152 152 Nd Pm Sm Eu Eu Eu Eu Eu Eu Eu E	Tb terbium 65	Bk berkelium 97
144 [147] 150 15	Gd gadolinium 64	(247) Cm curium 96
Nd Pm Nd Nd Nd Nd Nd Nd Nd N	152 Eu europium 63	[243] Am americium 95
Nd n neodymium pro 60 238 U uranium ne	Sm samarium 62	Pu Pu plutonium 94
- n	Pm promethium 61	Np neptunium 93
Ce Pr cerium praseodymium 58 59 232 [231] Th Pa thorium protactinium 90 91	Nd neodymium 60	238 U uranium 92
Ce cerium 58 232 Th thorium 90	Pr praseodymium 59	Pa protactinium 91
	Ce cerium	232 Th thorium 90