Write your name here Surname	Other I	names
Pearson Edexcel International Advanced Level	Centre Number	Candidate Number
Chemistry Advanced Unit 5: General Principles and Organic Nitro (including synoptic	s of Chemistry II – 1 gen Chemistry	Fransition Metals
Tuesday 21 January 2014 –	Afternoon	Paper Reference WCH05/01
Time: 1 hour 40 minutes		WC1103/01
You must have: Data Booklet Candidates may use a calculate	tor.	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.

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 2 9 9 3 A 0 1 2 8

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 In which of the following compounds does iron have the highest oxidation number?
 - A Fe₃O₄
 - B K₂FeO₄
 - ☑ C Na₄Fe(CN)₅
 - ☑ D Na₃Fe(CN)₆

(Total for Question 1 = 1 mark)

- **2** Which of the following is **not** a redox reaction?
 - \square A $3CrCl_2 + Na_2CrO_4 + 8HCl \rightarrow 4CrCl_3 + 4H_2O + 2NaCl$
 - \square **B** $2MnO_4^{2-} + C_8H_7O_3^{-} + 2OH^{-} \rightarrow 2MnO_4^{3-} + C_8H_5O_3^{-} + 2H_2O$
 - \square **C** $3MnO_4^{2-} + 4H^+ \rightarrow 2MnO_4^{-} + MnO_2 + 2H_2O$

(Total for Question 2 = 1 mark)

3 The standard reduction potentials of two systems are given below.

$$Cu^{2+}(aq) + 2e^{-} \rightleftharpoons Cu(s)$$
 $E^{\ominus} = +0.34 \text{ V}$

$$Ag^{+}(aq) + e^{-} \rightleftharpoons Ag(s)$$
 $E^{\oplus} = +0.80 \text{ V}$

What is E_{coll}^{\oplus} for the reaction between copper and silver nitrate?

$$Cu(s) + 2Ag^{+}(aq) \rightarrow Cu^{2+}(aq) + 2Ag(s)$$

- **B** −0.46 V
- C +0.46 V

(Total for Question 3 = 1 mark)

4	zero. (cell is set up with two metal-metal ion half cells and the digital voltmeter reads ero. Given that all the components of the cell have been included and are working roperly, what is the most likely explanation for the zero reading?		
	☑ A The cell has been set up the wrong way round.			
	$ □ $ B The entropy change, $ΔS_{\text{system}} = 0$.			
	 The activation energy for the reaction is very high. 			is very high.
	⊠ D	The rea	action system is at equilibrium.	
				(Total for Question 4 = 1 mark)
5	What is the electronic configuration of the Fe ³⁺ ion?			2 ³⁺ ion?
	⋈ A	[Ar]	3d	4s
	⊠ B	[Ar]	$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	
	⊠ C	[Ar]	$\begin{array}{ c c c c c c c c c c c c c c c c c c c$	1
	⊠ D	[Ar]	<u> </u>	$\uparrow\downarrow$
				(Total for Question 5 = 1 mark)
6	Ammonia gas is formed in the combustion of biomass fuels and is a significant pollutant when it is released directly into the atmosphere. One method of removing this ammonia involves its oxidation using a titanium(IV) oxide catalyst.			
	The b e	est expla	anation for the use of titanium((IV) oxide is that the titanium
	⊠ A	contain	ns many active sites on which t	he reaction can occur.
	⊠ B		lily oxidized to a higher oxidation state +4.	on state which can then be reduced back
	⊠ C		lily reduced to a lower oxidatior dation state +4.	n state which can then be oxidized back
	⊠ D	has par	rtially filled d orbitals in its +4 c	oxidation state.
				(Total for Question 6 = 1 mark)

7 A transition metal ion, M, forms a complex with a bidentate ligand, B. The formula the complex is MB ₃ so the shape of the complex is most likely to be		·	
	X		trigonal planar.
	×	В	pyramidal.
	×	C	trigonal bipyramidal.
	×	D	octahedral.
			(Total for Question 7 = 1 mark)
8			of the following lists all the types of bond that are present in a crystalline e of the compound tetraamminecopper(II) sulfate?
	×	A	lonic, covalent and dative covalent
	×	В	lonic and dative covalent
	×	C	lonic and covalent
	×	D	Covalent and dative covalent
_			(Total for Question 8 = 1 mark)
9	am	me	(Total for Question 8 = 1 mark) apound, X , is dissolved in water. Sodium hydroxide solution and dilute aqueous onia were added to different samples of this solution of X . In both, a precipitate d which dissolved in excess reagent. Compound X could be copper(II) sulfate.
9	am	me	apound, X , is dissolved in water. Sodium hydroxide solution and dilute aqueous onia were added to different samples of this solution of X . In both, a precipitate d which dissolved in excess reagent. Compound X could be
9	am	me Me	apound, X , is dissolved in water. Sodium hydroxide solution and dilute aqueous onia were added to different samples of this solution of X . In both, a precipitate d which dissolved in excess reagent. Compound X could be copper(II) sulfate.
9	am	nmo med A B	apound, X , is dissolved in water. Sodium hydroxide solution and dilute aqueous onia were added to different samples of this solution of X . In both, a precipitate d which dissolved in excess reagent. Compound X could be copper(II) sulfate. iron(II) sulfate.
9	am	nmo med A B	apound, X , is dissolved in water. Sodium hydroxide solution and dilute aqueous onia were added to different samples of this solution of X . In both, a precipitate d which dissolved in excess reagent. Compound X could be copper(II) sulfate. iron(II) sulfate. manganese(II) sulfate.
	am for	nmo A B C D	apound, X , is dissolved in water. Sodium hydroxide solution and dilute aqueous onia were added to different samples of this solution of X . In both, a precipitate d which dissolved in excess reagent. Compound X could be copper(II) sulfate. iron(II) sulfate. manganese(II) sulfate. zinc(II) sulfate.
	am for	nmo A B C D	apound, X , is dissolved in water. Sodium hydroxide solution and dilute aqueous onia were added to different samples of this solution of X . In both, a precipitate d which dissolved in excess reagent. Compound X could be copper(II) sulfate. iron(II) sulfate. manganese(II) sulfate. zinc(II) sulfate. (Total for Question 9 = 1 mark)
	am for some some some some some some some some	A B C D	apound, X , is dissolved in water. Sodium hydroxide solution and dilute aqueous onia were added to different samples of this solution of X . In both, a precipitate d which dissolved in excess reagent. Compound X could be copper(II) sulfate. iron(II) sulfate. manganese(II) sulfate. zinc(II) sulfate. (Total for Question 9 = 1 mark) arbon-carbon bonds in benzene are all the same length. The best evidence for omes from
	am for some some some some some some some some	A B C D	apound, X , is dissolved in water. Sodium hydroxide solution and dilute aqueous on were added to different samples of this solution of X . In both, a precipitate d which dissolved in excess reagent. Compound X could be copper(II) sulfate. iron(II) sulfate. manganese(II) sulfate. zinc(II) sulfate. (Total for Question 9 = 1 mark) arbon-carbon bonds in benzene are all the same length. The best evidence for omes from high resolution proton nmr spectroscopy.
	am for for	e cas co	apound, X , is dissolved in water. Sodium hydroxide solution and dilute aqueous onia were added to different samples of this solution of X . In both, a precipitate d which dissolved in excess reagent. Compound X could be copper(II) sulfate. iron(II) sulfate. manganese(II) sulfate. (Total for Question 9 = 1 mark) arbon-carbon bonds in benzene are all the same length. The best evidence for omes from high resolution proton nmr spectroscopy. X-ray diffraction.



11	Benzene reacts very slowly with chlorine but the reaction speeds up when finely divided iron is added. This is because		
	A the chlorine molecule donates an electron pair to the iron producing an electrophile.		
	■ B the iron reacts with chlorine to form iron(III) chloride which then acts as an electrophile in its reaction with benzene.		
	⊠ C	the iron reacts with chlorine to form iron(III) chloride which then acts as an electron pair acceptor.	
	\boxtimes D	iron is a transition metal and acts as a heterogeneous catalyst in the reaction.	
		(Total for Question 11 = 1 mark)	
12		culated that, as a result of delocalization, benzene has a stabilization energy of mol^{-1} . This means that	
		the enthalpy change for the conversion of benzene to cyclohexa-1,3,5-triene would be $+150~\rm kJ~mol^{-1}$.	
	■ B	the enthalpy change for the conversion of cyclohexa-1,3,5-triene to benzene would be $+150~\rm kJ~mol^{-1}$.	
	⊠ C	the enthalpy change for the conversion of cyclohexane to benzene is $+150~\rm kJ~mol^{-1}$.	
	⊠ D	the enthalpy change for the conversion of benzene to cyclohexane is $+150~\rm kJ~mol^{-1}$.	
		(Total for Question 12 = 1 mark)	
13	When a gas jar containing methylamine is opened near an open bottle of concentrated hydrochloric acid, white smoke is seen. The chemical formula of the white smoke is		
	⋈ A	CH ₃ NH ₄ Cl	
	⊠ B	CH ₃ NH ₃ Cl	
	⊠ C	CH ₃ NH ₂ CI	
	⋈ D	NH ₄ CI	
	(Total for Question 13 = 1 mark)		

- **14** When benzoyl chloride, C₆H₅COCl, is added to methylamine at room temperature, the main organic product of the reaction is

(Total for Question 14 = 1 mark)

- 15 In an aqueous solution with a pH of 3, the amino acid glycine exists mainly as
 - ✓ A✓ C—OH✓ H₂N—CH₂
 - B C—O⁻H₂N—CH₂
 - C C—OH
 H₃N—CH₂
 - D□ C—O

(Total for Question 15 = 1 mark)

	is the total number of peaks due to singly charged ions in the complete mass um of chlorine, Cl ₂ ?
⊠ A	Two
⊠ B	Three
⊠ C	Four
	E.
⊠ D	Five
The lo	(Total for Question 16 = 1 mark w resolution proton nmr spectrum of a compound contains two peaks. Which
' The lo	(Total for Question 16 = 1 mark we resolution proton nmr spectrum of a compound contains two peaks. Which following compounds could not give this spectrum?
' The lo	(Total for Question 16 = 1 mark w resolution proton nmr spectrum of a compound contains two peaks. Which
The loof the	(Total for Question 16 = 1 mark we resolution proton nmr spectrum of a compound contains two peaks. Which following compounds could not give this spectrum? Propane
The love of the A B C	(Total for Question 16 = 1 mark w resolution proton nmr spectrum of a compound contains two peaks. Which following compounds could not give this spectrum? Propane Butane

- **18** A compound, **P**, has the following properties:
 - **P** forms a red precipitate when heated with Fehling's or Benedict's solution.
 - **P** forms a pale yellow precipitate when warmed with iodine dissolved in aqueous sodium hydroxide.

P could be

(Total for Question 18 = 1 mark)

- 19 10 cm³ of a gaseous hydrocarbon was mixed with excess oxygen and ignited. The gas volumes were measured at room temperature and pressure before and after combustion and it was found that the total gas volume had contracted by 20 cm³. Given that combustion was complete, the formula of the hydrocarbon was

 - B C₄H₆
 - C C₄H₂
 - \square **D** C_4H_{10}

(Total for Question 19 = 1 mark)

20	O Steam distillation may be used in the purification of some compounds. The use of this technique depends on the compound		
	⊠ A	forming a single layer with water.	
	⊠ B	forming two layers with water.	
	⊠ C	having a lower boiling temperature than water.	
	⊠ D	being flammable.	
		(Total for Question 20 = 1 mark)	

TOTAL FOR SECTION A = 20 MARKS

SECTION B

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

- 21 Potassium manganate(VII) can be used in redox titrations to determine the concentration of iron(II) ions and ethanedioate ions in aqueous solution. Aqueous solutions of potassium manganate(VII) are unstable, so it is often standardized using solutions of iron(II) ammonium sulfate, freshly prepared from Mohr's salt, $Fe(NH_4)_2(SO_4)_2.6H_2O$.
 - (a) Use the relevant ionic half-equations, and standard reduction potentials on page 17 of the Data Booklet, to answer the following questions. State symbols are not required in the equations.

(i)	Write the ionic half-equation for the reduction of manganate(VII) ions in acid
	solution.

(1)

(ii) Write the ionic half-equation for the oxidation of water to form oxygen and hydrogen ions.

(1)

(iii) Combine your answers to (a)(i) and (a)(ii) to derive an equation to show the production of oxygen in acidified manganate(VII) solution.

(1)

(iv) Calculate $E_{\rm cell}^{\ominus}$ for the reaction in (a)(iii) and hence explain why aqueous solutions of potassium manganate(VII) are unstable.

(2)



(b) 250 cm³ of a solution containing 10.00 g of Mohr's salt was prepared. Separate 25.0 cm³ samples of this solution were pipetted into conical flasks, excess sulfuric acid added and then each mixture was titrated against potassium manganate(VII) solution.	
The mean titre was 25.85 cm ³ .	
*(i) Describe in outline how you would prepare the 250 cm³ of Mohr's salt solution, given 10.00 g of the solid.	(4)
(ii) State what you would see at the end-point of the titration.	(1)
(iii) Write the ionic equation showing that 1 mol of manganate(VII) ions reacts with 5 mol of iron(II) ions in acid conditions. State symbols are not required.	(1)



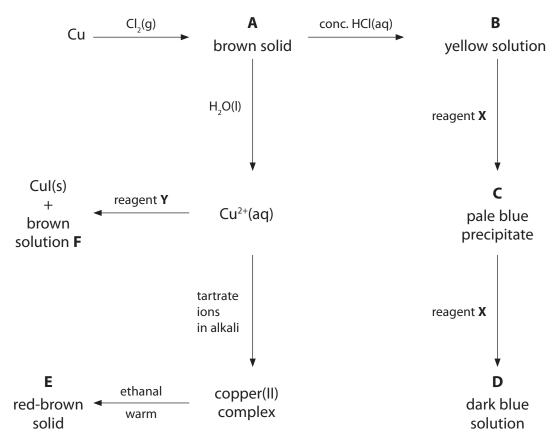
(iv) The concentration of the potassium manganate(VII) solution was stated to be $0.0200~\rm mol~dm^{-3}$. Calculate the percentage of the potassium manganate(VII) that had reacted between its preparation and the titration.

The molar mass of Mohr's salt is 392 g mol^{-1} .

(4)

(Total for Question 21 = 15 marks)

22 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**.

(6)

A	
В	
C	
D	
Ε	
F	
	(ii) Identify the reagents X and Y .
Χ.	

(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) Aqueous 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) Use the standard reduction potentials on page 17 of the Data Booklet to calculate $E_{\rm cell}^{\div}$ for this disproportionation. Hence show that this reaction is thermodynamically feasible.	(2)
(Total for Question 22 = 21 n	narks)



23 Compound **P** is a white crystalline solid with the following percentage composition by mass.

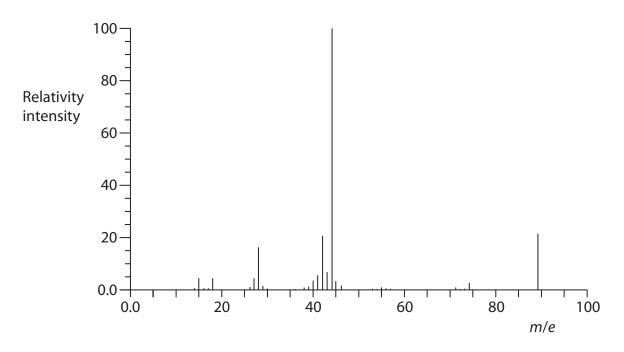
Element	% composition by mass
carbon	40.44
hydrogen	7.87
oxygen	35.96
nitrogen	15.73

(a) Calculate the empirical formula of **P**. You **must** show your working.

(3)



(b) The mass spectrum of ${\bf P}$ is shown below.



(i) Label the molecular ion on the mass spectrum using the symbol $\mathsf{M}^{\scriptscriptstyle{+}}.$

(1)

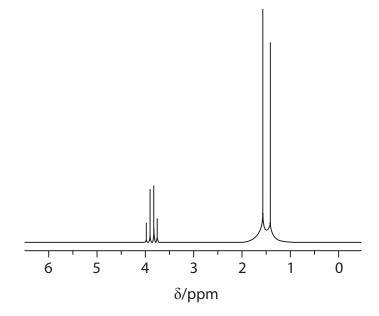
(ii) Deduce the molecular formula of **P**.

(1)

(c)	P was dissolved in sodium carbonate solution and, on heating, a colourless gas, which turned lime water cloudy, was very slowly evolved. When an aqueous solution of P was mixed with an aqueous solution of copper(II) sulfate, the blue colour of the copper(II) sulfate solution darkened.	
	(i) Use the results of these two experiments to deduce the functional groups that are present in P . Explain your answers.	(4)
	(ii) There are several compounds which have the formula you have deduced in (b)(ii) and contain the two functional groups you have identified in (c)(i).	
	Draw the structural or displayed formulae of two of these compounds.	(2)



(iii) The nuclear magnetic resonance spectrum of compound **P** is shown below.



This part of the spectrum only shows the peaks due to the hydrogen atoms which are attached directly to carbon atoms.

Use this spectrum to deduce the structure of **P**. Justify your answer.

(2)

(d) Explain why **P** is a solid at room temperature and pressure.

(1)

(Total for Question 23 = 14 marks)

TOTAL FOR SECTION B = 50 MARKS



SECTION C

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

24

Phthalates

The benzene dicarboxylic acids and their esters are important industrial compounds. The structures of two of these acids are shown below.

benzene-1,2-dicarboxylic acid

benzene-1,4-dicarboxylic acid

Benzene-1,2-dicarboxylic acid is manufactured by the catalytic oxidation of naphthalene to phthalic anhydride which is then hydrolysed. This reaction sequence is summarised below.

The use of naphthalene as a source of these compounds gave rise to the common names, phthalic acid for benzene-1,2-dicarboxylic acid and terephthalic acid for benzene-1,4-dicarboxylic acid.

The alkyl esters derived from benzene-1,2-dicarboxylic acid are known as phthalates.

Phthalates are used as plasticisers to increase the flexibility of many common plastics. However, some phthalates are known to be endocrine disruptors and recent studies have raised concerns about their role in the fall in human fertility rates. Because phthalates are used in making plastic drinks bottles and leach readily out of the structure, they are easily ingested.

The polyester *Terylene* is derived from benzene-1,4-dicarboxylic acid and ethane-1,2-diol.

(a) One method of preparing benzene dicarboxylic acids in the laboratory is from methylbenzene in the sequence shown below.

(i) Identify reagent **M**, by name or formula.

(1)

(ii) Write the equation for the reaction between ${\bf M}$ and ${\rm AlCl}_{_3}$ to form an electrophile.

(1)

(iii)	Give the mechanism for the reaction of methylbenzene with your electrophile in (a)(ii).	
		(3)
(iv)	Suggest why methylbenzene reacts faster than benzene in this type of	
	reaction.	(2)
		(-)
(v)	In the second step of the synthesis, the potassium manganate(VII) is an oxidizing agent.	
(v)	oxidizing agent.	
(v)		(1)
(v)	oxidizing agent.	



 (b) In the oxidation of naphthalene to phthalic anhydride, the catalyst is vanadium(V) oxide. With fresh catalyst, the reaction occurs at 360 °C but, over time, the temperature must be slowly increased as the catalyst activity decreases. (i) State the property which gives transition metal compounds, such as vanadium(V) oxide, catalytic properties. 	(1)
(ii) Suggest why the catalyst activity decreases over time.	(1)
 (c) The simplest phthalate of benzene-1,2-dicarboxylic acid is its dimethyl ester (1,2-dimethyl benzene-1,2-dicarboxylate). (i) Draw the structure of this phthalate. 	(1)



from plastic bottles.	/01
	(2)
*(iii) Suggest how a plasticiser works.	
	(2)
One way of making <i>Terylene</i> is by converting benzene-1,4-dicarboxylic active di-acyl chloride and then reacting it with ethane-1,2-diol.	id into
	oxvlic
(i) Suggest a reagent that could be used to convert benzene-1,4-dicarbo	
(i) Suggest a reagent that could be used to convert benzene-1,4-dicarbo acid into the di-acyl chloride.	
	(1)
acid into the di-acyl chloride.	
acid into the di-acyl chloride. (ii) Suggest an advantage of using the di-acyl chloride rather than the	
acid into the di-acyl chloride. (ii) Suggest an advantage of using the di-acyl chloride rather than the	(1)
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acid into the di-acyl chloride. (ii) Suggest an advantage of using the di-acyl chloride rather than the	(1)



	TOTAL FOR SECTION C = 20 MAR
	(Total for Question 24 = 20 mar
What would be the by-product of this re	eaction?
v) In practice, the manufacture of <i>Terylene</i> exchange in which ethane-1,2-diol reac benzene-1,4-dicarboxylic acid.	
ii) Draw the structure of the polyester, <i>Tery</i>	3 · · · · 3 · · · · · · · · · · · · · · · · · · ·



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7	(17)	19.0 F fluorine 9	35.5 Cl chlorine 17	79.9	Br bromine	126.9	-	iodine 53	[210]		astatine 85		oeen repo		175
9	(16)	16.0 O oxygen 8	32.1 S sulfur 16	79.0	Selenium	177.6	Ъ	tellurium 52	[509]	8	polonium 84	10000000	116 have	iticated	473
2	(15)	14.0 N nitrogen	31.0 P	74.9	As arsenic	171.8	Sb	antimony 51	209.0	Bi	bismuth 83		nbers 112-	but not fully autnenticated	140
4	(14)	12.0 C carbon 6	Si silicon	72.6	Ge germanium	118.7	S	tin 20	207.2	Ъ	lead 82		atomic nur	DUT NOT I	147
3	(13)	10.8 B boron 5	27.0 Al aluminium 13	2.69	Ga gallium	114.8	_	indium 49	204.4	F	thallium 81		Elements with atomic numbers 112-116 have been reported		145
	,		(12)	65.4	Zinc	112.4	3	cadmium 48	200.6	Нg	mercury 80		Elem		14.3
			(11)	63.5	Copper	6 201	Ag	silver 47	197.0	Αn	gold 79	[272]	Rg	oentgenium 111	150
			(01)	58.7	nickel	106.4	Р	palladium 46	195.1	ដ	platinum 78	_		damstadtium r 110	157
			(6)	58.9	Co	102.9	윤	rhodium 45	192.2	<u>-</u>	iridium 77	[368]		109	152
	1.0 Hydrogen		(8)	55.8	iron y	101.1	æ	ruthenium 44	190.2	os	osmium 76	[277]	- 8	108	150
			6	54.9	Мп папдалеѕе	C7		technetium 43	186.2	Re	rhenium 75	_		pohrnum 107	[447]
		nass ool	(9)	52.0	Cr Mn	95.9	Wo	molybdenum 42	183.8	>	tungsten 74	[597]	Sg	seaborgium 106	144
	Key	relative atomic mass atomic symbol name atomic (proton) number	(5)	50.9	Ę	67.9	£	niobium 41	180.9	Та	tantalum 73	_	g E	dubnium 105	171
		relati ato atomic	(4)	47.9	Ti titanium	91.2	Zr	zirconium 40	178.5		hafnium 72	[261]	₽,	rutherfordium 104	140
			(3)	45.0	Scandium	6.88	>	yttrium 39	138.9	La*	lanthanum 57	[227]		actinium 89	
2	(2)	9.0 Be beryllium 4	24.3 Mg magnesium 12	40.1	calcium	87.6	Sr	strontium 38	137.3		barium l	[526]	Ra	radium 88	
-	(1)	6.9 Li lithium	Na sodium 11	39.1	K potassium	85.5		rubidium 37	132.9	S	caesium 55	[223]	F.	rrancium 87	

* Lanthanide series
* Actinide series

				_			
175	Ľ	lutetium	71	[257]	۲	lawrencium	103
173	χp	ytterbium	70	[254]	8	nobelium	102
169	Ш	thulium	69	[526]	ΡW	mendelevium	101
167	д	erbium	89	[253]	Fm	fermium	100
165	유	holmium	67	[254]	Es	einsteinium	66
163	D	dysprosium	99	[251]	უ	californium	86
159	ТÞ	terbium	65	[245]	BK	berkelium	97
157	PS	gadolinium	64	[247]	5	aurium	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	09	238	_	uranium	92
141	P	praseodymium	26	[231]	Pa	protactinium	91
140	Ce	cerium	58	232	두	thorium	90