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Chemistry Unit 5 WCH05

Past Paper

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Write your name here Surname	Other	names
Pearson Edexcel	Centre Number	Candidate Number
(homistry		
Chemistry Advanced Unit 5: General Principle Metals and Orga (including synop	es of Chemistry I Inic Nitrogen Che	
Advanced Unit 5: General Principle Metals and Orga	es of Chemistry I Inic Nitrogen Che otic assessment)	Paper Reference
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Advanced Unit 5: General Principle Metals and Orga (including synop Monday 15 June 2015 – Aft	es of Chemistry I Inic Nitrogen Che otic assessment)	Paper Reference

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.

Turn over ▶



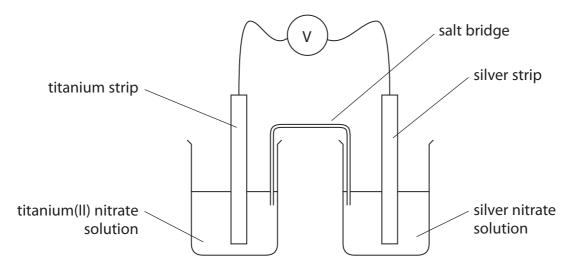
SECTION A

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	The	ОХ	idation number of oxygen in hydrogen peroxide, H ₂ O ₂ , is
	\times	Α	+2
	× E	В	0
	× (C	– 1
	X r	D	-2
			(Total for Question 1 = 1 mark)
2		-	r reacts with nitrate ions, NO_3^- , in acid conditions to form copper(II) ions and en(IV) oxide.
			sidering the changes in the oxidation numbers of copper and nitrogen, it can luced that the redox reaction involves
	\boxtimes /	Α	1 mol of copper reacting with 2 mol of nitrate ions.
	×	В	2 mol of copper reacting with 1 mol of nitrate ions.
	\boxtimes	C	1 mol of copper reacting with 4 mol of nitrate ions.
	\boxtimes	D	4 mol of copper reacting with 1 mol of nitrate ions.
			(Total for Question 2 = 1 mark)
3	The	Ece	for a reaction is $+1.00$ V. What can be deduced about this reaction?
	\times	A	At equilibrium there will be mainly reactants.
	X	В	At equilibrium there will be mainly products.
		C	At equilibrium there will be approximately equal amounts of reactants and products.
		D	It is impossible to estimate the relative amounts of reactants and products at equilibrium because the $E_{\rm cell}$ was not measured under standard conditions.
			(Total for Question 3 = 1 mark)

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4 The cell shown in the diagram below was set up.



The emf of this cell under standard conditions is +2.43 V. The E^{\ominus} value for the Ag⁺(aq)|Ag(s) half cell is +0.80 V.

What is the E^{\oplus} value for the $Ti^{2+}(aq)|Ti(s)$ half cell?

- **B** +1.63 V

(Total for Question 4 = 1 mark)

- 5 In the hydrogen-oxygen fuel cell, oxygen is
 - **A** oxidized at the negative electrode.
 - **B** oxidized at the positive electrode.
 - **C** reduced at the negative electrode.
 - **D** reduced at the positive electrode.

(Total for Question 5 = 1 mark)

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6	The el	ectronic c	onfigı	uratio	n of th	ne ele	ment	vanadium is
		3d					4s	
	⋈ A	[Ar]	1↓	↑↓	1			
	В	[Ar]	1	1	1	1	1	
	⊠ C	[Ar]	1	1	1	1		1
	⊠ D	[Ar]	1	1	1			$\uparrow\downarrow$
								(Total for Question 6 = 1 mark)
7	 solution. The best explanation for this property is that A the differences in the successive ionization energies are similar to the differences in hydration enthalpies of the ions. B all the ions are formed by the removal of electrons from the d subshell. C the ionization energies of transition metals are low. D the hydration enthalpies of transition metal ions are always more exothermic than those of ions of s and p block metals. 							
								(Total for Question 7 = 1 mark)
8	Which	of the fol	lowin	g spec	cies n e	ever (combi	ine with ligands to form complexes?
		Positivel	y char	ged ic	ons of	d blo	ck ele	ements.
	⊠ В	Neutral a	atoms	of d b	olock	eleme	ents.	
		Negative	ely cha	arged	ions c	of d bl	ock e	lements.
	D Positively charged ions of p block elements.							

(Total for Question 8 = 1 mark)

9	Transition metal complexes are formed when ethanedioate ions and ethanoate ions are added separately to aqueous solutions of transition metal ions.							
	The complexes formed by the bidentate ethanedioate ligands are more stable than the complexes formed by the monodentate ethanoate ligands. This is because							
	A ethanedioate ligands form stronger bonds with the metal ion of a complex than do ethanoate ligands.							
	В	the formation of ethanedioate complexes increases the number of particles in the solution.						
	⊠ C	ethanedioic acid is a stronger acid than ethanoic acid.						
	\boxtimes D	ethanedioic acid is a weaker acid than ethanoic acid.						
		(Total for Question 9 = 1 mark)						
10		temperature of the nitration of benzene is allowed to rise too high, dinitration initration can occur. This is evidence that the						
	\square A	nitro group is electron withdrawing.						
	⊠ B	nitro group is electron donating.						
	⊠ C	delocalisation energy of nitrobenzene is greater than that of benzene.						
	⊠ D	delocalisation energy of nitrobenzene is less than that of benzene.						
		(Total for Question 10 = 1 mark)						
11	The be	enzene molecule may be represented in two ways:						
		structure I structure II						
		of the following does not provide evidence that structure I is the better entation of benzene?						
	⊠ A	Infrared spectroscopy						

■ B High performance liquid chromatography

□ C Thermochemistry

 $\ \square$ **D** X-ray diffraction

(Total for Question 11 = 1 mark)

12 When phenol, C_6H_5OH , reacts with excess bromine water, the organic product is

0

A

⋈ B

⊠ C

Br

(Total for Question 12 = 1 mark)

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13 Ammonia, butylamine and phenylamine are dissolved in separate samples of water to form solutions of the same concentration. The pH of each solution was measured.						
The order o	of increasing pH will be	e				
🖾 A amı	monia < butylamine <	phenylamine				
■ B but	ylamine < ammonia <	phenylamine				
☑ C phe	enylamine < ammonia	< butylamine				
☑ D phe	enylamine < butylamine	e < ammonia				
		(Total for Question 13	s = 1 mark)		
	14 Aromatic amines may be converted into benzenediazonium ions. What are the most suitable reagents and conditions for this reaction?					
	Reagent 1	Reagent 2	Temperature / °C			
	NaNO ₂	sulfuric acid	+55			
⊠ B	■ B NaNO ₃ hydrochloric acid +5					
⊠ C	NaNO ₂	hydrochloric acid	+5			
⊠ D	NaNO ₃	sulfuric acid	+55			

(Total for Question 14 = 1 mark)

15	ne repeat unit oi	the polymer	formed from	etnane-1,2-	-diol and	ethanedioic acid	IS
----	-------------------	-------------	-------------	-------------	-----------	------------------	----

- A -OCH₂CH₂OOCCO-
- B -OCH₂CH₂OCOOCO-
- ☑ C -OCH₂OOCCH₂CO-
- ☑ D –OCCH₂CH₂OOCO−

(Total for Question 15 = 1 mark)

- **16** Poly(ethenol) is an example of
 - ☑ A an addition polymer that is soluble in water.
 - **B** an addition polymer that is insoluble in water.
 - **C** a condensation polymer that is soluble in water.
 - **D** a condensation polymer that is insoluble in water.

(Total for Question 16 = 1 mark)

17 A compound gives an orange precipitate with 2,4-dinitrophenylhydrazine, Brady's reagent, but does **not** react with ammonia in the cold. The compound could be

$$H_3C$$
 CH
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$$\begin{array}{c|c}
 & CH_2 \\
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(Total for Question 17 = 1 mark)

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

 \times D

18 Three compounds are possible monomers in the formation of a polymer:

	H ₂ C—HC=	CH—CH ₂
'	H ₂ N	NH ₂
II	O C—H₂C—	O CH ₂ —C CI
III	H₂C—H₂C—	CH ₂ —CH ₂ OH

Which of the following compounds could **not** react in the stated combination to form a polymer?

- **B** I in combination with II
- C II in combination with III
- D I in combination with III

(Total for Question 18 = 1 mark)

19 Pentan-3-one reacts with 2,4-dinitrophenylhydrazine to form a derivative which has a melting temperature of 156 °C.

A student attempted to synthesise pentan-3-one, and converted some of the product into the same derivative. The student's derivative melted gradually from 148 °C to 158 °C.

It is most likely that the student had synthesised

- A pure pentan-3-one.
- B impure pentan-3-one.
- approximately equal amounts of two carbonyl derivatives, one with a melting temperature of 148°C and the other with a melting temperature of 158°C.
- **D** a compound that was not a ketone.

(Total for Question 19 = 1 mark)

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20	O The purification of organic compounds prepared in aqueous mixtures often involves solvent extraction. Desirable properties of the solvent used include that it is						
	X	A	fully miscible in water and has a high boiling temperature.				
	×	В	fully miscible in water and has a low boiling temperature.				
	☐ C immiscible in water and has a high boiling temperature.						
	X	D	immiscible in water and has a low boiling temperature.				
			(Total for Question 20 = 1 mark)				

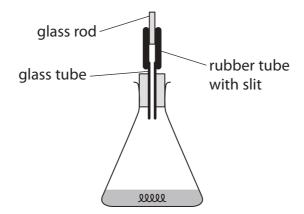
TOTAL FOR SECTION A = 20 MARKS

WCH05

SECTION B

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

- 21 Steel is the world's most important structural metal; it is strong and cheap but it corrodes rapidly if unprotected. In its simplest form, steel is an alloy of iron and carbon.
 - (a) The following method was used to determine the percentage of iron in a sample of wire. Exactly 1.25 g of the wire was placed in a conical flask and about 50 cm³ of dilute sulfuric acid (an excess) was added. The flask was closed as shown in the diagram below.



When all of the iron in the wire had been converted to iron(II) sulfate, the contents of the flask were used to make 250.0 cm³ of solution with distilled water.

25.00 cm³ portions of this final solution were placed in a conical flask, acidified with an equal volume of dilute sulfuric acid and then titrated with a potassium manganate(VII) solution of concentration 0.0195 mol dm⁻³. The mean titre was 22.15 cm³.

(i) Write the equation for the reaction between iron and dilute sulfuric acid. Include state symbols in your answer.

(1)

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the apparatus shown in the diagram works.	(2)
(iii) State the essential steps of the procedure for making to 250.0 cm³ for use in the titration.	
	(3)
(iv) Write the ionic equation for the titration reaction to shape and the contract with 1 mol of management (VIII) ions. State symbol	
react with 1 mol of manganate(VII) ions. State symbol	(1)

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three significant figures.

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(v) Calculate the percentage by mass of iron in the wire. Give your answer to

(4)

(vi) Describe the colour change at the end-point of the titration.

(1)

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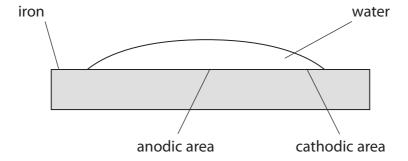
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MAC	۸	_

(vii) One student who carried out this experiment forgot to acidify the mixture in
the conical flask before the titration.
A brown precipitate formed before the end-point.

Identify the brown precipitate and explain how this error affects the titration

variation.	
	(3)

(b) The rusting of iron is an electrochemical process. A piece of iron with a droplet of water on its surface operates as an electrochemical cell. In the first stage of corrosion, iron is oxidized to iron(II) ions in the anodic area and the electrons produced travel to the cathodic area where oxygen from the air is reduced.



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	\circ	\mathbf{o}

(i)	From the information about standard electrode potentials on pages 14 and 15 of the Data Booklet, write the ionic half equations for the reactions taking place at the anodic area and at the cathodic area. State symbols are not required.	e
Anodic are		(2)
Cathodic a	area	
(::)	Calculate Γ^{\oplus} for the quantity of the (b)(i)	
(11)	Calculate E_{cell}^{\oplus} for the overall reaction in (b)(i).	(1)
(iii) By considering the rusting mechanism described in part (b), suggest why the	
(111)	presence of salt in the water droplet speeds up rusting.	(1)
(iv) The corrosion of oil pipelines made of steel is prevented by connecting the	
•	pipeline to magnesium blocks. Suggest how this method works.	(1)
	(Total for Question 21 = 20 ma	rks)

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	formula of the copper species in A , B and C . and spresent in each species.	You should include all
or the liga	ands present in each species.	(3)
/··> = 1 ·		
(II) Explain w	hy solution A is coloured.	(4)
(iii) Explain w	hy solution A is a different colour to solution	
		(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.
Step 1	The solid was dissolved in the minimum amount of hot ethanol.
Step 2	The hot solution was filtered .
Step 3	The filtrate was cooled in an ice bath .
Step 4	The mixture was filtered using suction filtration .
	(Total for Question 22 = 14 marks)

23 This question is about an unknown organic compound, Q.

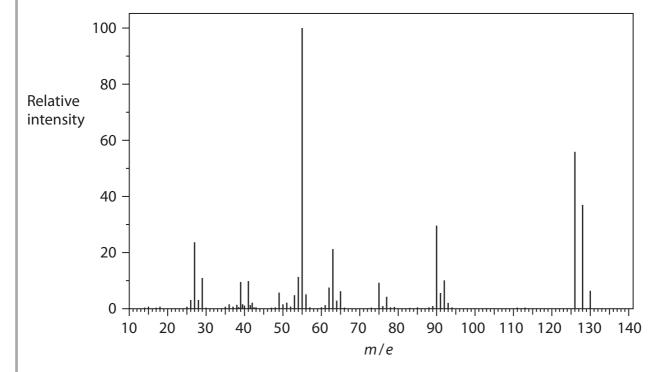
The percentage composition by mass of ${\bf Q}$ is

carbon = 37.8%; hydrogen = 6.30%; chlorine = 55.9%.

(a) Calculate the empirical formula of **Q**.

(2)

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



(i) Use your answer to part (a) and the mass spectrum to deduce the molecular formula of **Q**.

(1)



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(ii) Explain why there are three peaks in the molecular ion region (<i>m/e</i> from 126 to 130) of the mass spectrum of Q .	(2)
(iii) State why the peak at $m/e = 126$ is the highest of the three peaks in the molecular ion region.	(1)

(5)

*(iv) **Q** reacts with aqueous sodium hydroxide to form an organic compound, **R**.

The functional groups of compound **R** are on different carbon atoms.

Compound **R** reacted slowly with sodium, producing a total of one mole of hydrogen gas per mole of **R**.

When compound **R** was heated under reflux with excess acidified sodium dichromate(VI), an organic compound, **S**, was formed. Compound **S** reacted rapidly with sodium, producing a total of 0.5 mol of hydrogen gas per mole of **S**.

Draw the **displayed** formulae of the two possible structures for compound **S**. Explain how your structures are consistent with these data.

Structure I Structure II Explanation.....

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	TOTAL FOR SECTION B = 50 MA	RKS
	(Total for Question 23 = 16 ma	rks)
Equation		
R =	j	(3)
(vi)	Name compound R and give the equation for its reaction with sodium.	
	that you have given in (b)(iv) is compound S . Explain your reasoning.	(2)
	producing a pale yellow precipitate with an antiseptic smell. Identify the pale yellow precipitate and hence identify which of the structures	
	Compound S reacted with iodine dissolved in aqueous sodium hydroxide,	

SECTION C

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

24

Oil of cloves

Oil of cloves is a traditional remedy for toothache; it is a topical treatment, which means that it is applied externally. The main active ingredient of oil of cloves is eugenol, which also gives cloves their characteristic smell. The structure of eugenol is shown below.

(a) A student suggested that eugenol could be prepared by an electrophilic substitution of 2-methoxyphenol, which is produced in the gut of desert locusts. This compound is one of the main components of the pheromones that cause locust swarming. The structure of 2-methoxyphenol is shown below.

(i) Use your knowledge of electrophilic substitution reactions to suggest the structure of an electrophile that might be used in this reaction.

(1)

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(ii) Suggest an organic compound that could be used to produce the electrophile that you have given in (a)(i).

(1)

(iii) Write the mechanism for the electrophilic substitution to prepare eugenol from 2-methoxyphenol, using the electrophile that you have given in (a)(i).

(3)

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(b) Eugenol has been used in the manufacture of vanillin, the compound responsible for the flavour of vanilla. The process involves the conversion of eugenol to its isomer isoeugenol, which is then oxidized by hydrogen peroxide with a vanadium(V) oxide catalyst.

$$H_2O_2/V_2O_5$$
 + H_2C CH_3 + CH_3 vanillin methyl vanillyl ketone

(i) Isoeugenol exists as two stereoisomers, whereas eugenol has just one structure. State the type of stereoisomerism shown by isoeugenol and explain why it can show this type of stereoisomerism, whereas eugenol does not.



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ii) Discuss whether or not high resolution proton nmr spectroscopy could be used to distinguish between eugenol and a single isomer of isoeugenol. You should consider the numbers of peaks in the spectra, their splitting patterns and the areas under the peaks. You are not expected to suggest chemical shift values.								
Chemical stille values.	(4)							
(iii) Suggest, in outline, a mechanism by which vanadium(V) oxide acts as a catalyst for the reaction between hydrogen peroxide and isoeugenol to form								
vanillin. You are not expected to give equations for the mechanism.	(2)							

(iv) A student suggested that the infrared spectroscopy data on pag	je 5 of the
Data Booklet could be used to show whether the purified vanilli	n contains a
trace of methyl vanillyl ketone.	

Explain the basis of this idea and suggest why it may not work in practice.

(2)

(c) The Zeisel method for determining the purity of vanillin involves estimating the percentage by mass of methoxy group (CH₃O) in the sample and comparing this with the percentage in pure vanillin.

In a sequence of reactions, each methoxy group produces iodine which is estimated by titration with a sodium thiosulfate solution of known concentration. The Zeisel sequence is

Step 1
$$ROCH_3 + HI \rightarrow ROH + CH_3I$$

Step 2
$$CH_3I + Br_2 \rightarrow CH_3Br + IBr$$

Step 3 IBr +
$$3H_2O + 2Br_2 \rightarrow HIO_3 + 5HBr$$

Step 4
$$IO_3^- + 5I^- + 6H^+ \rightarrow 3I_2 + 3H_2O$$

Step 5
$$I_2 + 2S_2O_3^{2-} \rightarrow S_4O_6^{2-} + 2I^-$$

(i) Deduce the number of moles of thiosulfate ions that are equivalent to one mole of methoxy group. Explain your answer.

(2)

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(ii) A sample of vanillin was found to have 20.09% by mass of methoxy group. Calculate the percentage purity of the vanillin.

(3)

(Total for Question 24 = 20 marks)

TOTAL FOR SECTION C = 20 MARKS TOTAL FOR PAPER = 90 MARKS 0 (8)

9

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67

65

49

europium 63

61

9

58

Pr Pr 59

[242] 62

einsteinium

berkelium 97

Curium 96

neptunium plutonium

uranium

rotactinium

232 **Th** thorium

93

92

16

9

The Periodic Table of Elements

	-			_	_			_				_	_			_	_			_	_			_							
(18)	(0/)	4. J	helium	2	20.2	Ne	neon	10	39.9	Αr	argon	18	83.8	궃	krypton	36	131.3	Xe	xenon	24	[222]	R	radon	98		ted					
				(17)	19.0	L	fluorine	6	35.5	ರ	chlorine	17	6.62	Br	bromine	35	126.9	-	iodine	53	[210]	Αt	astatine	85		een repor			175	Ľ	lutetium
				(16)	16.0	0	oxygen	∞	32.1	s	sulfur	16	0.67	Se	selenium	34	127.6	Тe	tellurium	25	[506]	8	polonium	84		116 have b	ticated		173	ХÞ	ytterbium
				(12)	14.0	z	nitrogen	7	31.0	۵	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		169	Ħ	
				(14)	12.0	U	carbon	9	28.1	Si	silicon	14	72.6	Ge	germanium	32	118.7	Sn	tin	20	207.2	Ъ	lead	82		atomic nur	but not fi		167	Ę	erbium
				(13)	10.8	В	boron	2	27.0	¥	aluminium	13	2.69	ga	gallium	31	114.8	드	indium	49	204.4	F	thallium	81		ents with			165	유	holmium
											ć	(71)	65.4	Zu	zinc	30	112.4	В	cadmium	48	200.6	Ę	mercury	80					163	D	dysprosium holmium
											(**)	(11)	63.5	J	copper	29	107.9	Ag	silver	47	197.0	Αn	plog	79	[272]	Rg	roentgenium	111	159	4	terbium
											0	(01)	58.7	ź	nickel	28	106.4	Pd	palladium	46	195.1	చ	platinum	78		S D	Ē	110	157	В	gadolinium
											ę	(6)	58.9	ပိ	cobalt	27	102.9	뫈	rhodium	45	192.2	<u>-</u>	iridium	77	[568]	¥	Ę	109	152	En	europium
	1.0	Œ	hydrogen 1	-							Ó	(8)	55.8	Fe			101.1	Ru	ruthenium	44	190.2	õ	osmium	76	[277]	꿈	hassium	108	150	Sm	samarium
				100							Ĺ	2	54.9	W	manganese	25	[86]	2	technetium	43	186.2	Re	rhenium	75	100		ğ	107	[147]	Pm	promethium
					mass	pol	1	umber				(9)	52.0		Ę	24	95.9	Wo	molybdenum	42	183.8	>	tungsten	74	[392]	Sg	seaborgium	106	144	PN	praseodymium promethium samarium europium gadolinium
			1	Key	relative atomic mass	atomic symbol	name	atomic (proton) number			í	(5)	50.9	>	vanadium	23	92.9	g	niobium	41	180.9	Тa	tantalum	73	_		Ε	105	141	P	praseodymium
				Ç.	relati	ato		atomic			5	(4)	47.9	ï	titanium	22	91.2	Zr	zirconium	40	178.5	Ŧ	hafnium	72	[261]	¥	nutherfordium	104	140		cerium
											Ç	(3)	45.0	Sc	scandium	21	88.9	>	yttrium	39	138.9	La*	lanthanum	57	[227]	Ac*	actinium	88	•	sa	
				(2)	0.6	Be	beryllium	4	24.3	Mg	magnesium	12	40.1	Ç	calcinm	20	9.78	Sr	strontium	38	137.3	Ba	barium	26	[526]	Ra	radium	88		* Lanthanide series	* Actinide series
				(1)	6.9	'n	lithium	3	23.0	Na	sodium	1	39.1	×	potassium	19	85.5	&	rubidium	37	132.9	ర	caesium	22	[223]	Ė	francium	8/		* Lant	* Actin