



Cambridge International AS & A Level

CANDIDATE NAME				
CENTRE NUMBER		CANDIE NUMBE		

CHEMISTRY 9701/41

Paper 4 A Level Structured Questions

October/November 2022

2 hours

You must answer on the question paper.

No additional materials are needed.

INSTRUCTIONS

- Answer all questions.
- Use a black or dark blue pen. You may use an HB pencil for any diagrams or graphs.
- Write your name, centre number and candidate number in the boxes at the top of the page.
- Write your answer to each question in the space provided.
- Do not use an erasable pen or correction fluid.
- Do not write on any bar codes.
- You may use a calculator.
- You should show all your working and use appropriate units.

INFORMATION

- The total mark for this paper is 100.
- The number of marks for each question or part question is shown in brackets [].
- The Periodic Table is printed in the question paper.
- Important values, constants and standards are printed in the question paper.

 ${\bf 1} \quad \hbox{Potassium chloride, KC} \it{l}, \hbox{ and magnesium chloride, MgC} \it{l}_{\it{2}}, \hbox{ are both ionic solids}.$

Table 1.1

energy change	value/kJ mol ⁻¹
standard enthalpy change of solution, $\Delta H_{\text{sol}}^{\text{e}}$, of KC1	+15
lattice energy, $\Delta H_{\text{latt}}^{e}$, of KC $l(s)$	-7 01
standard enthalpy change of hydration, $\Delta H_{\text{hyd}}^{\text{e}}$, of K ⁺	-322
standard enthalpy change of hydration, $\Delta H_{\text{hyd}}^{\text{e}}$, of C l^-	-364
standard enthalpy change of solution, $\Delta H_{\mathrm{sol}}^{\mathrm{e}}$ of MgC l_2	–155
lattice energy, $\Delta H_{\text{latt}}^{\text{e}}$, of MgC l_2 (s)	-2493

		standard enthalpy c	hange of hydration,	$\Delta H_{\text{hyd}}^{\Theta}$, of C l^-	-364	
		standard enthalpy c	hange of solution, Δa	H_{sol}^{Θ} , of MgC l_2	–155	
		lattice energy, ∆H ^e _{latt}	, of MgC l_2 (s)		-2493	
(a)	potas	plete the energy cyclessium chloride, KC <i>l</i> , a	and the relevant enth			
	State	e symbols should be u	used.			
					KCl(s)	
						[2]
(b)		the data in Table 1.1 . Show your working.	to calculate the en	thalpy change o	of hydration of mag	gnesium ions,

 $\Delta H_{\text{hyd}}^{\text{e}}$ of magnesium ions, $Mg^{2+} = \dots kJ \, \text{mol}^{-1}$ [2]

(c)		plain the reasons why the lattice energy of ${\rm MgC} l_2$ is more exothermic than the lattice energy KC l .
		[2]
(d)	Def	fine the following terms.
	(i)	enthalpy change of atomisation
		[1]
	(ii)	first electron affinity
		[1]
(e)	(i)	Explain what is meant by entropy, S.
		[1]
	(ii)	Potassium chloride is very soluble in water at 20 °C.
		Explain the solubility of potassium chloride by reference to change in entropy, ΔS .
		[1]
	(iii)	Use the Gibbs equation and your answer to (e)(ii) to predict whether potassium chloride is more soluble in water at 20 °C or at 80 °C. Explain your answer.
		[1]
		[Total: 11]

4

2	(a) (i)	Explain what is meant by the following terms:
		homogeneous catalyst
		heterogeneous catalyst
		[1]
	(ii)	lodide ions react with peroxydisulfate ions.
		$2I^{-} + S_{2}O_{8}^{2-} \rightarrow I_{2} + 2SO_{4}^{2-}$
		This reaction is slow, but it is catalysed by Fe ²⁺ ions.
		Write two equations to explain how this reaction is catalysed by Fe ²⁺ ions.
		1
		2[2]
	(iii)	Suggest why the alternative route in the presence of Fe ²⁺ ions has a lower activation energy than the route in the absence of a catalyst.
		[1]
	(b) Nit	rogen monoxide reacts with oxygen.
		$2NO(g) + O_2(g) \rightarrow 2NO_2(g)$
		is reaction is second order with respect to nitrogen monoxide and first order with respect to ygen.
	Un	der certain conditions the value of the rate constant, k , is $8.60 \times 10^6 \mathrm{mol^{-2}}\mathrm{dm^6s^{-1}}$.
	(i)	Construct the rate equation for this reaction.
		rate = [1]
	(ii)	Calculate the initial rate of the reaction under these conditions when the initial concentration
	(11)	of nitrogen monoxide is $7.20 \times 10^{-4} \text{mol dm}^{-3}$ and the initial concentration of oxygen is $1.90 \times 10^{-3} \text{mol dm}^{-3}$.
		rate of rocation and dece-3 1 F43
		rate of reaction = mol dm ⁻³ s ⁻¹ [1]

(c)	The drug	cisplatin,	$Pt(NH_3)_2Cl_2$,	hydrolyses	in wate
\ -/			3/2 - 21	,	

$$Pt(NH_3)_2Cl_2 + H_2O \rightarrow [Pt(NH_3)_2Cl(H_2O)]^+ + Cl^-$$

The rate equation is shown.

rate =
$$k[Pt(NH_3)_2Cl_2]$$

The value of k is 2.50×10^{-5} s⁻¹ under certain conditions.

(i)	This reaction has a constant half-life.
	Explain why this is the case

Explain why this is the case.	
[[1]
	_

(ii) Use the information in this question to show that the half-life of this reaction is $2.77 \times 10^4 \, s$.

[1]

(iii) 8.00×10^{-6} moles of Pt(NH₃)₂C l_2 are added to $100\,\mathrm{cm^3}$ of water. Calculate the time taken for the concentration of Pt(NH₃)₂C l_2 to fall to $2.50 \times 10^{-6}\,\mathrm{mol\,dm^{-3}}$.

time taken = s [2]

[Total: 10]

3 Data should be selected from Table 3.1 in order to answer some parts of this question.

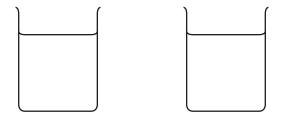
Table 3.1

electrode reaction	E [⊕] /V
Cl₂ + 2e⁻ ⇌ 2Cl⁻	+1.36
$2\text{HOC}l + 2\text{H}^+ + 2\text{e}^- \rightleftharpoons \text{C}l_2 + 2\text{H}_2\text{O}$	+1.64
$ClO^- + H_2O + 2e^- \rightleftharpoons Cl^- + 2OH^-$	+0.89
Sn⁴+ + 2e⁻ ⇌ Sn²+	+0.15
Sn²+ + 2e⁻ ⇌ Sn	-0.14
V²+ + 2e⁻ ⇌ V	-1.20
V³+ + e⁻ ← V²+	-0.26
$VO^{2+} + 2H^{+} + e^{-} \rightleftharpoons V^{3+} + H_{2}O$	+0.34
$VO_2^+ + 2H^+ + e^- \rightleftharpoons VO^{2+} + H_2O$	+1.00

(a) Standard electrode potentials are measured under standard conditions.

(i)	Describe the standard conditions used in the Sn ⁴⁺ /Sn ²⁺ half-cell.			
	[1]			

(ii) Complete the diagram below to show how E° (Sn⁴⁺/Sn²⁺) can be measured experimentally. Your diagram should be fully labelled to identify all apparatus and substances.



[3]

	(iii)	Equal volumes of 1.0 mol dm ⁻³ Sn ²⁺ (aq) and 1.0 mol dm ⁻³ C l^- (aq) are mixed.	
		Use relevant E^{e} values to explain whether a reaction occurs between these two ions.	
		[2]	
	(iv)	Equal volumes of 1.0 mol dm ⁻³ of Sn ²⁺ (aq) and acidified 1.0 mol dm ⁻³ VO ²⁺ (aq) are mixed.	
		Write an equation for the reaction that takes place in the resulting mixture.	
		[2]	
(b)	A s	olution of $SnCl_2(aq)$ is electrolysed for a measured time using a steady current.	
A mass of 2.95g of tin metal is produced at the cathode.			
	O ₃ (I) is electrolysed for the same time by the same current.		
		culate the mass of aluminium metal produced at the cathode. Give your answer to three nificant figures. Show your working.	
		mass of aluminium metal = g [2]	
		[Total: 10]	

[1]

4	(a)	sol ^v bet	A sample of butanoic acid, $CH_3(CH_2)_2COOH$, is shaken with a mixture of two immiscible solvents, ethoxyethane and water. The solvents form two layers. The butanoic acid is distributed between the two layers, its concentration in ethoxyethane being higher than its concentration in water.			
		(i)	State what is meant by partition coefficient.			
			[1]			
		(ii)	The partition coefficient, K_{pc} , for butanoic acid between ethoxyethane and water is 3.50.			
			A solution of 2.00 g of butanoic acid in 100 cm³ ethoxyethane is added to water. This mixture is left until there is no further change in the concentration of butanoic acid in either solvent. The mass of butanoic acid dissolved in the ethoxyethane layer is now 1.62 g.			
			Calculate the volume of water used.			
			volume of water used =cm³ [2]			
	(b)	An	aqueous solution of butanoic acid can be used to make a buffer solution.			
		(i)	Define buffer solution.			
			[1]			
		(ii)	Suggest one organic compound, and one inorganic compound, that can be added to two different samples of aqueous butanoic acid to produce buffer solutions.			
			organic compound			
			inorganic compound			

(c)	The solubility of a	aluminium hydroxide	$Al(OH)_3$, in	n water is 2.47	\times 10 ⁻⁹ mol dm ⁻³
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(i) Give the expression for the solubility product, K_{sn} , of aluminium	. nyaroxiae.
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$$K_{sp} =$$

[1]

(ii) Calculate the numerical value of the $K_{\rm sp}$ of aluminium hydroxide. Include the units of $K_{\rm sp}$ in your answer.

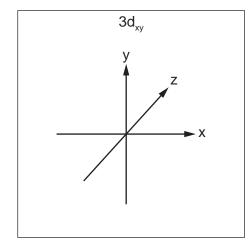
 K_{sp} =

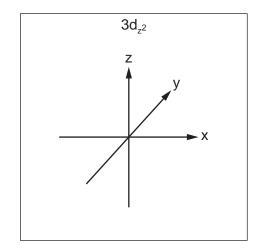
units =[3]

[Total: 9]

- **5** A transition element is a d-block element which forms one or more stable ions with incomplete d-orbitals.
 - (a) Two of the 3d orbitals are the $3d_{xy}$ orbital and the $3d_{z^2}$ orbital.

Sketch the shapes of these two orbitals.





[1]

(b) The Ni²⁺ ion forms many different complexes. A solution containing the $[Ni(H_2O)_6]^{2+}$ complex ion is green. When an excess of 1,2-diaminoethane, en, $H_2NCH_2CH_2NH_2$, is added, the colour of the solution changes to blue. This is due to the formation of the $[Ni(en)_3]^{2+}$ complex ion.

1	i۱	Explain wh	v the two	solutions a	re coloured	and why th	ne colours are	different
v	'/	Explain will	iy tilo tivo	oolationo a	no ocioaroa,	aria writy ti	io obloaro aro	anio on c

	ΓΛ

	(ii)	The $[Ni(en)_3]^{2+}$	complex can exist	as a mixture	of two stereois	somers.			
		Complete the three-dimensional diagram to show one of the stereoisomers.							
		Each en ligand	can be represent	ed using N	N .				
				1					
				 Ni					
						I	[1]		
	(iii)	Name the geor shown by [Ni(e	metry of the comp $n)_3]^{2+}$.	lex ion drawn	in (b)(ii) and	the type of ste	ereoisomerism		
		geometry							
		stereoisomeris	m shown						
							[1]		
(c)	Iror	n(II) carbonate, F	FeCO ₃ , and nickel	(II) carbonate	, NiCO ₃ , both	decompose w	hen heated.		
	FeC	CO ₃ decomposes	s at a lower tempe	erature than N	iCO ₃ .				
	Sug	gest a possible	reason for this dif	ference. Expla	ain your answe	er.			
							[2]		

(d)	A is a pale green salt containing Fe2+ ions. A sample of 2.62 g of A is dissolved in water and the
	solution is made up to exactly 100 cm ³ with water. 25.0 cm ³ samples of this solution are placed
	in conical flasks and titrated against 0.0100 mol dm ⁻³ acidified potassium manganate(VII).

The equation for the only reaction that occurs is shown.

$$5 Fe^{2+} \ + \ MnO_4^{-} \ + \ 8H^+ \ \rightarrow \ 5 Fe^{3+} \ + \ Mn^{2+} \ + \ 4H_2O$$

The average titre value is 35.0 cm³ of 0.0100 mol dm⁻³ acidified potassium manganate(VII).

- (ii) Calculate the percentage by mass of iron in A.

[Total: 12]

Question 6 starts on the next page.

(a) (i)	Write an equation for the reaction bet	tween $[Cu(H_2O)_6]^{2+}$ ions and NaOH(aq).
(ii)	Write an equation for the reaction bet	tween $[Cu(H_2O)_6]^{2+}$ ions and an excess of conc.
		excess of ammonia is added to a sol NH_3 ₄ $(H_2O)_2$ ²⁺ complex ions is formed	lution of $[Cu(H_2O)_6]^{2+}$ a deep blue solution contad.
(i)	There are two possible stereoisomers	s with the formula $[Cu(NH_3)_4(H_2O)_2]^{2+}$.
		Complete the diagrams to show the t	wo stereoisomers in the boxes below.
		isomer 1	isomer 2
		Cu	Cu
(ii)	Use your answer in (b)(i) to deduce v	whether each of these isomers is polar or non-p
		polarity of isomer 1	
		polarity of isomer 2	

	(ii) Compare the stabilities of the $[Cu(H_2O)_6]^{2+}$ and $[Cu(NH_3)_4(H_2O)_2]^{2+}$ complex ions. Ex your answer.					
		[1]				
(iii)	Write an expression for the stability constant, K_{stab} , of the $[\text{Cu}(\text{NH}_3)_4(\text{H}_2\text{O})_2]^{2+}$ complex ion. State the units of the stability constant.				
		$K_{\text{stab}} =$				
		units =[2]				
(iv)	In a particular solution the concentration of the $[Cu(NH_3)_4(H_2O)_2]^{2+}$ complex ion is 0.0074 mol dm ⁻³ and the concentration of NH_3 is 0.57 mol dm ⁻³ .				
		Use your expression in (c)(iii) and the K_{stab} value of 1.40×10^{13} to calculate the concentration of the $[\text{Cu}(\text{H}_2\text{O})_6]^{2^+}$ complex ion in this solution.				
		concentration of $[Cu(H_2O)_6]^{2+} =$ moldm ⁻³ [1]				
(d)	Phe	enanthroline, $C_{12}H_8N_2$, and ethanedioate ions, $C_2O_4^{2-}$, are bidentate ligands.				
		henium(III) ions, Ru³+, form an octahedral complex with phenanthroline and chloride ions. complex ion contains two phenanthroline molecules.				
	Iron	(III) ions, Fe ³⁺ , form an octahedral complex with ethanedioate ions only.				
	Dec	duce the formula and charge of each of these complex ions.				
	Ru³	† complex				
	Fe ^{3.}	† complex				
		[2]				
		[Total: 11]				

7 (a) The structure of compound P is shown in Fig. 7.1.

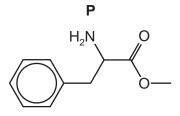


Fig. 7.1

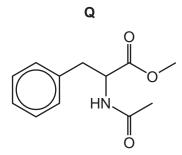
(i) P is optically active.

Use an asterisk (*) to identify all chiral carbon atoms on the structure of **P** in Fig. 7.1. [1]

(ii) Plane polarised light is passed through a pure sample of one enantiomer of **P**. This is then repeated with a pure sample of the other enantiomer of **P**.

the results.	suits of these two	experiments, s	tating the simila	rities and differe	nces of
					[2]

(b) ${\bf P}$ can be used to make compound ${\bf Q}$ in a single step reaction.



(i) Give the structural formula of the compound that is added to **P** to make **Q** and give the formula of the other product of this reaction.

compound added to P	
other product	
·	[1]

(ii)	When an ester is treated with LiAlH4 in dry ether the ester linkage is cleaved by the
	addition of four hydrogen atoms and two alcohols are produced.

Draw the structures of the compounds formed when ${\bf Q}$ is treated with an ${\bf excess}$ of ${\rm LiA} l{\rm H_4}$ in dry ether.

		[3]	
(iii)	·	ties of compound P , compound Q and phenylamine.	
	least basic	most basic	
	Explain your answer.		
		[31	

(c) P can be used to make compound R in a two-step reaction, shown in Fig. 7.2.

Fig. 7.2

(i)	Identify the reagents and conditions used for the two steps of the reaction.	
	step 1	
	step 2	
		[2

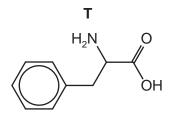
(ii) Complete Table 7.1 by drawing the structures of the organic products formed when ${\bf R}$ is treated separately with the reagents given.

Table 7.1

reagent	product
HNO₂(aq) at 4°C	
an excess of Br ₂ (aq) at room temperature	

[2]

(d) ${\bf P}$ can be used to produce compound ${\bf T}$.



(i)	In aqueous solution, T has a property called an isoelectric point.
	Explain what is meant by isoelectric point.
	[1]
(ii)	T can polymerise under suitable conditions. No other monomer is involved in this reaction.
	Draw a section of the polymer chain formed that includes three T monomers. Identify the repeat unit on your diagram.

[2]

[Total: 17]

[1]

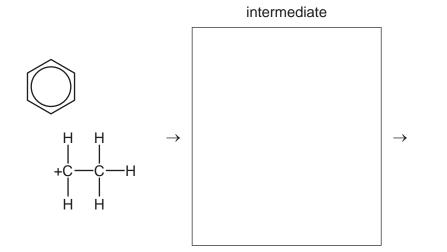
- **8** Benzene, C₆H₆, is an aromatic molecule.

 - **(b)** Benzene reacts with chloroethane in the presence of a catalyst. The reaction mechanism is called electrophilic substitution.
 - (i) The first step in the reaction is the generation of the ⁺CH₂CH₃ electrophile.

Write an equation for the reaction that generates this electrophile.

.....[1]

(ii) Describe the mechanism for the reaction between benzene and the ⁺CH₂CH₃ electrophile. Include all relevant curly arrows and charges.



[3]

products

(c)		orobenzene and chloroethane have different reactivities in nucleophilic substitution ctions.
	(i)	Identify a suitable reagent to illustrate this difference in reactivity.
		The reagent chosen should give visibly different results with chlorobenzene and chloroethane.
		[1]
	(ii)	Write equations to describe any reactions that occur.
		[1]
(iii)	Explain the difference in the reactivities of chlorobenzene and chloroethane in nucleophilic substitution reactions.
		[1]
		[Total: 8]

[1]

9	(a)	Gas-liquid	chromatography	involves a	stationary	phase	and a	mobile	phase
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(i)	Name, or describe in detail, a suitable substance that could be used for each phase.
	stationary
	mobile

A mixture of three organic compounds is separated by gas-liquid chromatography. The chromatogram obtained is shown in Fig. 9.1. The amount of each substance is proportional to the area under its peak.

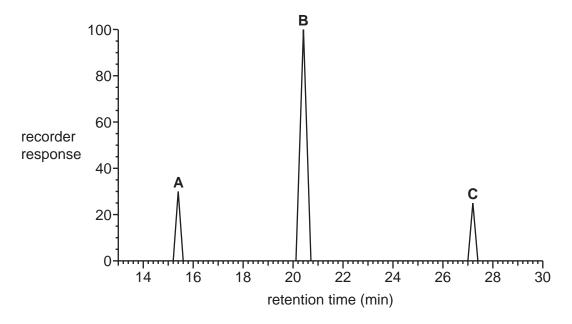


Fig. 9.1

(ii)	Explain the meaning of retention time.
	[1

(iii) Calculate the percentage of **B** in the mixture. Show your working.

percentage of **B** = % [2]

(b) Complete Table 9.1 to give the number of peaks in the carbon-13 NMR spectrum of each of the five isomers of $C_5H_{10}O_2$ that has an ester group.

Table 9.1

structural formula	number of peaks
CH ₃ CH ₂ CH ₂ CO ₂ CH ₃	
CH ₃ CH ₂ CO ₂ CH ₂ CH ₃	
CH ₃ CO ₂ CH ₂ CH ₂ CH ₃	
(CH ₃) ₂ CHCO ₂ CH ₃	
CH ₃ CO ₂ CH(CH ₃) ₂	

[2]

(c)	State the number of peaks that would be seen in the proton (1 H) NMR spectrum of methyl butanoate, $CH_{3}CH_{2}CO_{2}CH_{3}$. Name all the splitting patterns seen in this spectrum.
	number of peaks
	splitting patterns[2]

(d) **D** and **E** are both esters with the molecular formula $C_5H_{10}O_2$. Their proton (¹H) NMR spectra are shown in Fig. 9.2 and Fig. 9.3.

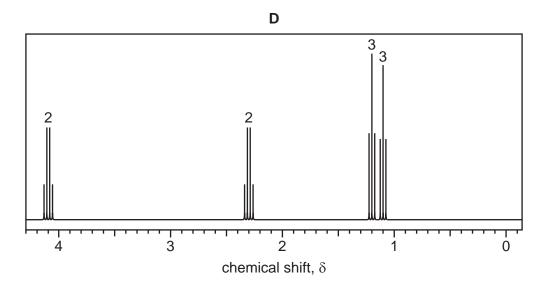


Fig. 9.2

Е

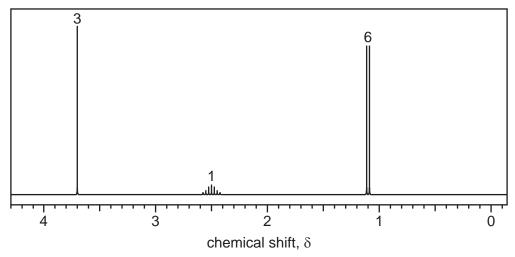


Fig. 9.3

Table 9.2

environment of proton	example	typical chemical shift range, δ/ppm
alkane	-CH ₃ , -CH ₂ -, >CH-	0.9–1.7
alkyl next to C=O	CH ₃ -C=O, -CH ₂ -C=O, >CH-C=O	2.2–3.0
alkyl next to aromatic ring	CH ₃ -Ar, -CH ₂ -Ar, >CH-Ar	2.3–3.0
alkyl next to electronegative atom	CH ₃ -O, -CH ₂ -O, -CH ₂ -C <i>l</i>	3.2–4.0
attached to alkene	=CHR	4.5–6.0

(i)	Deduce the structures of the two esters D and E and draw their displayed formulae in the
	boxes below.

	D C ₅ H ₁₀ O ₂		E C ₅ H ₁₀ O ₂	
				[2]
(ii)	The spectrum of D includes a quarter	t at a	8 4.1.	
	Identify the protons responsible for this quartet on your structure in (i) by labelling these protons with the letter F .			
	Explain why this peak is split into a q	luart	et.	
				[1]
(iii)	The spectrum of ${\bf E}$ has a doublet at δ	5 1.1		
	Identify the protons responsible for the protons with the letter G .	his d	doublet on your structure in (i) by labelling th	iese
	Explain why this peak has a chemica	al sh	ift of 1.1.	

[Total: 12]

[1]

26

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Important values, constants and standards

molar gas constant	$R = 8.31 \mathrm{J}\mathrm{K}^{-1}\mathrm{mol}^{-1}$
Faraday constant	$F = 9.65 \times 10^4 \mathrm{C}\mathrm{mol}^{-1}$
Avogadro constant	$L = 6.022 \times 10^{23} \mathrm{mol^{-1}}$
electronic charge	$e = -1.60 \times 10^{-19} \mathrm{C}$
molar volume of gas	$V_{\rm m} = 22.4 {\rm dm^3 mol^{-1}}$ at s.t.p. (101 kPa and 273 K) $V_{\rm m} = 24.0 {\rm dm^3 mol^{-1}}$ at room conditions
ionic product of water	$K_{\rm w} = 1.00 \times 10^{-14} \rm mol^2 dm^{-6} (at 298 \rm K (25 {}^{\circ} C))$
specific heat capacity of water	$c = 4.18 \mathrm{kJ kg^{-1} K^{-1}} (4.18 \mathrm{J g^{-1} K^{-1}})$

The Periodic Table of Elements

1 2 2	- 1																						
The color of the		18	2	He	helium 4.0	10	Ne	neon 20.2	18	Ar	argon 39.9	36	궃	krypton 83.8	54	Xe	xenon 131.3	98	Rn	radon	118	Og	oganesso
1		17				6	Щ	fluorine 19.0	17	Cl	chlorine 35.5	35	Ŗ	bromine 79.9	53	Ι	iodine 126.9	82	Ą	astatine -	117	<u>S</u>	tennessine
14 Handburner Handburner		16				80	0	oxygen 16.0	16	S	sulfur 32.1	34	Se	selenium 79.0	52	Те	tellurium 127.6	84	Ъо	polonium –	116	_	livermorium -
13 H H H H H H H H H		15				7	z	nitrogen 14.0	15	۵	phosphorus 31.0	33	As	arsenic 74.9	51	Sp	antimony 121.8	83	<u>.</u>	bismuth 209.0	115	Mc	moscovium -
2 1 1 1 1 1 1 1 1 1		14				9	O	carbon 12.0	14	S	silicon 28.1	32	Ge	germanium 72.6	20	Sn	tin 118.7	82	Pb	lead 207.2	114	ŁΙ	flerovium
1		13				2	В	boron 10.8	13	ΝI	aluminium 27.0	31	Ga	gallium 69.7	49	In	indium 114.8	18	11	thallium 204.4	113	R	nihonium
A											12	30	Zu	zinc 65.4	48	පි	cadmium 112.4	80	Нg	mercury 200.6	112	ပ်	copernicium
A											1	59	Cn	copper 63.5	47	Ag	silver 107.9	62	Au	gold 197.0	111	Rg	roentgenium
1	dno										10	28	Z	nickel 58.7	46	Pd	palladium 106.4	78	₫	platinum 195.1	110	Ds	darmstadtium -
A	Gro										0	27	ဝိ	cobalt 58.9	45	R	rhodium 102.9	77	ľ	iridium 192.2	109	Ĭ	meitnerium -
A			-	I	hydrogen 1.0						∞	26	Pe	iron 55.8	44	Ru	ruthenium 101.1	92	SO	osmium 190.2	108	¥	hassium -
A											7	25	Mn	manganese 54.9	43	ည	technetium -	75	Re	rhenium 186.2	107	Bh	bohrium –
A							pol	ass			9	24	ပ်	chromium 52.0	42	Mo	molybdenum 95.9	74	≥	tungsten 183.8	106	Sg	seaborgium -
A					Key	atomic number	mic sym	name ative atomic ma			2	23	>	vanadium 50.9	41	g	niobium 92.9	73	<u>n</u>	tantalum 180.9	105	o D	dubnium -
2 Be benyilium 9.0							ato	rela			4	22	F	titanium 47.9	40	Zr	zirconium 91.2	72	Ξ	hafnium 178.5	104	Ŗ	rutherfordium -
									-		က	21	လွ	scandium 45.0	39	>	yttrium 88.9	57-71	lanthanoids		89–103	actinoids	
1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1		2				4	Be	beryllium 9.0	12	Mg	magnesium 24.3	20	Ca	calcium 40.1	38	လွ	strontium 87.6	56	Ba	barium 137.3	88	Ra	radium
		1				က	:-	lithium 6.9	11	Na	sodium 23.0	19	¥	potassium 39.1	37	Rb	rubidium 85.5	55	S	caesium 132.9	87	ŗ	francium -

						_
71	2	lutetium 175.0	103	ت	lawrencium	ı
		ytterbium 173.1			_	I
69	T	thulium 168.9	101	Md	mendelevium	ı
89	Ē	erbium 167.3	100	Fm	ferminm	ı
29	웃	holmium 164.9	66	Es	einsteinium	-
99	۵	dysprosium 162.5	86	ŭ	californium	1
99	Р	terbium 158.9	26	益	berkelium	1
64	В	gadolinium 157.3	96	Cm	curium	ı
63	En	europium 152.0	95	Am	americium	ı
62	Sm	samarium 150.4	94	Pu	plutonium	I
19	Pm	promethium —	93	g	neptunium	ı
09	PN	neodymium 144.4	92	\supset	uranium	238.0
59	Ą	praseodymium 140.9	91	Ра	protactinium	231.0
58	Ce	cerium 140.1	06	Th	thorium	232.0
22	Гa	lanthanum 138.9	89	Ac	actinium	1

lanthanoids actinoids

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