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School: .....

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**P525/1**  
**CHEMISTRY**  
**Paper 1**  
**July/August 2015**  
**2 ¾ hours**



## **WAKISSHA JOINT MOCK EXAMINATIONS**

**Uganda Advanced Certificate of Education**

**CHEMISTRY**

**Paper 1**

**2 hours 45 minutes**

### **Instructions to Candidates**

- Attempt **all** questions in section **A** and any **six** questions from section **B**.
- All questions are to be answered in the spaces provided.
- A Periodic Table with relevant atomic masses is supplied at the end of the paper.
- Mathematical tables (3 figures) and non-programmable silent scientific calculators may be used.
- Illustrate your answers with equations where applicable.

For Examiner's Use Only																	
1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	Total

## SECTION A (46 marks)

Answer **all** questions in this section.

1. (a) X, Y and Z are structural isomers of  $C_4H_{10}O$ . The isomers react with sodium metal liberating hydrogen gas.

(i) Name the functional group in the isomers of  $C_4H_{10}O$ . (1/2 mark)

(ii) Write an equation for the general reaction that occur between  $C_4H_{10}O$  and sodium metal. (1 mark)

- (b) Isomer x reacts with hot concentrated sulphuric acid but it gives no observable change on addition of warm acidified potassium dichromate solution.

(i) identify x. (1 mark)

(ii) Suggest a suitable mechanism for the reaction between x and hot concentrated sulphuric acid. (2 marks)

(iii) State the molecularity of the reaction in b(ii) above. Give a reason for your answer. (1 1/2 mark)

2. (a) Copper like other transition elements forms compounds in oxidation states: +1 and +2.

Write the electronic configuration of:

(i) Copper (I) ions. (1/2 mark)

(c) State any two practical applications of radioactive isotopes. (1 mark)

4. Name the reagent(s) that can be used to distinguish between each of the following pairs of compounds. State the observation(s) when each compound is separately treated with the reagent(s).

(a)  $\text{CH}_3\text{COONH}_4(\text{aq})$  and  $\text{NH}_4\text{F}(\text{aq})$  (1½ marks each)

Reagent(s)

Observations.

(b)  $\begin{array}{c} \text{CH}_3\text{CHCH}_2\text{NH}_2 \\ | \\ \text{OH} \end{array}$  and  $\begin{array}{c} \text{CH}_3\text{CHNHCH}_3 \\ | \\ \text{OH} \end{array}$

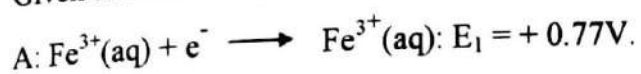
Reagent(s)

Observations.

(c) NiO and FeO.  
Reagent(s)

Observations.

5. Given the following reduction reactions and their corresponding electrode potentials.



The two half cell reactions; A and B form a feasible cell reaction with reaction A at the cathode and  $E^\theta_{\text{cell}} = +0.23\text{V}$ .

(a) (i) Write the equation for the reaction that occurs at the anode. (1 mark)

(ii) Calculate the value of  $E_2$ . (1mark)

(b) Draw a well labelled diagram to show how the  $E^\theta_{\text{cell}} = +0.23\text{V}$  can be measured practically from the half-cell reactions. (3 marks)

Turn Over  
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- (c) State two conditions under which the cell in (b) above is set-up. (1 mark)

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6. Iron and lead form mixed oxides  $\text{Fe}_3\text{O}_4$  and  $\text{Pb}_3\text{O}_4$  respectively.

(a) Write the;

- (i) IUPAC name of the oxide of lead ( $\text{Pb}_3\text{O}_4$ ) that represents the ratio in which the oxides mix.

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- (ii) Equation to show that  $\text{Fe}_3\text{O}_4$  is a mixed oxide. (1 mark)

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(b)  $\text{Pb}_3\text{O}_4$  oxidises hot concentrated hydrochloric acid to a pale green gas and itself reduced to a colourless solution which forms a white precipitate on cooling.

(i) Identify the;

Pale green gas. ( $\frac{1}{2}$  mark)

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White precipitate. ( $\frac{1}{2}$  mark)

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- (ii) Write the equation for the reaction between  $\text{Pb}_3\text{O}_4$  and hot concentrated hydrochloric acid. (1 mark)

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- (c) State any two other chemical properties to show the similarity between iron and lead. (1 mark)

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7. At  $100^\circ\text{C}$ , the dissociation pressures of carbon dioxide in equilibrium with the carbonates of Lithium, sodium and potassium are given in the table below.

$\text{Li}_2\text{CO}_3$	$\text{Na}_2\text{CO}_3$	$\text{K}_2\text{CO}_3$
90mmHg	19mmHg	8mmHg

- (a) Which one of these carbonates is thermally most stable at  $100^\circ\text{C}$ ?

Give a reason for

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(b) (i) Identify the

(ii) Write an equation for the carbon

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(iii) Explain why the reaction in b(i) above

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(c) State any two conditions for the reaction in b(ii) above

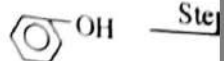
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8. The figure below shows the reaction of acetic acid and water. Step



(a) Name the compounds:

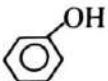
(i) Q \_\_\_\_\_ (1/2 mark)

(ii) W \_\_\_\_\_ (1/2 mark)

(b) State the condition(s) necessary for the formation of:

(i) Q from  $\text{CH}_3\text{COOH}$  (1/2 mark)

\_\_\_\_\_  
\_\_\_\_\_

(ii) W from 

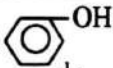
\_\_\_\_\_  
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(c) (i) Name the type of reaction that occurs in step III. (1/2 mark)

\_\_\_\_\_  
\_\_\_\_\_

(ii) Explain why W reacts with Q to form Z under the conditions in step III. (1 1/2 mark)

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\_\_\_\_\_  
\_\_\_\_\_  
\_\_\_\_\_

(d) (i) State what would be observed when compound Q is reacted with  (1/2 mark)

\_\_\_\_\_  
\_\_\_\_\_

(ii) Write an equation for the reaction in d(i) above. (1 mark)

\_\_\_\_\_  
\_\_\_\_\_  
\_\_\_\_\_  
\_\_\_\_\_

9. Pure water boils at  $100^\circ\text{C}$  at  $1013.25\text{KNm}^{-2}$  pressure. However aqueous solutions of nitric acid and ethanol boil at temperatures shown below.

Solution	Composition	Boiling point ( $^\circ\text{C}$ )
Aqueous nitric acid	68.0% $\text{HNO}_3$	120.5
Aqueous ethanol	95.6% $\text{CH}_3\text{CH}_2\text{OH}$	78.2

- [illegible]

- (i) Distimate.

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- (i) Intermolecular forces in water.

- \_\_\_\_\_

- \_\_\_\_\_
- \_\_\_\_\_



## SECTION B

Answer only **six** questions from this section.

Any additional questions answer **will not be marked**.

10. In an experiment to determine the formula of an alkylhalide,  $\text{RCH}_2\text{I}$ , the alkylhalide is refluxed with concentrated aqueous sodium hydroxide followed by acidified silver nitrate. The mixture is filtered and the mass of dry residue measured.

- (a) (i) Write equation(s) for the reaction(s) that take place in the experiment. (2 marks)

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- (ii) Why is the silver nitrate solution acidified? (1 mark)

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- (iii) Name the type of reaction that occurs on reflux. ( $\frac{1}{2}$  mark)

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- (b) 3.4g an alkylhalide  $\text{RCH}_2\text{I}$  were refluxed with excess 2M sodium hydroxide solution. To the resultant mixture, excess acidified silver nitrate was added and the product filtered. The mass of the dry residue was found to be 4.70g.

- (i) Calculate the formula mass of R. (2 marks)

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- (ii) Deduce the structural formula of  $\text{RCH}_2\text{I}$  from your answer in b(i) above.

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- (c) Using equations, show any alkylhalide  $\text{RCH}_2\text{I}$  can be converted to  $\text{RI}$ . ( $2\frac{1}{2}$  marks)

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11. Explain each of the following observations. Illustrate your answer with equations of reactions (where applicable).

- (a) When potassium methanoate solution is added to copper (II) sulphate solution, a blue precipitate is formed. (3 marks)

- (b) When boiled with Fehling's solution, aliphatic alkanals form a reddish brown precipitate. (3 marks)

**Turn Over**

- (c) Helium can be separated from Argon by diffusion however a mixture of ethane and nitrogen cannot be separated by diffusion. (3 marks)

12. (a) 3.0g of solid calcium iodate(v) were vigorously shaken with water in a stoppered bottle and the mixture allowed to stand at  $25^{\circ}\text{C}$  for about 30 minutes. The resultant mixture was filtered to obtain exactly  $200\text{cm}^3$  of the filtrate. Given that calcium iodate (v) is sparingly soluble in water;

(i) Name the filtrate. ( $\frac{1}{2}$ mark)

(ii) Write an equation for the reaction that occurs at  $25^{\circ}\text{C}$  at the end of the 30 minutes. (1marks)

- (b)  $20\text{cm}^3$  of the filtrate were pipetted into excess acidified potassium iodide solution and the mixture titrated with exactly  $6.75\text{cm}^3$  of 0.1M sodium thiosulphate solution in the presence of starch indicator.

Calculate the percentage by mass of calcium iodate(V) in  $200\text{cm}^3$  of the filtrate. ( $4\frac{1}{2}$  marks)

(c) The experiment in (a) above was repeated with potassium iodate solution instead of water at 25°C.

(i) State whether the percentage of calcium iodate(v) in the filtrate would be less, equal to or greater than the calculated value in (b) above. (1 mark)

(ii) Explain your answer in c(i) above. (2 marks)

13. (a)  $F_2$ ,  $Cl_2$ ,  $Br_2$  and  $I_2$  are diatomic molecules of group VII elements of the periodic table. Briefly describe how the elements react with;

(i) Potassium bromide solution, (2 marks)

(ii) Hydrogen sulphide. (2 marks)

- (b) The hydride of the elements in (a) above boil at different temperatures shown in the table below.

HF	HCL	HBR.	HI
+20 <sup>0</sup> C	-85 <sup>0</sup> C	-67 <sup>0</sup> C	-35 <sup>0</sup> C

State and explain the trend in the boiling points of the hydrides.

(3 marks)

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- (d) (i) State the order in reducing strength of the hydrides.

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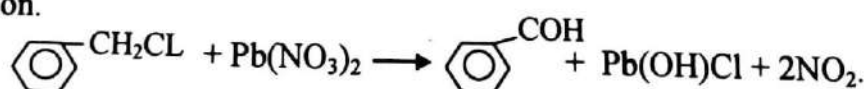
- (ii) Write a general equation for the reaction between aqueous solutions of halogen acids (hydrides) with solid sodium carbonate.

(1 mark)

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14. (a) In the hydrolysis – oxidation reaction, benzyl chloride is boiled under air-reflux with lead (II) nitrate solution for about 3 hours to form Benzaldehyde according to the equation.



Benzaldehyde is obtained by steam distillation from the reaction mixture and extracted by ether from the distillate. State the reason(s) why:

- (i) the basic chloride of lead  $\text{Pb}(\text{OH})\text{Cl}$  is not found in the steam distillate?

( $\frac{1}{2}$  mark)

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- (ii) ether is used to extract benzaldehyde from the distillate.

(1 mark)

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- (b) Benzaldehyde is separately treated with cold concentrated potassium hydroxide followed by dilute hydrochloric acid and hydroxylamine in acidic medium.

17. Write an equation for the reaction between benzoaldehyde and potassium hydroxide followed by dilute hydrochloric acid. (1 mark)

18. Suggest a suitable mechanism for the reaction between benzoaldehyde and hydrocyanic acid in acidic medium. (3 marks)

19. Write equations which show how benzoaldehyde can be:  
(a) Prepared from methyl benzoate. (1½ marks)

- (b) Converted to butyraldehyde. (2 marks)

20. (a) Define the following terms as applied to ionic equilibria. (1 mark)

(i) Ionisation

- (ii) Salt hydrolysis. (1 mark)

Turn Over  
15

(b) Write an equation for the;

(1 mark)

(i) Ionisation of ammonia in water.

(ii) hydrolysis of ammonium sulphate.

(1 mark)

(c) You are provided with solid anhydrous ammonium sulphate, water and 250ml volumetric flask. Briefly outline how a solution of ammonium sulphate of pH = 5.20 can be prepared in the laboratory. ( $K_b$  for ammonia =  $1.78 \times 10^{-5} \text{ mol dm}^{-3}$  and ionic product of water at  $25^\circ\text{C}$ ,  $K_w = 1.0 \times 10^{-14} \text{ mol}^2 \text{ dm}^{-6}$ ) (5 marks)

(5 marks)

16. (a) Powdered chlomite ( $\text{FeCr}_2\text{O}_4$ ) is roasted with anhydrous sodium carbonate in air and calcium oxide.



Hot water is added to the products of the reaction and a yellow solid crystallises out.

(i) Name the yellow solid.

( $\frac{1}{2}$  mark)

- (ii) Draw the structure and name the shape adopted by the anion in the ionic crystal lattice of the yellow solid in a(i) above. (1 mark)

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- (b) The yellow solid dissolves in excess dilute sulphuric acid to form an orange solution which is a strong oxidizing agent. State what would be observed and write an equation for the REDOX reaction (where applicable) that occurs when the orange solution is reacted with;

- (i) Concentrated sodium hydroxide solution, (1½ marks)

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- (ii) A saturated solution of potassium Sulphite. (2 marks)

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- (iii) Excess warm ethanol. (2 marks)

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- (c) Other than acting as an oxidizing agent in volumetric analysis, the aqueous solution of the yellow solid in a(i) above is used in precipitation titration involving silver nitrate solution.

- (i) State the role of the yellow aqueous solution in precipitation titration. (½ mark)

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- (ii) Write an equation for the reaction that occurs and state the observation at the end point. (1 mark)

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- 17 (a) (i) Define the term vapour pressure. (1 mark)

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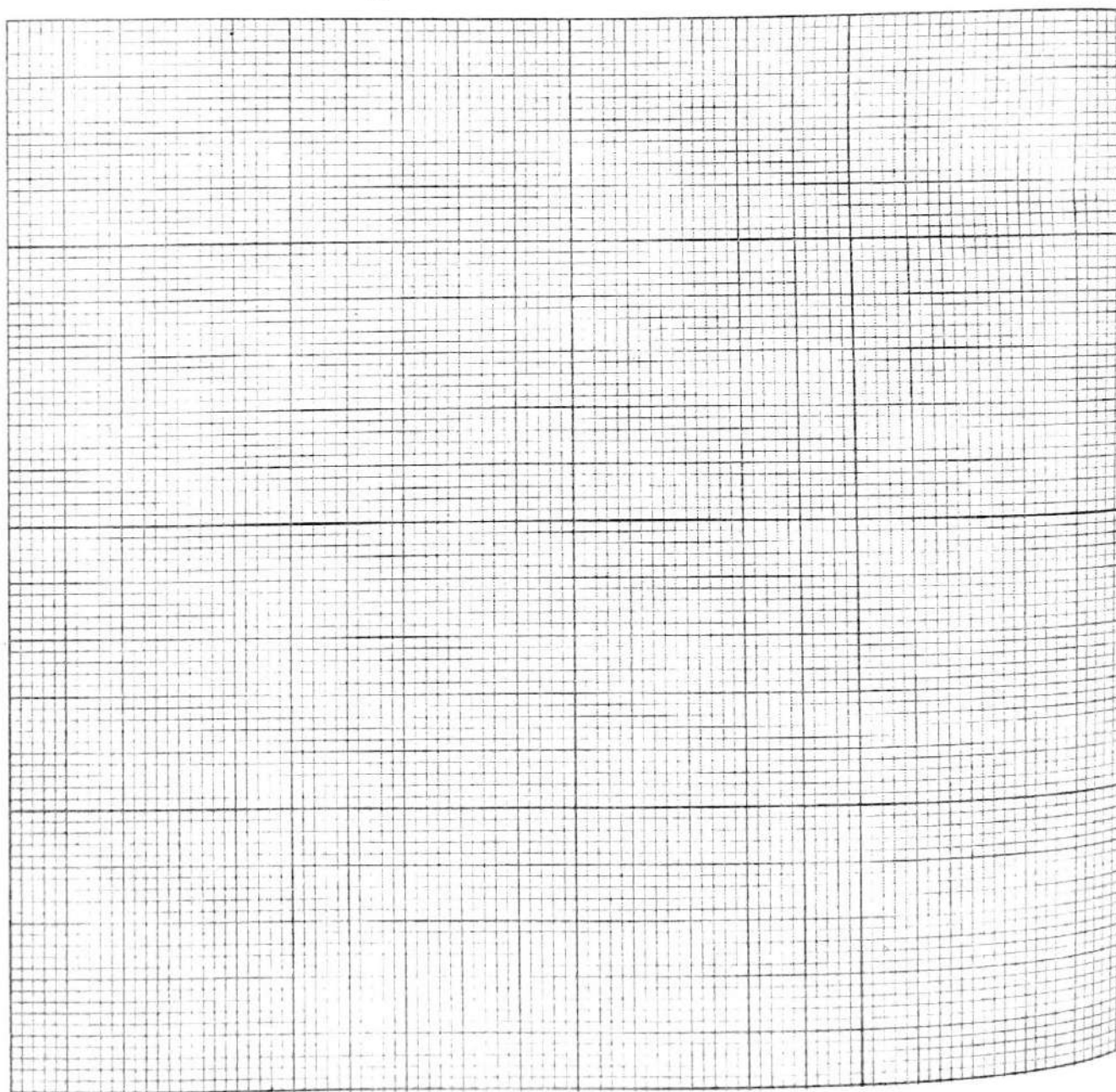
(ii) State Raoult's law of vapour pressure lowering.

(1 mark)

- (b) The vapour pressure of aqueous solutions of glucose containing 9.0g of water at 27°C varies with the mass of glucose dissolved as shown in the table below.

Mass of glucose dissolved in 9.0g of water	0.00	0.45	0.90	1.80	3.60	4.50	7.20
Vapour pressure of solution (mmHg)	31.82	31.66	31.50	31.32	30.55	30.23	29.27

- (i) Plot a graph of lowering in vapour pressure ( $\Delta P$ ) against mass of glucose dissolved in 9.0g of water. (4 marks)



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**END**

1 H		hydrogen																2 He		helium											
1.00794																		4.0026													
3 Li		4 Be																		5 B		6 C		7 N		8 O		9 F		10 Ne	
6.941		9.0122																		10.811		12.011		14.007		15.999		18.998		20.180	
sodium		magnesium																		aluminium		silicon		phosphorus		sulphur		chlorine		argon	
11		12																		13		14		15		16		17		18	
Na		Mg																		Al		Si		P		S		Cl		Ar	
22.990		24.305																		26.982		28.086		30.974		32.065		35.453		39.948	
potassium		calcium																		gallium		germanium		arsenic		selenium		bromine		krypton	
19		20																		31		32		33		34		35		36	
K		Ca																		Ga		Ge		As		Se		Br		Kr	
39.098		40.078																		69.723		72.61		74.922		78.96		79.904		83.80	
yttrium		strontium																		indium		tin		antimony		tellurium		iodine		xenon	
37		38																		49		50		51		52		53		54	
Rb		Sr																		In		Sn		Sb		Te		I		Xe	
85.468		87.62																		114.82		118.71		121.76		127.60		126.90		131.29	
barium		lanthanum																		thallium		lead		bismuth		polonium		astatine		radon	
56		57-70																		81		82		83		84		85		86	
Cs		Ba																		Tl		Pb		Bi		Po		At		Rn	
132.91		137.33																		204.38		207.2		208.98		209		210		222	
francium		actinium																		unermium		ununbium		ununium		ununeptium		unustatium		unuectium	
87		88																		110		111		112		113		114		115	
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																				Sc		Ti		V		Cr		Mn		Fe	
																				21		22		23		24		25		26	
																				63.546		64.942		65.94		66.94		67.94		68.94	
																				47.88		48.94		49.94		50.94		51.94		52.94	
																				40		41		42		43		44		45	
																				Zr		Nb		Mo		Tc		Ru		Rh	
																				39		40		41		42		43		44	
																				Y		Zr		Nb		Mo		Tc		Ru	
																				88.906		89.907		90.908		91.909		92.910		93.911	
																				lanthanum		cerium		praseodymium		neodymium		promethium		samarium	
																				71		72		73		74		75		76	
																				Lu		Hf		Ta		W		Re		Os	
																				174.967		178.49		180.948		183.84		186.207		188.905	
																				Lr		Rf		Db		Sg		Bh		Hs	
																				123.1		123.1		123.1		123.1		123.1		123.1	
																				Y		Zr		Nb		Mo		Tc		Ru	
																				Sc		Ti		V		Cr		Mn		Fe	
																				21		22		23		24		25		26	
																				63.546		64.942		65.94		66.94		67.94		68.94	
																				47.88		48.94		49.94		50.94		51.94		52.94	
																				40		41		42		43		44		45	
																				Zr		Nb		Mo		Tc		Ru		Rh	
																				39		40		41		42		43		44	
																				Y		Zr		Nb		Mo		Tc		Ru	
																				88.906		89.907		90.908		91.909		92.910		93.911	
																				lanthanum		cerium		praseodymium		neodymium		promethium		samarium	
																				71		72		73		74		75		76	
																				Lu		Hf		Ta		W		Re		Os	
																				1											

Lanthanide series

Actinide series

57 La 138.91	58 Ce 140.12	59 Pr 140.91	60 Nd 144.24	61 Pm 145	62 Sm 150.36	63 Eu 151.96	64 Gd 157.25	65 Tb 158.93	66 Dy 162.50	67 Ho 164.93	68 Er 167.26	69 Tm 168.93	70 Yb 173.04
89 Ac 227	90 Th 232.04	91 Pa 231.04	92 U 238.03	93 Np 237	94 Pu 244	95 Am 243	96 Cm 247	97 Bk 247	98 Cf 251	99 Es 252	100 Fm 257	101 Md 258	102 No 259