

**UNIVERSITY OF SWAZILAND  
SUPPLEMENTARY EXAMINATION 2007-08**

---

**TITLE OF PAPER** : Organic Chemistry

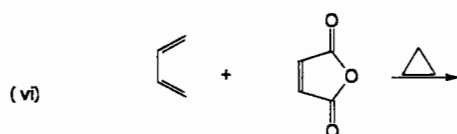
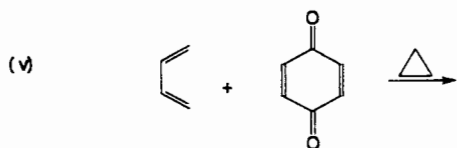
**COURSE NUMBER** : C303

**TIME** : Three Hours

**INSTRUCTIONS** : Answer any **FOUR** questions.  
Each question carries **25** marks.

---

*You are not supposed to open this paper until permission to do so has been granted by the Chief Invigilator.*

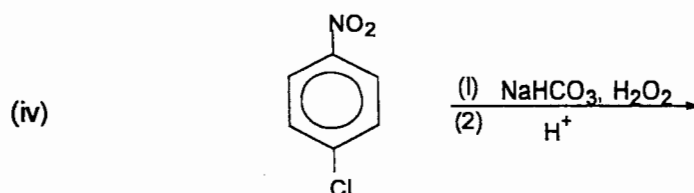
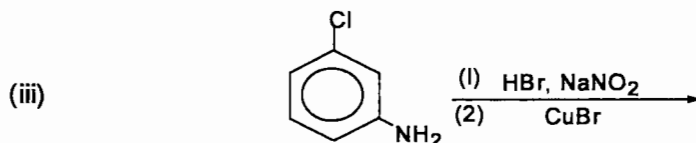
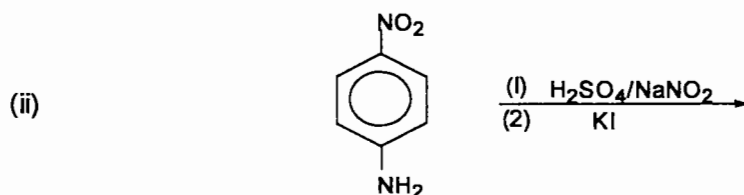
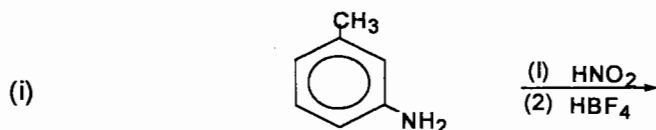


(b) What is meant by the term “disrotatory motion” in electrocyclic reactions? Illustrate your answer with any diene. (4)

(c) Why is thermal cycloaddition of ethene difficult? (3)

#### QUESTION 4

(a) Write the structure of the product of each step of the following: (16)

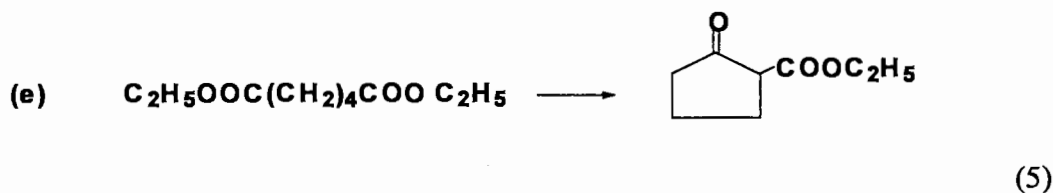


- (b) Write all steps involved in the mechanism of nucleophilic aromatic substitution involving Meisenheimer complex. What is the effect of electron-withdrawing groups on the substitution? (9)

### QUESTION 5

Write down the likely steps in the following transformations:

- (a)  $\text{CH}_3\text{COCH}_3 + \text{Br}_2 / \text{H}^+ \rightarrow \text{CH}_3\text{COCH}_2\text{Br}$  (4)
- (b)  $\text{CH}_3\text{CHO} + \text{OH}^- \rightarrow \text{CH}_3\text{CH}(\text{OH})\text{CH}_2\text{CHO}$  (3)
- (c)  $\text{CH}_3\text{CH}_2\text{CH}_2\text{CH}_2\text{OH} \rightarrow \text{CH}_3\text{CH}_2\text{CH}_2\text{CH}_2\text{CH}(\text{C}_2\text{H}_5)\text{CH}_2\text{OH}$  (7)
- (d)  $\text{CH}_3\text{COO C}_2\text{H}_5 \rightarrow \text{CH}_3\text{COCH}_2\text{COO C}_2\text{H}_5$  (6)



### QUESTION 6

- (a) (i) Write the three isomeric ethers with the molecular formula  $\text{C}_4\text{H}_{10}\text{O}$  and name them.
- (ii) State how many signals will arise in the  $^{13}\text{C}$  nmr spectrum of each named compound. (12)
- (b) (i) How many ions would you expect in the mass spectrum of 2-hexene by  $\beta$ -fragmentation?
- (ii) What are the masses of the ions in (i) above? (6)
- (c) Predict the structure of a compound with molecular formula  $\text{C}_7\text{H}_8\text{O}$  that has  $^1\text{H}$  nmr signals at  $\delta = 7.3, 4.4$  and  $3.7$  ppm with relative intensities of 7:2.9:1.4 respectively. (7)

**DEPARTMENT OF CHEMISTRY  
UNIVERSITY OF SWAZILAND**

C304

**INSTRUMENTAL ANALYSIS**

**JULY 2008 SUPPLEMENTARY EXAMINATION**

**Time Allowed:**

**Three (3) Hours**

---

---

**Instructions:**

1. This examination has six (6) questions and two (2) data sheets. The total number of pages is six (6) including this page.
2. Answer any four (4) questions fully; diagrams should be clear, large and properly labeled. Marks will be deducted for improper units and lack of procedural steps in calculations.
3. Each question is worth 25 marks.

**Special Requirements**

1. Data sheets.
2. Graph paper.

**YOU ARE NOT SUPPOSED TO OPEN THIS PAPER UNTIL PERMISSION TO DO SO HAS BEEN GIVEN BY THE CHIEF INVIGILATOR.**

**QUESTION 1 [25]**

- a. i) Use diagrams to explain what is meant by resistance to mass transfer in the stagnant mobile phase. [3]
- ii) State the HETP equation for resistance to mass transfer in the stagnant mobile phase, and explain all terms appearing in it. [4]
- iii) For each of any four (4) of these terms above, explain how it influences bandbroadening in practical terms. [4]
- b. With regards to liquid chromatography,
- i) What does the acronym "HPLC" stand for? [1]
- ii) Describe three (3) ways of degassing solvents in HPLC. [3]
- iii) Explain why two solvent reservoirs are used instead of one in HPLC. [2]
- iv) Give two (2) reasons for using a pre-column in HPLC. [2]
- v) State one advantage and one disadvantage of using microparticle packings in HPLC. [2]
- c. Use diagrams to explain how a "flow-through" ultraviolet detector works in HPLC. [4]

**QUESTION 2 [25]**

- a. In the table below,

Spectral Region	Energy (J)	Type of Transition	Wavelength (nm)	Frequency ( $\text{sec}^{-1}$ )	Wavenumber ( $\text{cm}^{-1}$ )	Energy (eV)
gamma rays		A		$>10^{20}$		
x-ray		B				120
uv-visible	F	C	700	H		I
infra-red		D			4,000	
micro-wave			G	$10^8$	J	

- i. State A, B, C, D [4]
- ii. Calculate F, G, H, I, J [5]
- b. On a single plot, draw the blackbody radiation profiles of sources heated at 500K and at 2000K. [2]
- c. Explain why the 500K plot in (b) above is the ideal one for use as an IR source, and explain its implication on resolution. [2]
- d. Draw and label the "Bunsen" arrangement of optical components in a spectrometer. [5]
- e. Explain how a single wavelength is practically selected in the spectrometer in (d) above. [3]
- f. Explain how the slit width affects resolution in infra-red spectroscopy. [4]

**QUESTION 3 [25]**

- a. For each of the five (5) desirable properties of a solid support for gas chromatography,
- i) Discuss the property desired [5]
- ii) Explain how each property relates to the Van Deemter equation [5]
- b. Describe the solid support "Chromosorb T". [2]

- c. In GC, the column is kept in an oven, whereas in LC, the column is kept at ambient temperature. Explain why this is so. [2]
- d. Use diagrams and equations to explain how a flame ionization detector works. List three (3) compounds that can not be detected by this method and explain why. [5]
- e. Use diagrams and equations to explain how an electron capture detector works. List three (3) compounds that can not be detected by this method and explain why. [6]

#### **QUESTION 4** [25]

- a. What does the acronym “ICP-OES” stand for? [1]
- b. Give operational definitions of:
- i) a plasma [2]
  - ii) an inductively coupled plasma [2]
- c. The ICP is normally referred to as a “non-chemical” flame. Explain the meaning and significance of this. [2]
- d. Draw the ICP torch and label all its components. [4]
- e. Analytical ICP operates under a range of experimental parameters. Give the optimum range for the following:
- iii) radio frequency of coil in MHz [1]
  - iv) operating power in kW [1]
  - v) main argon gas flow in mL/min [1]
  - vi) auxiliary gas flow in mL/min [1]
- f. List and describe three (3) disadvantages of graphite furnace AAS when compared to flame AAS. [3]
- g. Explain why at low frequencies the plasma shape in ICP-OES assembles a “teardrop”, whereas at high frequencies the plasma resembles a “doughnut”. [2]
- h. In the ICP, “calibration curves cover almost 5 orders of magnitude”. Explain the meaning and significance of this phrase. [2]
- i. Use diagrams to explain why ICP-OES can make the simultaneous determination of up to 35 elements possible. [3]

#### **QUESTION 5** [25]

- a. In the following table:

Wavelength	Colour absorbed	Colour observed
380-420	Violet	A
470-500	D	Red
620-680	Red	C
680-780	B	Green

Complete the table by stating A, B, C, D [4]

- b. State Bragg’s Law and draw the Czerny-Turner arrangement of opticals in a spectrometer. [4]
- c. With the aid of a large diagram, explain how the “barrier layer cell” detector works in spectroscopy. [3]

- d. What is meant by a “resonance line” in atomic spectroscopy? [1]
- i) Explain why atomic spectra appear as lines when compared to molecular spectra. [2]
- ii) The 228.8nm cadmium line corresponds to a  $^1S_1 \rightarrow ^1S_0$  transition. Would emission or absorption be chosen for its analysis at 2250°C flame operating temperatures? Explain why. [3]
- e. For each of the following interferences in flame atomic absorption spectroscopy, describe the interference and explain how it is eliminated.
- i) spectral [3]
- ii) physical [3]
- f. List and describe two (2) disadvantages of graphite furnace atomic absorption spectroscopy over flame methods. [2]

### **QUESTION 6 [25]**

- b. i) What is meant by “chromophore” in uv-visible spectroscopy? [1]
- ii) State the difference between a “bathochromic” and “hypsochromic” shift in uv-visible spectroscopy. [2]
- c. Explain how the following cause deviations in Beer’s Law
- i) slit width [2]
- ii) pH [2]
- iii) temperature [2]
- d. Explain why IR spectroscopy is performed almost exclusively using double beam instruments. [3]
- e. Use diagrams to explain how a “circular cage” photomultiplier tube works. [4]
- f. Use diagrams to explain how the Molar Ratio method is used to determine the stoichiometry of the Fe- bipyridine complex, given that bipyridine does not absorb at the wavelength of analysis at 522 nm. [3]
- g. Explain how uv-visible spectroscopy can be used to simultaneously measure more than one component in a mixture. [3]
- h. Use equations to explain why components in a spectrometer are kept in a darkened compartment. [3]





1. PERIODIC CHART OF THE ELEMENTS

1 1A 3 Li 6.941	2 2A 4 Be 9.0122											13 3A 5 B 10.81	14 4A 6 C 12.011	15 5A 7 N 14.0067	16 6A 8 O 15.9994	17 7A 9 F 18.99840	18 2 He 4.00260				
11 3B 23 Na 22.98977	12 4B 24 Mg 24.305	3 3B 39 K 39.0983	4 4B 40 Ca 40.08	5 5B 44 Sc 44.9559	6 6B 46 Ti 47.88	7 7B 50 V 50.9415	8 8B 52 Cr 51.996	9 8B 54 Mn 54.9380	10 8B 56 Fe 55.847	11 8B 58 Co 58.9332	12 8B 59 Ni 58.69	13 9B 63 Cu 63.546	14 10B 65 Zn 65.38	15 10B 69 Ga 69.72	16 10B 72 Ge 72.59	17 10B 74 As 74.9216	18 10B 76 Se 78.96	19 10B 79 Br 79.904	20 10B 81 Kr 83.80		
37 11B 85 Rb 85.4678	38 12B 86 Sr 87.62	39 11B 88 Y 88.9058	40 12B 90 Zr 91.22	41 12B 92 Nb 92.9064	42 12B 94 Mo 95.94	43 12B 96 Tc (98)	44 12B 98 Ru 101.07	45 12B 100 Rh 102.9055	46 12B 102 Pd 106.42	47 12B 106 Ag 107.8682	48 12B 108 Cd 112.411	49 12B 114 In 114.82	50 12B 116 Sn 118.69	51 12B 121 Sb 121.75	52 12B 127 Te 127.60	53 12B 126 I 126.9054	54 12B 131 Xe 131.29				
55 13B 132 Cs 132.9055	56 14B 137 Ba 137.33	57 13B 138 La 138.9055	58 14B 178 Hf 178.49	59 14B 180 Ta 180.9479	60 14B 182 W 183.85	61 14B 186 Re 186.207	62 14B 190 Os 190.2	63 14B 192 Ir 192.22	64 14B 195 Pt 195.08	65 14B 197 Au 196.9665	66 14B 200 Hg 200.59	67 14B 204 Tl 204.383	68 14B 207 Pb 207.2	69 14B 208 Bi 208.9804	70 14B (209)	71 14B 209 Po (209)	72 14B 210 At (210)	73 14B 210 Rn (222)			
87 15B 223 Fr (223)	88 16B 226 Ra (226)	89 15B 227 Ac (227)	104 Unq (261)	105 Unp (262)	106 Unh (263)	107 Uns (264)	108 Uno (265)	109 Une (266)	A value in brackets denotes the mass number of the longest lived or best known isotope.												
★ Lanthanide series		58 140.12 Ce	59 140.9077 Pr	60 144.24 Nd	61 (145) Pm	62 150.36 Sm	63 151.96 Eu	64 157.25 Gd	65 158.9254 Tb	66 162.50 Dy	67 164.9304 Ho	68 167.26 Er	69 168.9342 Tm	70 173.04 Yb	71 174.967 Lu						
▲ Actinide series		90 232.0381 Th	91 231.0369 Pa	92 238.0289 U	93 237.0482 Np	94 244 Pu	95 247 Am	96 251 Cm	97 257 Bk	98 261 Cf	99 267 Es	100 271 Fm	101 277 Md	102 285 No	103 289 Lr						

4. NET STABILITY CONSTANTS

Ag(CN) <sub>2</sub> <sup>-</sup>	5 × 10
Ag(NH <sub>3</sub> ) <sub>2</sub> <sup>+</sup>	1.6 × 10
Ag(S <sub>2</sub> O <sub>3</sub> ) <sub>2</sub> <sup>-3</sup>	4.7 × 10
Al(OH) <sub>4</sub> <sup>-</sup>	1.0 × 10
Ca(EDTA)	1.0 × 10
Cd(CN) <sub>4</sub>	8.3 × 10
Cd(NH <sub>3</sub> ) <sub>4</sub> <sup>++</sup>	5.5 × 10
Co(NH <sub>3</sub> ) <sub>6</sub> <sup>+3</sup>	2 × 10
Cr(OH) <sub>4</sub> <sup>-</sup>	4 × 10
Cu(CN) <sub>4</sub> <sup>-3</sup>	1 × 10
Cu(NH <sub>3</sub> ) <sub>4</sub> <sup>+2</sup>	1.2 × 10
Fe(CN) <sub>6</sub> <sup>-4</sup>	4.0 × 10
Fe(CN) <sub>6</sub> <sup>-4</sup>	2.5 × 10
Fe(SCN) <sub>6</sub> <sup>++</sup>	1.0 × 10
HgCl <sub>4</sub>	1.3 × 10
Hg(CN) <sub>4</sub>	8.3 × 10
Hg(SCN) <sub>4</sub>	5.0 × 10
HgI <sub>4</sub>	6.3 × 10
Mg(EDTA)	1.3 × 10
Ni(NH <sub>3</sub> ) <sub>6</sub> <sup>++</sup>	4.7 × 10
Pb(OH) <sub>3</sub> <sup>-</sup>	7.9 × 10
Zn(CN) <sub>4</sub> <sup>-2</sup>	4.2 × 10
Zn(NH <sub>3</sub> ) <sub>4</sub> <sup>++</sup>	7.8 × 10
Zn(OH) <sub>4</sub> <sup>-2</sup>	6.3 × 10

2. IONIZATION CONSTANTS (K<sub>a</sub>) FOR WEAK ACIDS

Acetic	1.9 × 10 <sup>-5</sup>	Hypochlorous	3.7 × 10 <sup>-8</sup>
2-Amino-pyridinium Ion	2 × 10 <sup>-7</sup>	H <sub>2</sub> S	K <sub>1</sub> 9 × 10 <sup>-8</sup> K <sub>2</sub> 1 × 10 <sup>-15</sup>
Ammonium Ion	5.6 × 10 <sup>-10</sup>	Imidazolium Ion	1.1 × 10 <sup>-7</sup>
Anilinium Ion	2.3 × 10 <sup>-5</sup>	Lactic	1.4 × 10 <sup>-4</sup>
Arsenic	K <sub>1</sub> 5.6 × 10 <sup>-3</sup>	Methylammonium Ion	2.7 × 10 <sup>-11</sup>
Benzoic	6.7 × 10 <sup>-5</sup>	Monoethanol-ammonium Ion	3 × 10 <sup>-10</sup>
Boric	K <sub>1</sub> 5 × 10 <sup>-10</sup>	Nicotinium Ion	9.6 × 10 <sup>-10</sup>
Carbonic	K <sub>1</sub> 4.3 × 10 <sup>-7</sup> K <sub>2</sub> 5.6 × 10 <sup>-11</sup>	Oxalic	K <sub>1</sub> 6 × 10 <sup>-2</sup> K <sub>2</sub> 6 × 10 <sup>-5</sup>
Chloroacetic	1.5 × 10 <sup>-5</sup>	Phenol	1.3 × 10 <sup>-10</sup>
Chromic	K <sub>2</sub> 3.2 × 10 <sup>-7</sup>	Phthalic	K <sub>2</sub> 4 × 10 <sup>-6</sup>
Citric	K <sub>1</sub> 8.7 × 10 <sup>-4</sup> K <sub>2</sub> 1.8 × 10 <sup>-5</sup> K <sub>3</sub> 4 × 10 <sup>-6</sup>	Phosphoric	K <sub>1</sub> 7.5 × 10 <sup>-3</sup> K <sub>2</sub> 6.2 × 10 <sup>-8</sup> K <sub>3</sub> 4.7 × 10 <sup>-13</sup>
Dichloroacetic	5 × 10 <sup>-2</sup>	Phosphorous	K <sub>1</sub> 1.0 × 10 <sup>-2</sup> K <sub>2</sub> 2.6 × 10 <sup>-7</sup>
EDTA	K <sub>1</sub> 7 × 10 <sup>-3</sup> K <sub>2</sub> 2 × 10 <sup>-3</sup> K <sub>3</sub> 7 × 10 <sup>-7</sup> K <sub>4</sub> 6 × 10 <sup>-11</sup>	Pyridinium Ion	1 × 10 <sup>-5</sup>
Formic	2 × 10 <sup>-4</sup>	Succinic	K <sub>1</sub> 7 × 10 <sup>-5</sup> K <sub>2</sub> 2.5 × 10 <sup>-6</sup>
α-D(+)-Glucose	5.2 × 10 <sup>-13</sup>	Sulfuric	K <sub>2</sub> 1.2 × 10 <sup>-2</sup>
Glycinium Ion	K <sub>1</sub> 4.6 × 10 <sup>-3</sup> K <sub>2</sub> 2.5 × 10 <sup>-10</sup>	Sulfurous	K <sub>1</sub> 2 × 10 <sup>-2</sup> K <sub>2</sub> 6 × 10 <sup>-8</sup>
Hydrazinium Ion	5.9 × 10 <sup>-9</sup>	Trimethyl-ammonium Ion	1.6 × 10 <sup>-10</sup>
Hydrocyanic	7 × 10 <sup>-10</sup>	Uric	1.3 × 10 <sup>-4</sup>
Hydrofluoric	7 × 10 <sup>-4</sup>	Water, K <sub>w</sub> , 24°C	1.0 × 10 <sup>-14</sup>
Hydroxyl-ammonium Ion	9.1 × 10 <sup>-7</sup>		

3. SOLUBILITY PRODUCT CONSTANTS

AgBr	4 × 10 <sup>-13</sup>	BaCO <sub>3</sub>	2 × 10 <sup>-8</sup>	KClO <sub>4</sub>	2 × 10 <sup>-2</sup>
Ag <sub>2</sub> CO <sub>3</sub>	6 × 10 <sup>-12</sup>	BaSO <sub>4</sub>	1 × 10 <sup>-10</sup>	MgCO <sub>3</sub>	1 × 10 <sup>-5</sup>
AgCl	1 × 10 <sup>-10</sup>	CaCO <sub>3</sub>	5 × 10 <sup>-9</sup>	MgC <sub>2</sub> O <sub>4</sub>	9 × 10 <sup>-5</sup>
Ag <sub>2</sub> CrO <sub>4</sub>	2 × 10 <sup>-12</sup>	CaF <sub>2</sub>	4 × 10 <sup>-11</sup>	MgNH <sub>4</sub> PO <sub>4</sub>	2 × 10 <sup>-13</sup>
Ag[Ag(CN) <sub>2</sub> ] <sub>4</sub>	4 × 10 <sup>-12</sup>	CaC <sub>2</sub> O <sub>4</sub>	2 × 10 <sup>-9</sup>	Mg(OH) <sub>2</sub>	1 × 10 <sup>-11</sup>
AgI	1 × 10 <sup>-16</sup>	CdS	1 × 10 <sup>-28</sup>	MnS	1 × 10 <sup>-15</sup>
Ag <sub>3</sub> PO <sub>4</sub>	1 × 10 <sup>-19</sup>	Cu(OH) <sub>2</sub>	2 × 10 <sup>-20</sup>	PbCrO <sub>4</sub>	2 × 10 <sup>-14</sup>
Ag <sub>2</sub> S	1 × 10 <sup>-50</sup>	CuS	1 × 10 <sup>-36</sup>	PbS	1 × 10 <sup>-28</sup>
AgCNS	1 × 10 <sup>-12</sup>	Fe(OH) <sub>3</sub>	1 × 10 <sup>-36</sup>	PbSO <sub>4</sub>	2 × 10 <sup>-8</sup>
Al(OH) <sub>3</sub>	2 × 10 <sup>-32</sup>	Hg <sub>2</sub> Br <sub>2</sub>	3 × 10 <sup>-23</sup>	ZnCO <sub>3</sub>	4 × 10 <sup>-5</sup>
BaCO <sub>3</sub>	5 × 10 <sup>-9</sup>	Hg <sub>2</sub> Cl <sub>2</sub>	6 × 10 <sup>-19</sup>	Sr(OH) <sub>2</sub>	3.6 × 10 <sup>-16</sup>
BaCrO <sub>4</sub>	1 × 10 <sup>-10</sup>	HgS	1 × 10 <sup>-52</sup>	ZnS	1 × 10 <sup>-24</sup>

5. FIRST IONIZATION ENERGIES, e.v.

1A	2A											14					
5.4	9.3											8.3	11	15	14	7.7	
5.1	7.6	38	48	58	68	78	88	18	28	6.0	8.1	11	10	9.8	12		
4.3	6.1	6.6	6.8	6.7	6.8	7.4	7.9	7.9	7.6	7.7	9.4	6.0	8.1	10	9.8	12	
4.2	5.7	6.6	7.0	6.8	7.2	7.5	7.7	8.3	7.6	9.0	5.8	7.3	8.6	9.0	10		
3.9	5.2	5.0	5.5	6	6.0	7.9	8.7	9.2	9.0	9.2	10	6.1	7.4	8			

6. ELECTRONEGATIVITIES, Pauling

1A	2A											21					
1.0	1.5											2.0	2.5	3.0	3.5	4.0	
0.9	1.2	38	48	58	68	78	88	18	28	1.5	1.8	2.1	2.5	3.0			
0.8	1.0	1.3	1.5	1.6	1.6	1.5	1.8	1.8	1.8	1.9	1.6	1.8	1.8	2.0	2.4	2.8	
0.8	1.0	1.2	1.4	1.6	1.8	1.9	2.2	2.2	2.2	1.9	1.7	1.7	1.8	1.9	2.1	2.5	
0.7	0.9	1.1	1.3	1.5	1.7	1.9	2.2	2.2	2.2	2.4	1.9	1.8	1.8	1.9	2.0	2.2	

7. ATOMIC RADII picometers

1A	2A											37					
155	112											98	91	92	73	71	
190	160	38	48	58	68	78	88	18	28	143	132	128	127	99			
235	197	162	147	134	130	135	126	125	124	128	138	141	137	139	140	114	
248	216	178	160	148	139	136	134	134	137	144	154	166	162	169	160	133	
287	222	187	167	149	141	137	135	136	139	146	157	171	175	170	176		

8. IONIC RADII pm

Li <sup>+</sup>	60	Sr <sup>+2</sup>	113	S <sup>-2</sup>	184
Na <sup>+</sup>	95	Ba <sup>+2</sup>	135	Se <sup>-2</sup>	198
K <sup>+</sup>	133	B <sup>+3</sup>	20	Te <sup>-2</sup>	221
Rb <sup>+</sup>	148	Al <sup>+3</sup>	50	F <sup>-</sup>	136
Be <sup>+2</sup>	31	N <sup>+3</sup>	171	Cl <sup>-</sup>	181
Mg <sup>+2</sup>	65	P <sup>+3</sup>	212	Br <sup>-</sup>	195
Ca <sup>+2</sup>	99	O <sup>-2</sup>	140	I <sup>-</sup>	216

9. LATTICE ENERGIES

Li <sup>+</sup>		K <sup>+</sup>		Cs <sup>+</sup>	
1030	840	718	602	502	400
Na <sup>+</sup>		Rb <sup>+</sup>		Fr <sup>+</sup>	
788	704	648	548	448	348
Mg <sup>+2</sup>		Ca <sup>+2</sup>		Sr <sup>+2</sup>	
2466	1735	1462	1170	934	744
Zn <sup>+2</sup>		Ba <sup>+2</sup>		Pb <sup>+2</sup>	
2051	1462	1170	934	744	594

10. HALF LIVES

H <sup>3</sup>	12.3 years	K <sup>40</sup>	1.28 × 10 <sup>9</sup> y	Fr <sup>213</sup>	8.1 day
F <sup>20</sup>	11.4 secs	Ca <sup>45</sup>	165 days	Cs <sup>137</sup>	30 yea
C <sup>14</sup>	5730 years	Fe <sup>59</sup>	45 days	Au <sup>198</sup>	2.69 da
Na <sup>24</sup>	15.0 hours	Co <sup>60</sup>	5.26 y	Ra <sup>226</sup>	1620 y
P <sup>32</sup>	14.3 days	Br <sup>82</sup>	35.5 hours	U <sup>235</sup>	7.1 × 10 <sup>8</sup>
S <sup>35</sup>	88 days	Sr <sup>90</sup>	28 years	U <sup>238</sup>	4.51 × 10 <sup>9</sup>
Cl <sup>36</sup>	3.1 × 10 <sup>5</sup> y	I <sup>129</sup>	1.7 × 10 <sup>7</sup> y	Pu <sup>239</sup>	24,400

Electrode Potentials, E<sup>0</sup>

H<sup>+</sup> + e<sup>-</sup> ⇌ 1/2 H<sub>2</sub> E<sup>0</sup> = 0.000V

Cd<sup>2+</sup> + 2e<sup>-</sup> ⇌ Cd(s) E<sup>0</sup> = -0.246V

" " ⇌ Ag(s) + Cl<sup>-</sup> E<sup>0</sup> = -0.023V

Cu<sup>2+</sup> + 2e<sup>-</sup> ⇌ Cu(s) E<sup>0</sup> = +0.34