

Today: Finish ch. 6 (except for Born-Haber cycle section)

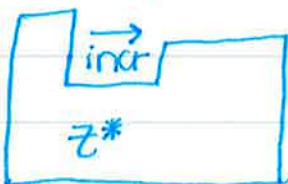
October 11<sup>th</sup>

Over Break: checklist available for CK-3, prelab  
expt 8 titration

\*Problem Club w/ kendall → Sunday October 20<sup>th</sup>  
→ Tuesday October 22<sup>nd</sup>  
↳ 7:30 - 9pm, Eppley 211

## Periodic Trends

$\exists^*$  or  $\exists^{\text{eff}}$



due to energy diagram

→ due to filling new shells

E	$\frac{1\downarrow}{\text{1s}} \frac{1\uparrow}{\text{2s}} \frac{1}{\text{2p}}$
	$\frac{1\downarrow}{\text{1s}}$

Graph showing atomic radius ( $r_e$ ) vs. ionization energy ( $E_i$ ). The y-axis is  $r_e$  and the x-axis is  $E_i$ . The curve shows a general decrease in  $r_e$  from left to right across the periodic table, with a notable peak for Helium (He).

Points marked on the curve:

- Li
- Be
- B
- C
- N
- O
- F
- Ne
- Na

Arrows above the curve point to the points for Li and Be.

Below the curve, arrows point to the electron loss processes:

$$\text{Li} + e^- \rightarrow \text{Li}^+$$

$$\text{B} + e^- \rightarrow \text{B}^+ + e^-$$

$$2s^1 \rightarrow 2s^0$$

$$2s^2 2p^1 \rightarrow 2s^2 2p^0$$

lower than

Li because of ionization energies decreasing as you go down the periodic table

$$L_i \rightarrow L_i^+ + e^- \quad B \rightarrow B^+ + e^-$$

$$Li \rightarrow Li^+ + e^- \quad B \rightarrow B^+ + e^-$$

$$2s^1 \rightarrow 2s^0 \quad 2s^2 2p^1 \rightarrow 2s^2 2p^0$$

$$n=2 \rightarrow n=\infty$$

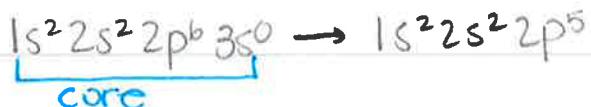
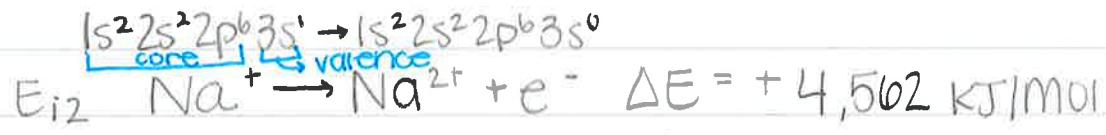
Ne / Na  
is more ionized than Li because  
of ionization energy decreasing as  
you go down the periodic table

\* All elements want to keep their electrons therefore it takes energy for EVERY ONE!

October 11<sup>th</sup>

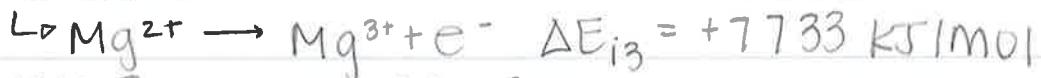
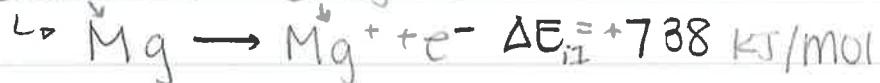


### Na



\* elements can ONLY truly lose their valence electrons, you cannot eat into the core!!!

### Mg



Electron Affinity:  $E + e^- \rightarrow E^-$



$\text{2s}^1 \quad \text{2s}^2$   $\uparrow$  energy given off/released

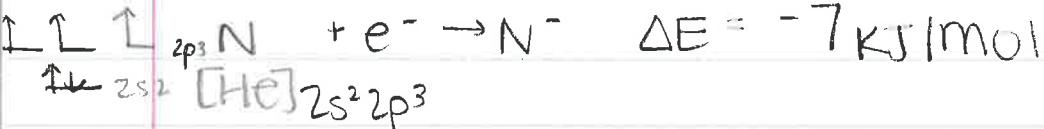


B

$$\Delta E = -27 \text{ kJ/mol}$$

C

$$\Delta E = -154 \text{ kJ/mol}$$



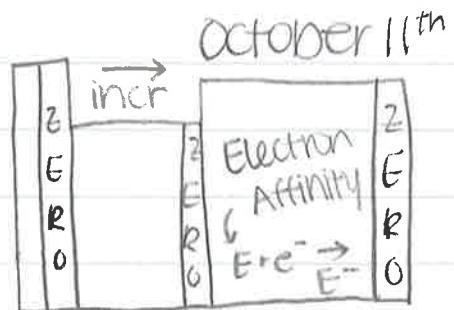
$$\Delta E_{EA}$$

$$\text{F}^- - 328 \text{ kJ/mol}$$

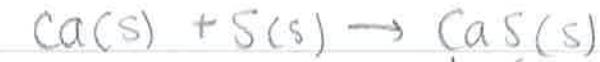
$$\text{Cl}^- - 349 \text{ kJ/mol}$$

$$\text{Br}^- - 224 \text{ kJ/mol}$$

$$\text{I}^- - 295 \text{ kJ/mol}$$



\* If a subshell is filled, they don't want another  $e^-$



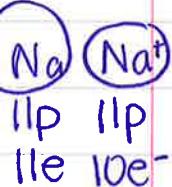
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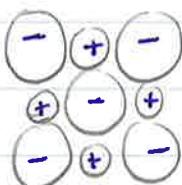
 $\text{Ca}^{2+}$  $\text{S}^{2-}$ 

reduced  
oxidized

## Lattice Energy

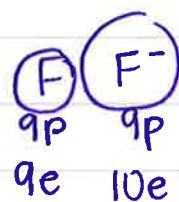


### Ionic Lattice

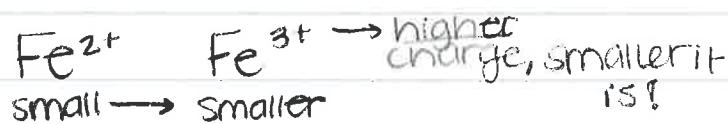


\* cations = SMALL

\* anions = BIG



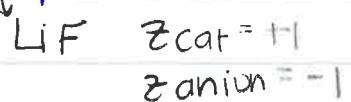
charge on cation  
↓



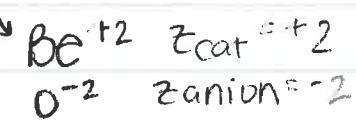
\* Forming lattice always releases energy!

Lattice Energy  $\propto Z_{\text{cation}}^2 Z_{\text{anion}}^2$

$$\rightarrow E_{\text{lattice}} = 1036 \text{ kJ/mol}$$



charge on anion  
↓



$$\rightarrow E_{\text{lattice}} = -4443 \text{ kJ/mol}$$

## Chapter 6 Day 2 (Sections 6.5 – 6.8)

1. The sequential ionization energies for silicon are (all in kJ/mol):  $E_{i1} = 787$ ;  $E_{i2} = 1577$ ;  $E_{i3} = 3231$ ;  $E_{i4} = 4356$ ;  $E_{i5} = 16091$ ;  $E_{i6} = 19784$ ; and  $E_{i7} = 23783$ . Plot these data on the graph paper below. (Make each horizontal line an increment of 5000 kJ) Write the electron configuration of the resulting ion near each data point – so for 1<sup>st</sup>, you would write  $1s^2 2s^2 2p^6 3s^2 3p^1$ , and so on.



1<sup>st</sup>    2<sup>nd</sup>    3<sup>rd</sup>    4<sup>th</sup>    5<sup>th</sup>    6<sup>th</sup>    7<sup>th</sup>

2a. What is the solubility rule involving Group 1 cations?

2b. What is the solubility rule involving halides?

2c. From the table in the book (and on the handout) what is the range of lattice energy values for Group 1 halides? Is this the energy associated with forming the lattice from gas phase ions or breaking the lattice into gas phase ions?

2d. What is energy, E, for this process? (Think carefully about the sign)  $\text{Na}^+(\text{g}) + \text{Cl}^-(\text{g}) \rightarrow \text{NaCl}(\text{s})$     E =

3a. The lattice energies of  $\text{CaSO}_4$ ,  $\text{SrSO}_4$ , and  $\text{BaSO}_4$  are 2653, 2603 and 2423 kJ/mol, respectively. Why do you think these lattice energies are larger than for Group 1 halides?

## (Unit 3) 11 October 2019

3b. How does lattice energy seem to relate to solubility?

4a. The charge on the cation is called  $Z^+$  and the charge on the anion is  $Z^-$ . Lattice energies are  $\propto Z^+ \times Z^-$ . Complete this table of  $Z^+ \times Z^-$  and give an example of each combination. Circle those that you know or suspect are insoluble from a solubility rule.

$ Z^+ \times Z^- $	$Z^+ = 1$	$Z^+ = 2$	$Z^+ = 3$
$Z^- = -1$	-1, $\text{NaCl}$		
$Z^- = -2$		-4, $\text{CaCO}_3$	
$Z^- = -3$			-9, $\text{AlPO}_4$

4b. Can you make a generalization about the product of charges and lattice energies?

### Questions in final exam format (multiple choice):

5. Which liberates the most energy?

- A.  $\text{Br}(\text{g}) + \text{e}^- \rightarrow \text{Br}^-(\text{g})$
- B.  $\text{Cl}(\text{g}) + \text{e}^- \rightarrow \text{Cl}^-(\text{g})$
- C.  $\text{F}(\text{g}) + \text{e}^- \rightarrow \text{F}^-(\text{g})$
- D.  $\text{I}(\text{g}) + \text{e}^- \rightarrow \text{I}^-(\text{g})$

6. What is the ground-state electron configuration of the ion  $\text{Cu}^{2+}$ ?

- A. [Ar] 3d9
- B. [Ar] 4s1 3d8
- C. [Ar] 4s2 3d7
- D. [Ar] 4s2 3d10 4p1

7. Which ionic compound would be expected to have the highest lattice energy?

- A.  $\text{Li}_2\text{O}$
- B.  $\text{Na}_2\text{O}$
- C.  $\text{K}_2\text{O}_2$
- D.  $\text{Ca}_3\text{PO}_4$

### Now try these problems from the book:

Section 6.5. (Electron affinity) Problems 9, 10, and 66 – 72, even.

Section 6.6. (Octet Rule) Problems 11, 12, 30, and 74 – 78, even.

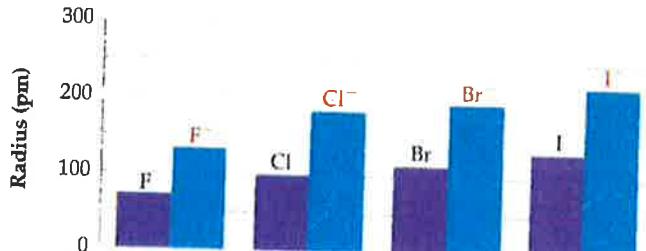
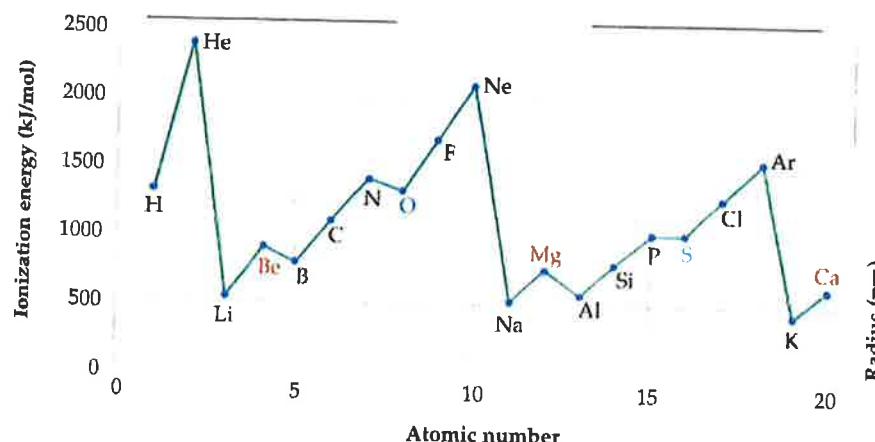
Section 6.7. Skip for now. This section will be discussed after we finish Chapter 9.

Section 6.8. (Lattice energy) Problems 15, 16, and 98.

Practice Quiz (pg. 231 – 232) Problems 9 – 10, 13.

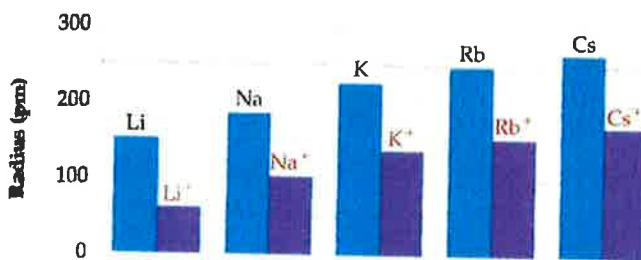
**Figure 6.4**

**Ionization energies of the first 20 elements.**

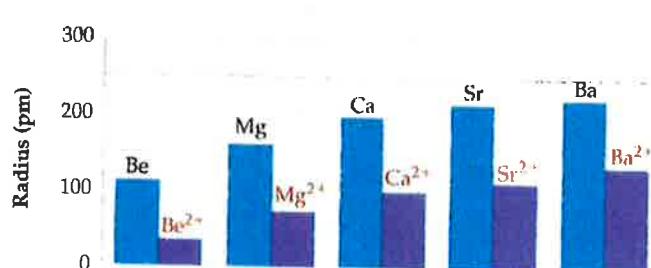


**Anions** are larger than their **neutral atoms** because of additional electron–electron repulsions and a decrease in  $Z_{\text{eff}}$ .

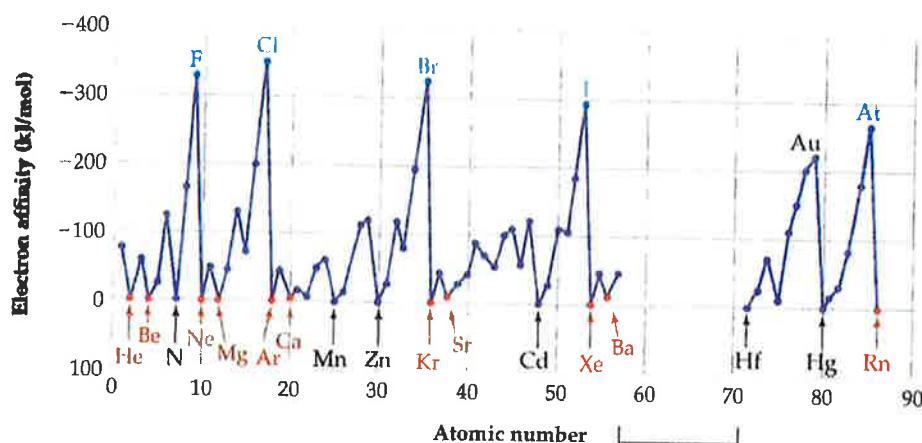
(a)



(b)



**Cations** are smaller than the corresponding **neutral atoms**, both because the principal quantum number of the valence-shell electrons is smaller for the cations than it is for the neutral atoms and because  $Z_{\text{eff}}$  is larger.



A negative value for  $E_{\text{ea}}$ , such as those for the **group 7A elements (halogens)**, means that energy is released when an electron adds to an atom.

A value of zero, such as those for the **group 2A elements (alkaline earths)** and **group 8A elements (noble gases)**, means that energy is absorbed but the exact amount can't be measured.

**TABLE 6.3 Lattice Energies of Some Ionic Solids (kJ/mol)**

Cation	Anion				
	F <sup>-</sup>	Cl <sup>-</sup>	Br <sup>-</sup>	I <sup>-</sup>	O <sup>2-</sup>
Li <sup>+</sup>	1036	853	807	757	2925
Na <sup>+</sup>	923	787	747	704	2695
K <sup>+</sup>	821	715	682	649	2360
Be <sup>2+</sup>	3505	3020	2914	2800	4443
Mg <sup>2+</sup>	2957	2524	2440	2327	3791
Ca <sup>2+</sup>	2630	2258	2176	2074	3401
Al <sup>3+</sup>	5215	5492	5361	5218	15,916