# Chapter 6 CHEMICAL PERIODICITY

## Effective nuclear charge -

### Shielding -

### GENERALIZATIONS about atomic behavior & properties

(1) Atomic radii (size)

Across a period, from left to right

Down a group,

from top to bottom

<u>(2) Io</u>	nizatio	on En	<u>ergy</u> (IE) ·	- the minimum amount of energy required to remove outermost electron(s) from an atom
А	$\rightarrow$ A	<b>^</b> +	+ e-	1st ionization energy
$A^+$	$\rightarrow$ A	2+	+ e-	2nd ionization energy
Across a period,				from left to right
Down a group,				from top to bottom

(3) Electron Affinity (EA) - energy absorbed when an atom gains an electron

$A + e \rightarrow A^{-}$	Energy released - exothermic, negative number. <u>Most negative</u> refers to element most likely to gain an e
Across a period,	from left to right
Down a group,	from top to bottom

#### (4) Ionic Radii (size)

# for atoms of SAME CHARGE, trend is exactly the same as atomic radii

cations are SMALLER that	Na <sup>+</sup>	Na	
anions are LARGER than	F	F⁻	
		$Cs^+$	F⁻
increasing + charge:	$Cr^+$	Cr <sup>2+</sup>	Cr <sup>3+</sup>

\*(5) Electronegativity \* (EN) - the relative tendency of an atom to attract electrons to itself when chemically combined with another atom ("pull")

ex.	NaCl,	Na <sup>+</sup> & CI ions metal/nonmetal	electron is <i>transferred</i> from Na to Cl
ex.	SO <sub>3</sub>	covalent bonds nonmetal/nonmetal electron pair sharing	electron is shared <i>unequally</i> between S and O
Acros	s a period,	from left to	o right

Down a group,	from top to bottom
0 17	1

How to memorize:

# Dr. Heising's method for Lewis Dot Structures

Note: These rules will not work if

- a) **hydrogen** is present in the molecule, as H does not obey the octet rule
- b) the molecule has more than one central atom

If the molecule falls into one or both of these categories, flip over for some alternate ways to get the Lewis structure.

- 1. <u>Add up the valence electrons</u>. Remember the charge (subtract for +, add for -).
- 2. <u>Divide by eight</u>. The *integer* is the number of sigma ( $\sigma$ ) bonds around the central atom. The *remainder* (if any) is the number of electrons around the central atom.
- 3. <u>Arrange the atoms with the least electronegative atom at the center</u>. Symmetric is often (but not always) correct.
- 4. <u>Draw the sigma ( $\sigma$ ) bonds</u>. Should be equal to the *integer* in step 2, and every atom should have one bond to the central atom by the time you are finished.
- 5. <u>Put the *remainder* electrons as lone pairs on the central atom</u>. If there are an odd number of electrons (infrequent but not impossible), just put the odd one by itself.
- 6. Does the central atom have 8 electrons around it? Include electrons in bonds and electrons in lone pairs in the count. If it has less than 8, make additional bonds, called pi  $(\pi)$  bonds, with the surrounding atoms to complete the octet. More than 8 electrons is OK for atoms in row 3 or higher of Periodic Table.
- 7. Fill the octets of the surrounding atoms using electron pairs.
- 8. <u>Check yourself</u>. Count the total # of electrons in bonds and lone pairs in the entire molecule. It should match the value you found in step 1.
- 9. <u>Calculate the formal charge on the central atom</u>. Choose the resonance structure with the least formal charges as your best answer. If central atom is in row 3 or higher, you may expand the octet with additional  $\pi$  bonds to minimize the formal charge.

#### The method must be modified if

#### a) Hydrogen is present.

- To which atom is the hydrogen connected? Often H is bonded to the atom next to it in the chemical formula. Ex.
  - o CHCl<sub>3</sub> has one H bonded to C, the central atom
  - $\circ$  HCO<sub>2</sub>H has one H bonded to C (central atom) and one H bonded to O.
  - $\circ$  In ternary oxoacids, (H<sub>2</sub>SO<sub>4</sub>, H<sub>3</sub>PO<sub>4</sub>, HNO<sub>3</sub>, etc.) H is bonded to O.
- Substitute a minus charge in the formula for every H present. For example, CHCl<sub>3</sub>  $\rightarrow$  CCl<sub>3</sub><sup>-</sup>, H<sub>3</sub>CCN  $\rightarrow$  CCN<sup>3-</sup>, HCO<sub>2</sub>H  $\rightarrow$  CO<sub>2</sub><sup>2-</sup>
- Follow the rules. After step 8, substitute a sigma bond to an H atom for one lone pair (each) on the appropriate atom. Avoid putting H on O atoms that form  $\pi$  bonds.

#### b) the structure has more than one 'central' atom.

- If the extra atom not bonded to the central atom is just hydrogen (Ex.,  $H_2SO_4$ ,  $H_3PO_4$ , etc.), follow the directions in method a).
- Diatomic molecules (ex.  $N_2$ ,  $O_2$ ,  $C_2$ ). Follow the book, or treat BOTH as central atoms. Complete octet with  $\pi$  bonds.
- Otherwise (organic molecules,  $P_2O_7^{4-}$ ), follow the method in the book, or do the following:
  - Do steps 1, 3, and 4. As there is more than one central atom, arrange the atoms in the most symmetric manner possible.
  - Subtract bonding electrons (step 4) from valence electrons (step 1). Distribute remainder over the peripheral atoms until the octet rule is fulfilled for all atoms (except H, of course).
    - Some left over put as lone pairs on central atoms
    - Not enough go back and turn some of the lone pairs into  $\pi$  bonds.
  - o Do steps 8 and 9.

Determine Lewis structures for the following examples: PCI3, CO2, CH2CI2, N2 PCI3 i)  $P: s^2p^3 = 5$ ci:  $s^*p^5 = 3(7)$  $75a^-$ 2) 26/8= 3 w/ 2et left 30 bonds & 1 lone pair 3) CI P CI CI (4,5)  $(1-\ddot{p}-C)$ C02 i) C: 4 O:  $\frac{2(6)}{16e^{-}/8} = 2 \sigma$  bonds 3),4),5) 0-C-0  $6) \qquad O=C=O \qquad 2\pi \ bonds$ 7,8) 0=c=0 16e

3,4,5) 
$$(1 - \ddot{C} - C)$$
  
6,7),8)  $:\ddot{C}_{1} - \ddot{C}_{2} - \dot{C}_{1}:$  20e<sup>-</sup>  
**\*** put H backin.  
**:** $\ddot{C}_{1} - \ddot{C}_{2} - \ddot{C}_{1}:$   
**H**

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