

# California State Polytechnic University, Pomona

Organic Chemistry I, CHM 3140, Dr. Laurie S. Starkey

## Ch. 1 Summary (Klein 4<sup>th</sup> edition) Chemistry Review & Intro to Organic Molecules

### I. Review of General Chemistry concepts (1.1 – 1.8)

- A) atomic structure; energy of atomic orbitals ( $s$ ,  $p$ )
- B) electronegativity - ability of an atom to attract electron density
  - i) fluorine is most electronegative element (oxygen is second-most!), periodic trends
  - ii)  $C \approx H$  electronegativity,  $N = Cl$  electronegativity
- C) a filled valence shell (full octet) imparts stability
- D) covalent vs. ionic bonds
- E) bond polarity ( $\delta^+$  and  $\delta^-$ ) **SkillBuilder 1.4**

### II. Reading Line Drawings (1.6) **SkillBuilder 1.5**

### III. Lewis Structures (1.3, 1.4) **SkillBuilder 1.2**

- A) structures show  $\sigma$ ,  $\pi$  and nonbonded electrons
- B) formal charges (1.4) **SkillBuilder 1.3**
- C) recognize "typical" configurations for common atoms (H, C, N, O, X)

### IV. Atomic Orbitals (AO's) combine to give Molecular Orbitals (MO's) (1.7 – 1.9)

- A) Bonding MO's ( $\sigma$ ,  $\pi$ ) contain electrons in covalent bonds
- B) Antibonding MO's ( $\sigma^*$ ,  $\pi^*$ ) are usually empty, can contain excited electrons
- C) Relative energies, stabilities of MO's

### V. Hybrid Orbitals (1.10) and Shape/Geometry (1.11) **SkillBuilders 1.7, 1.8**

- A)  $sp^3$  hybridization: 4 regions of electron density, tetrahedral geometry
- B)  $sp^2$  hybridization: 3 regions of electron density, trigonal planar geometry, contains an unhybridized  $p$  orbital
- C)  $sp$  hybridization: 2 regions of electron density, linear geometry, contains two unhybridized  $p$  orbitals

### VI. 3-D sketches (2.6)

- A) determine hybridization to learn geometry about each atom
- B) draw aligned  $p$  orbitals to show  $\pi$  bonds

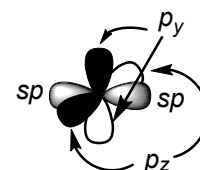
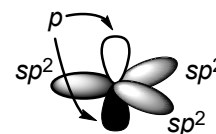
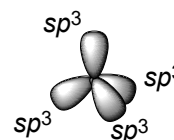
### VII. Molecular Polarity (1.12) & Physical Properties (1.13, 1.14) **SkillBuilders 1.9, 1.10**

- A) Nonbonding (intermolecular) Interactions affect bp, mp
  - i) dipole-dipole for polar molecules ( $\delta^+$ ,  $\delta^-$ )
  - ii) hydrogen bonding for molecules containing NH, OH or HF (STRONG dipole)
  - iii) van der Waals (London dispersion) temporary dipole moments
    - a) explains why bp varies by MW (higher MW, higher bp)
    - b) straight vs. branched molecules (greater surface area, higher bp)
- B) mp increases for molecules that can pack tighter (more spherical, higher mp)
- C) water solubility increases with polarity, hydrogen-bonding ability

### VIII. Isomerism (1.2) **SkillBuilder 1.1**

- A) structural (constitutional): same molecular formula, different connectivity
- B) cis-trans (stereoisomers): structures vary only by orientation in space

#### Hybridization



#### Predicting Boiling Points

Formula	C <sub>3</sub> H <sub>8</sub>	C <sub>4</sub> H <sub>10</sub>	C <sub>4</sub> H <sub>10</sub>	C <sub>5</sub> H <sub>12</sub>
BP (°C)	- 42.1	- 11.7	- 0.6	27.9
↑ Molecular Weight, ↑ BP. If MW is the same, THEN ↑ branching, ↓ BP				