

Ch. 15 Conjugated Systems

The General Stabilization Effect of Conjugation (Section 15.1, 2, 3, 8, 9)

	Conjugated (more stable)	Isolated (less stable)	Notes:
1 Cations			
2 Radicals			
3 Anions			
4 Dienes			
5 Ethers			An N or O next to a double bond becomes sp^2 . An isolated N or O is sp^3
6 Amines			
7 Esters			
8 Amides			Very special, chapter 23, all of biochemistry, proteins, enzymes, etc.
9 Oxyanions (Carboxylates)			Very special, chapter 21
10 Carbanions (Enolates)			Very special, chapter 22
11 Aromatics			Very special, chapters 16 + 17

Conjugation: Anything that is or can be sp^2 hybridized is stabilized when next to π bonds.

- oxygen, nitrogen, cations, radicals, and anions

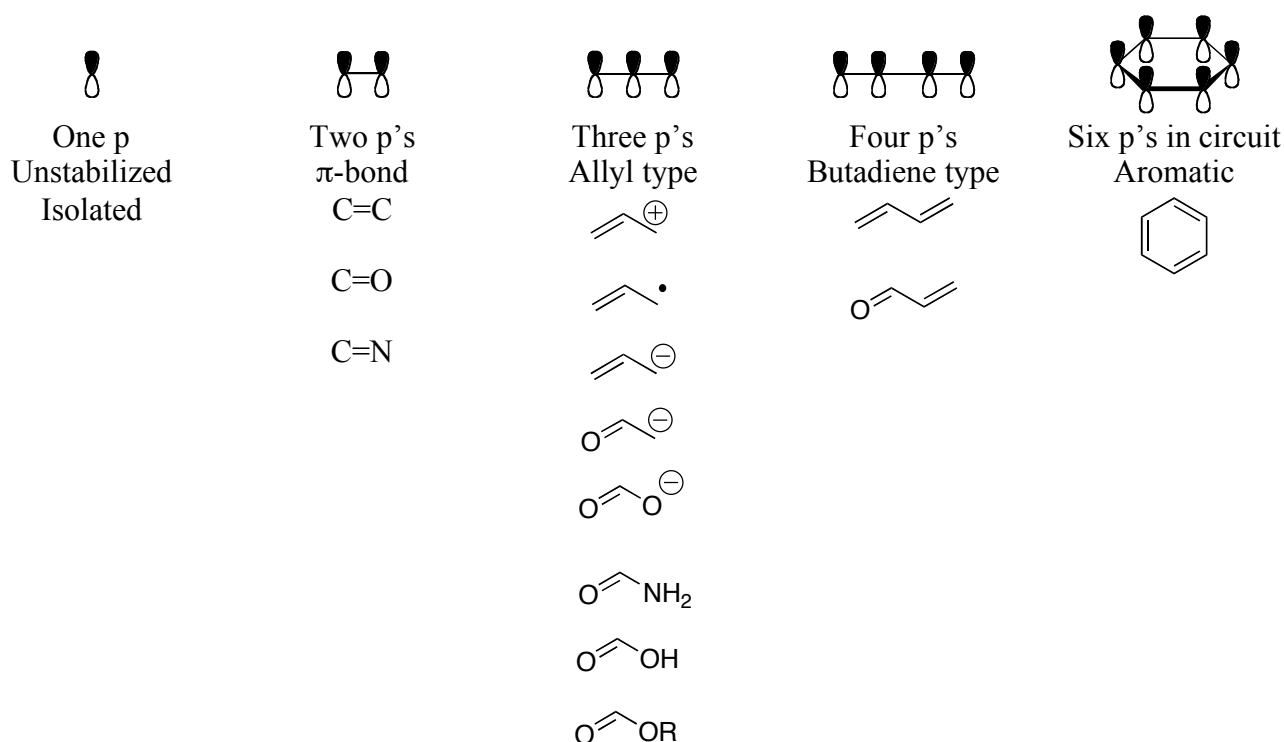
Notes:

- Any atom that can be sp^2 will be sp^2 when next to a double bond
- "Conjugation" is when sp^2 centers are joined in an uninterrupted series of 3 or more, such that an uninterrupted series of p-orbitals is possible
- Any sp^2 center has one p orbital

Impact of Conjugation

4. **Stability:** Conjugation is **stabilizing** because of p-orbital overlap (Sections 15.2, 4, 7)

- Note: In the allyl family, **resonance = conjugation**

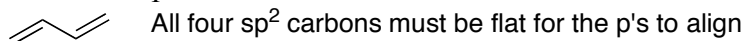


5. **Reactivity:** Conjugation-induced stability impacts **reactivity** (Sections 15.4-7)

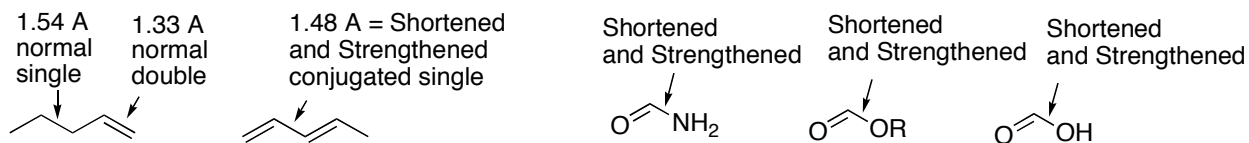
- If the **product** of a rate-determining step is stabilized, the reaction rate will go **faster** (product stability-reactivity principle)
 - Common when allylic cations, radicals, or carbanions are involved
- If the **reactant** in the rate-determining step is stabilized, the reaction rate will go **slower** (reactant stability-reactivity principle)
 - Why aromatics are so much less reactive
 - Why ester, amide, and acid carbonyls are less electrophilic than aldehydes or ketones

6. **Molecular shape** (Sections 15.3, 8, 9)

- The p-orbitals must be aligned in parallel for max overlap and max stability
- The sp^2 centers must be coplanar



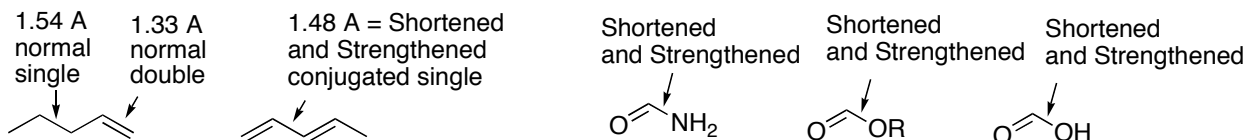
7. **Bond Length:** Bonds that look like singles but are actually between conjugated sp^2 centers are **shorter** than ordinary single bonds



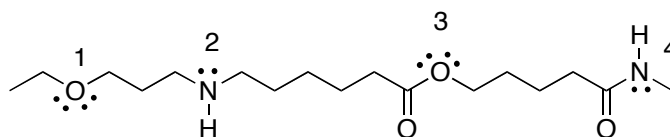
- In amides, esters, and acids, the bond between the carbonyl and the heteroatom is shortened

8. **Bond Strength:** Bonds that look like singles but are actually between conjugated sp^2 centers are **stronger** than ordinary single bonds

9. **Bond Rotation Barrier:** Bonds that look like singles but are actually between conjugated have much larger rotation barriers than ordinary single bonds
- Because in the process of rotating, the p-overlap and its associated stability would be temporarily lost



10. **Hybridization:** Conjugated sp^2 atoms have both sp^2 and p orbitals. You should always be able to classify the hybridization of **lone pairs on nitrogen and oxygen**.
- Isolated** oxygens or nitrogens: sp^3 atom hybridization, sp^3 lone-pair hybridization, and tetrahedral, 109° bond angles
 - Conjugated nitrogens:** sp^2 atom hybridization, **p lone-pair hybridization (needed for conjugation)**, and 120° bond angles
 - Conjugated oxygens:** sp^2 atom hybridization, **one p lone-pair hybridization** (needed for conjugation), **one sp^2 lone-pair**, and 120° bond angles



Atom

O-1

N-2

O-3

N-4

Isolated vs.

Conjugated

Atom

Hybridization

Lone-Pair(s)

Hybridization

Bond Angles