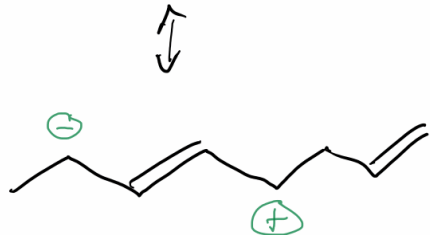
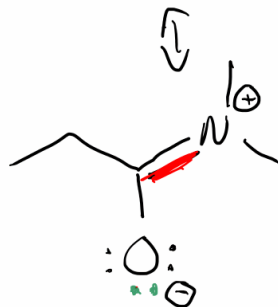
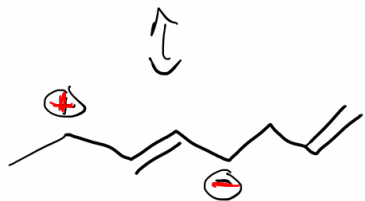
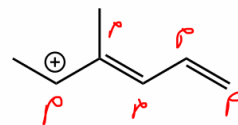
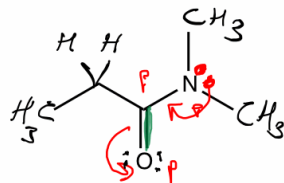
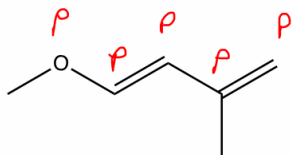
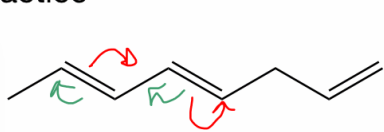


Practice



doesn't work
can't put a pi bond here because of the 2 H atoms

can't charge the charge of the molecule

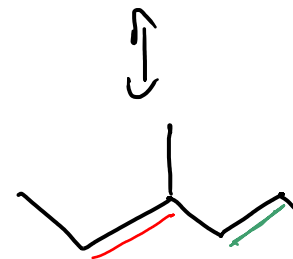
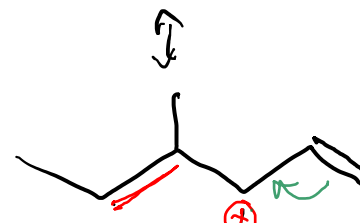
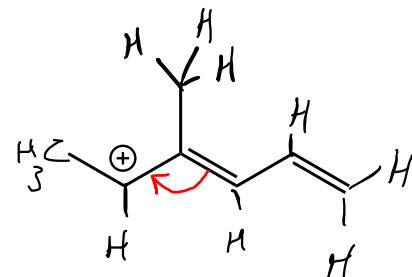
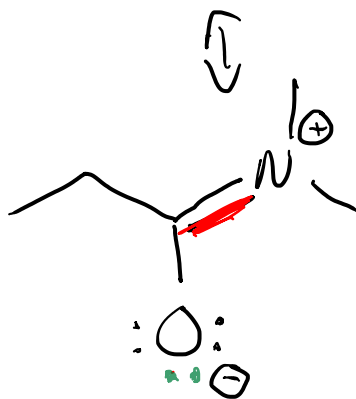
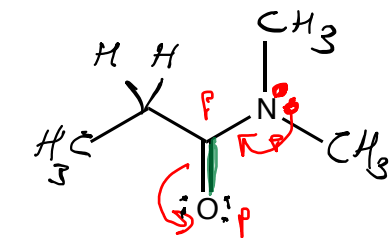
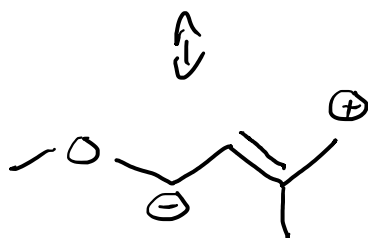
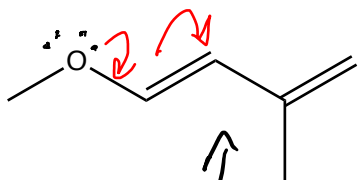
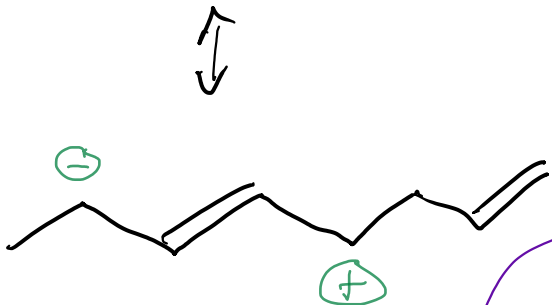
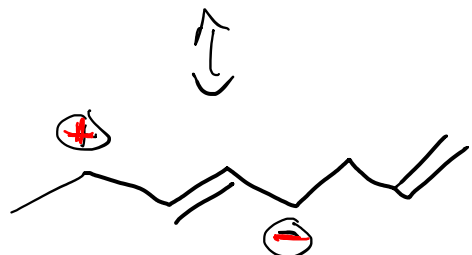
- move lp e-'s towards pi bonds
- move lp e-'s towards empty P orbitals

Strategies of p orbitals

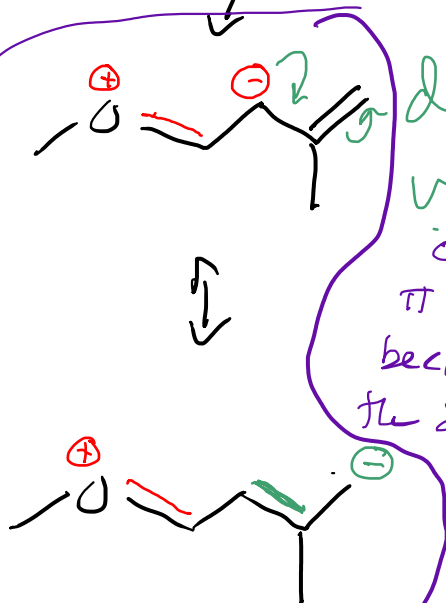
- move pi bonds towards pi bonds
- move pi bonds towards empty p orbitals.



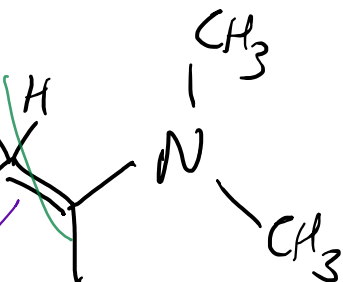
Practice



most important

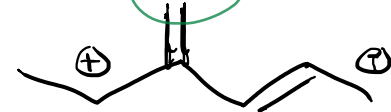


doesn't work
can't put a pi bond here because of the 2H atoms



can't charge the charge of the molecule

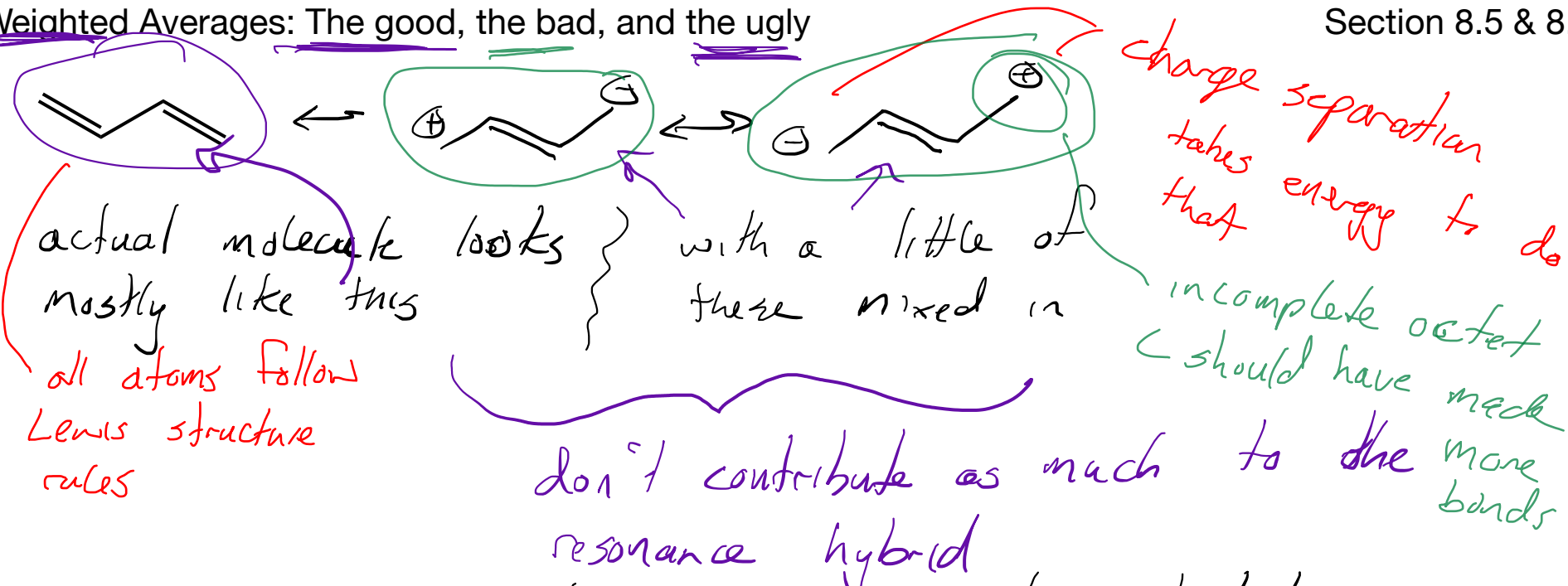
too many bonds



2+ charges

Weighted Averages: The good, the bad, and the ugly

Section 8.5 & 8.6



The "better" the contributor the more it contributes to the resonance hybrid.

Creating charge separation

Creating atoms with incomplete octets

Putting the "wrong" charge on an atom

least eneg atom with \ominus

most eneg atom with \oplus

all three problems would create a structure that would be an insignificant contributor to the resonance hybrid
don't draw it