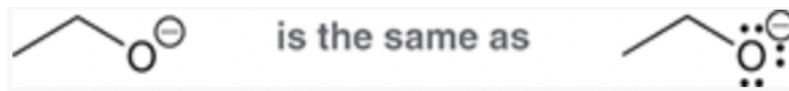


Session 2 Worksheet

You do not have to write lone pairs if you don't want to, however, you **MUST** include a formal charge (if applicable)



Resonance

• Shows distribution of charge

Resonance structures:

• used when

We represent resonance structures with brackets and double sided arrows



Note: the resonance structures are not switching back and forth! The hybrid is a mixture of both structures



Resonance hybrid
of 1 and 2

Curved Arrows:

Shows e^- as if they are moving, but they're not

Use a double-barbed arrow, single-barbed arrows show the movement of radicals (single e^-)

Molecular Orbitals

Molecular Orbital (MO):

- Represents the region of space where one or two electrons of a molecule are likely to be found
- Have a wave-like behavior with + and - lobes

Remember

The +/- lobes are not an indicator of charge, simply, they are an illustration of displacement

Bonding MO:

- Result of constructive interference
- forms bonds
- lower in energy

Anti-bonding MO:

- Result of destructive interference
- does not form bonds
- higher in energy

Which molecular orbital will be formed first?

bonding MO

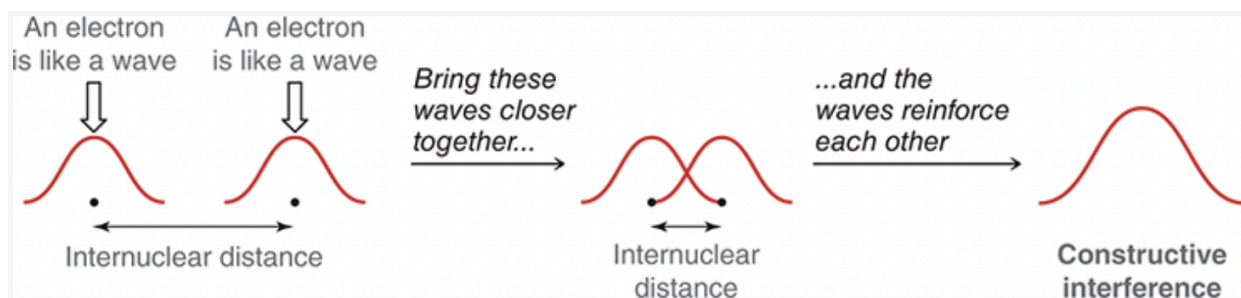
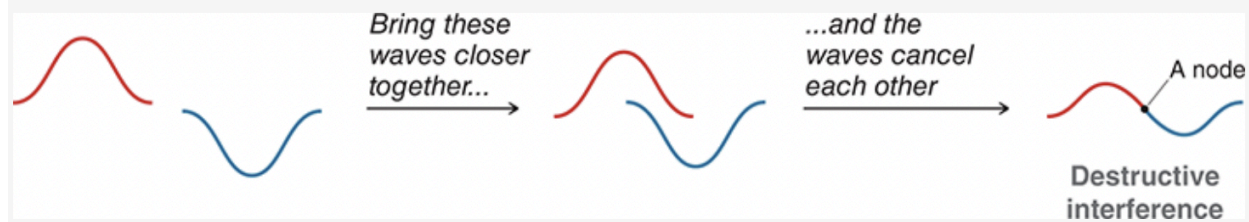
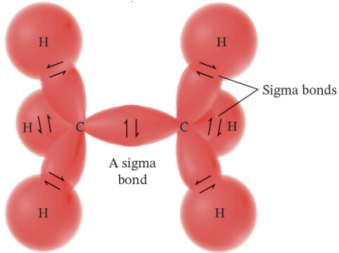
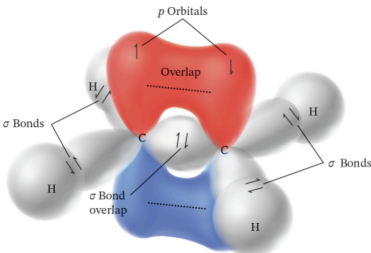
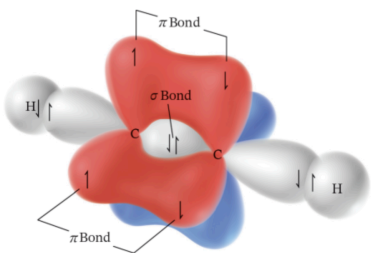





FIGURE 1.11 Constructive interference resulting from the interaction of two electrons.

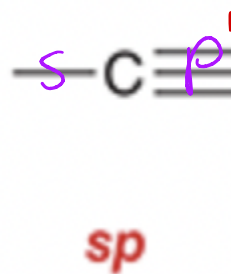
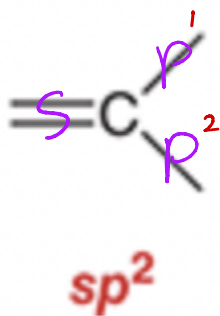
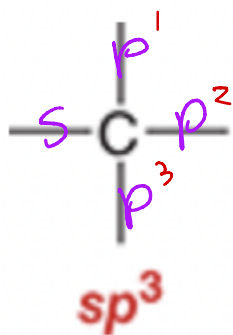


Hybridized Orbitals:

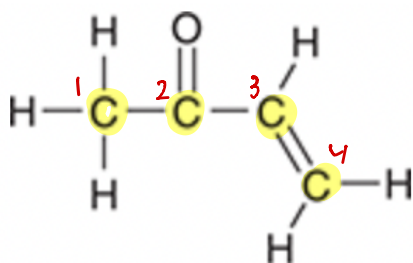
- A combination of pure atomic orbitals
- for this class we only focus on s and p

	Sp ³	Sp ²	Sp
Diagram			
What's Happening	A combination of 1s orbital and 3p orbitals	An overlap of p orbitals forms 1 pi bond	An overlap of p orbitals forms 2 pi bonds
Bond-line			
Geometry	tetrahedral	trigonal planar	linear
Angles	109.5°	120°	180°

Hybridization life hack!!!



Determine the hybridization state of each carbon:



$$C1 = sp^3$$

$$C2 = sp^2$$

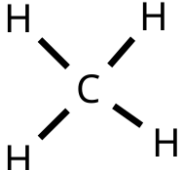
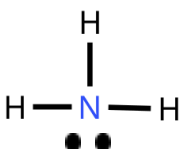

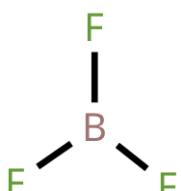

$$C3 = sp^2$$

$$C4 = sp^2$$

VSEPR Theory:

- Valence Shell electron pair repulsion
- Used to predict the geometry of an atom

Common Molecular Shapes:

Compound	Bonding e- pairs	Lone e- pairs	Steric number	Arrangement of e- pairs	Molecular Geometry
	4	0	4	tetrahedral	tetrahedral
	3	1	4	tetrahedral	trigonal pyramidal
	2	2	4	tetrahedral	Bent
	3	0	3	Trigonal planar	trigonal planar
	2	0	2	Linear	Linear

Steric # = # single bond + # lone pairs

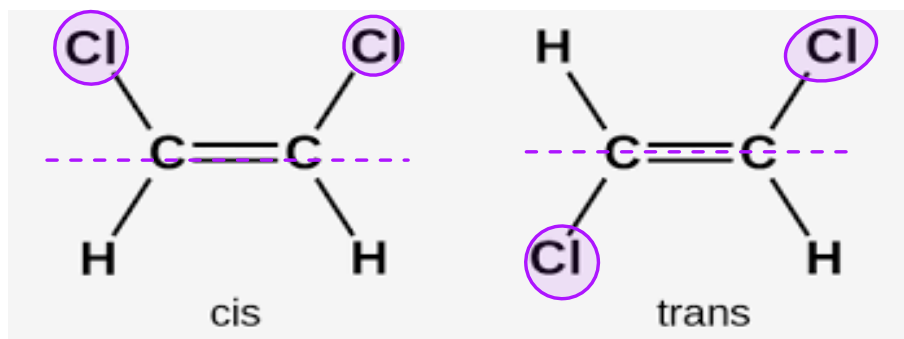
Cis/trans Stereoisomerism:

Cis: *on the same plane*

Trans: *on opposite ends of the plane*

We can think of the molecule as being on a plane and separating this plane evenly either through the molecule itself or through a double/triple bond

Ex:



Restricted Rotation:

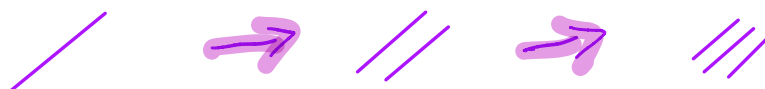
AKA, the properties of a single, double, and triple bond

Order the bonds:

Length

longest

shortest



Energy

high

low



Strength

stronger

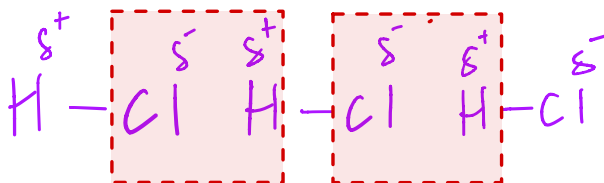
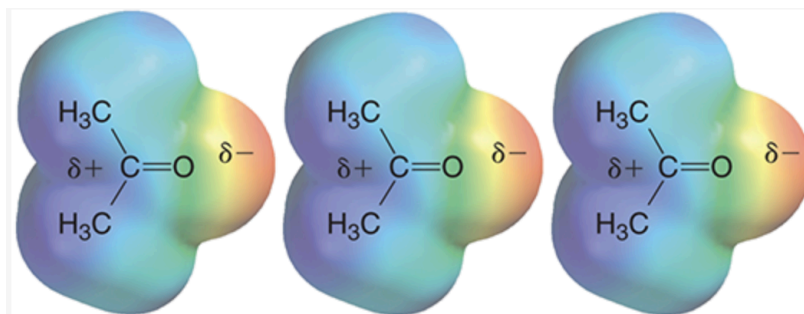
weaker



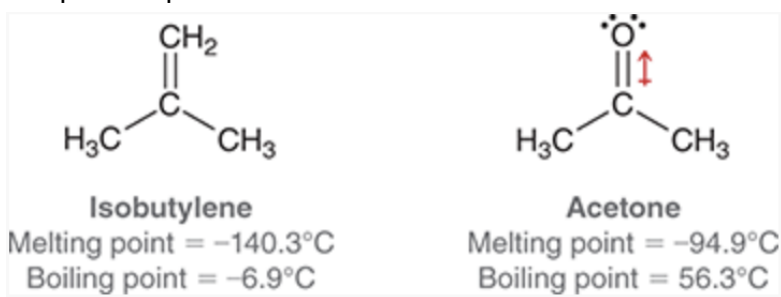
Intermolecular Forces

Dipole-Dipole:

The resulting net attraction between two dipoles



How does this affect bp and mp?



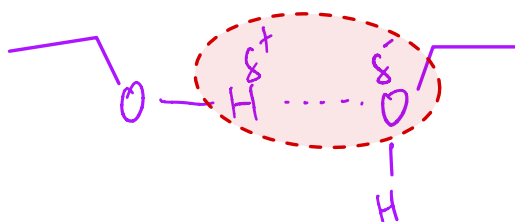
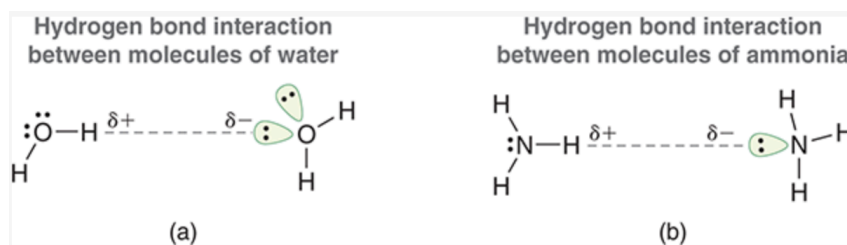
Isobutylene lacks a significant dipole moment, so the mp and bp are much lower compared to Acetone, which has strong net dipole

Hydrogen Bonding:

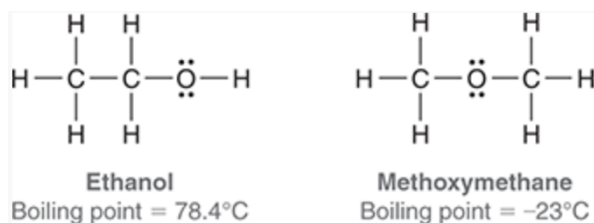
- Not technically a "bond", more like another form of attraction

A hydrogen is connected to an EN atom (O or N)

In most cases, H-bonding takes priority

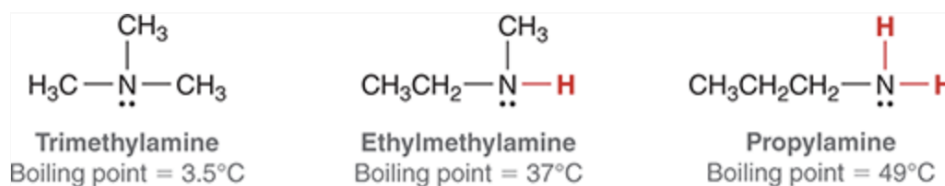


How does this affect bp & mp?



Ethanol has a higher bp because it has a hydrogen bonded to, versus Methoxymethane, which

only has a C-O bond



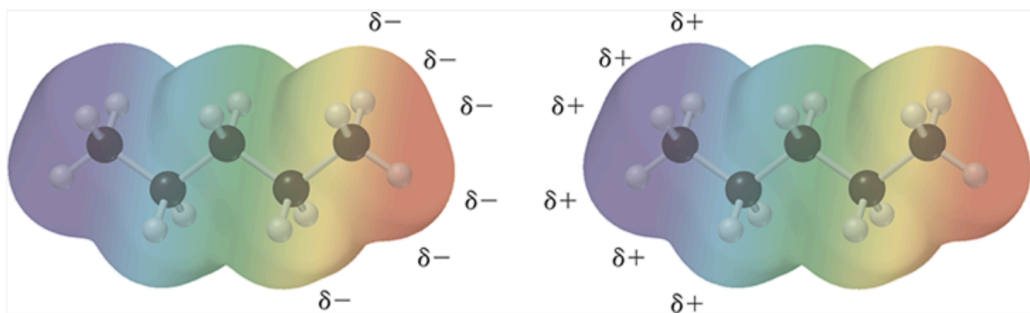
Notice how as more hydrogens are bonded to the Nitrogen atom, the higher the bp gets

London Dispersion Forces:

A consideration of the positive and Negative charges on a whole molecule, rather than the entire atom

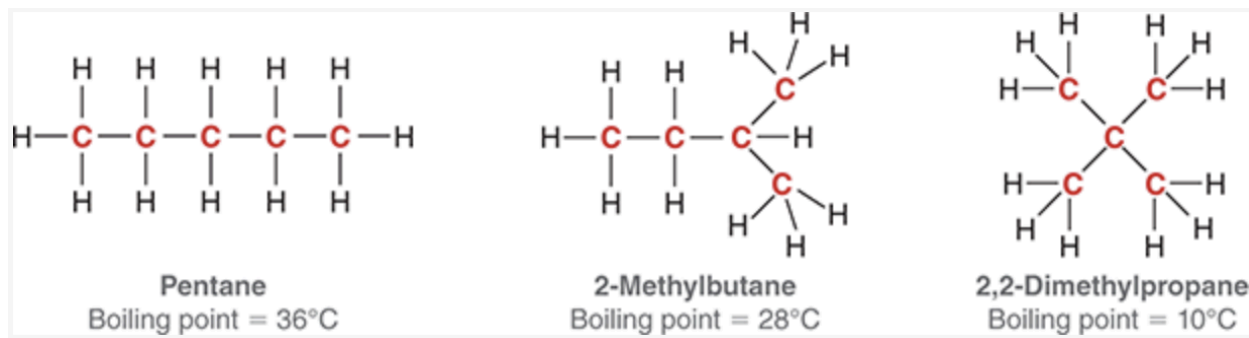
Usually observed in large hydrocarbons

This force is transient, or temporary



How does this affect bp and mp?

The longer the carbon chain, the higher ~~more~~ the bp



The more branching, the lower the bp