Chapter 4 - Hybridization Problems #1

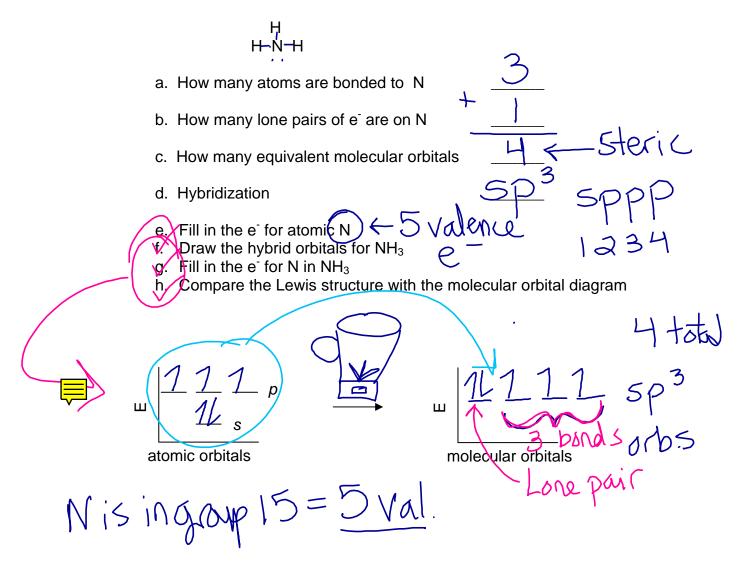
Click on the pink circles to listen to the written portions and get helpful hints! Use the Drawing and

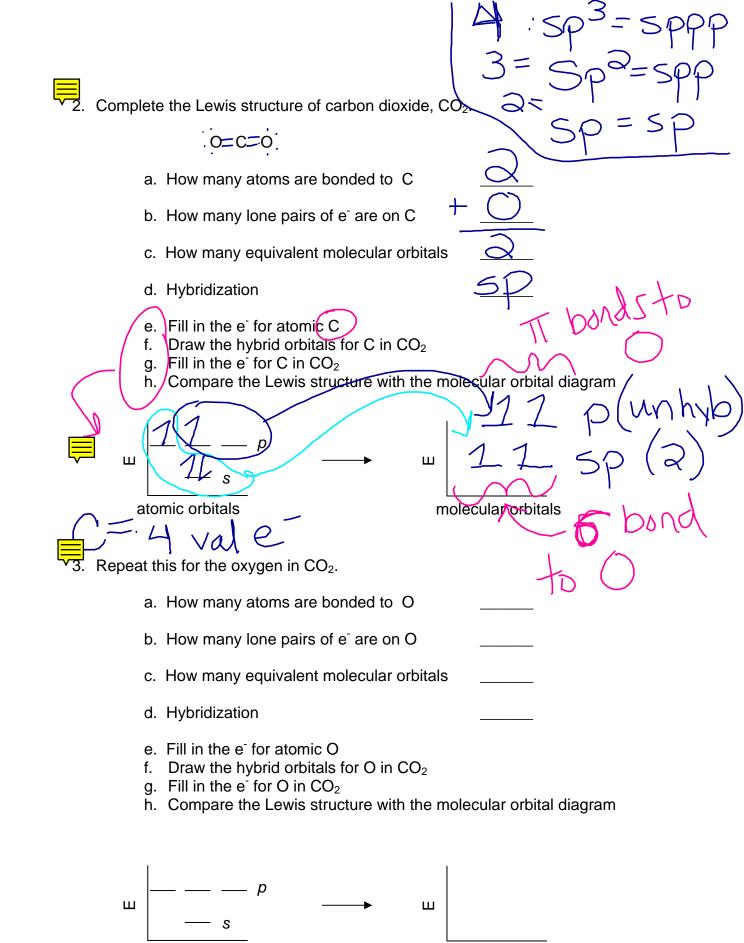
Text box tools to complete the questions!

One model to explain bonding in molecules is called the **valence bond theory**. It is based on the concept of **hybridized molecular orbitals**. Atomic orbitals (s,p,d) are mathematically mixed to form molecular orbitals with defined shapes and energy levels. In organic chemistry, we need only look at the hybridization of the one 2s and three 2p orbitals containing the valence electrons of C,O and N.

When **four** equivalent molecular orbitals are needed, all **four** atomic orbitals are mixed to give sp^3 hybridization. When three equivalent molecular orbitals are needed, three of the atomic orbitals are mixed to give sp^2 hybridization. In this case, one unhybridized *p* atomic orbital remains, with its shape and energy level unchanged. When two equivalent molecular orbitals are needed, only two of the four atomic orbitals are mixed, giving *sp* hybridization. Two unhybridized atomic *p* orbitals remain.

. Lewis structures help to determine how many equivalent molecular orbitals are present. Complete the Lewis structure of ammonia, NH₃.

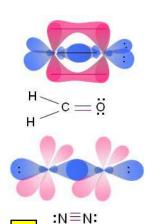




atomic orbitals

molecular orbitals

Bonds made with hybridized orbitals are called σ (sigma) bonds. All single bonds are σ -bonds. Bonds made with unhybridized p orbitals are called π (pi) bonds. In multiple bonds (= or =) one bond is a σ -bond and the others are π -bonds. Sigma bonds result from the end-to-end overlap of hybridized orbitals. The bonding electrons are held directly between the two nuclei. Pi bonds result from the side-to-side overlap of p orbitals. The bonding electrons are held directly between the two nuclei.



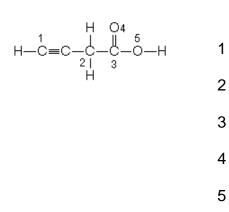
The top figure shows the electronic structure of CH_2O . The C and O are both sp^2 hybridized. Two of the orbitals on O are populated with lone pair e⁻. The bond angle between the hybridized orbitals is **120°** making this a **trigonal planar** structure. The unhybridized *p* orbital is at 90° to the sp^2 orbitals

The bottom figure shows the electronic structure of N₂. Both N are *sp* hybridized, with **180°** between these orbitals. One hybridized orbital forms a σ -bond and a lone pair of e occupies the other. The two *p* orbitals on each N form two π bonds between the atoms.

The four sp^3 orbitals lie at **109.5°** to each other and form a **tetrahedron**.

For the compound below, determine the hybridization, bond angles and lone pair e⁻ at each numbered atom.

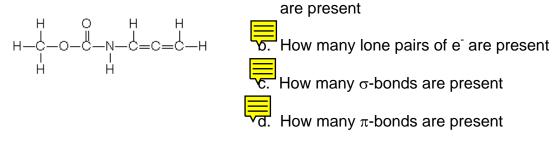
hybridization



bond angle

lone pair e

5. For the compound below determine a. How many sp^3 , sp^2 and sp hybridized atoms



It is often possible to draw more than one Lewis structure for a compound. For example, N_2O can be drawn as:

:N::N:::O: N:::N::Ö: :N:::N:Ö:

To determine which structure is the most reasonable, the **formal charge** on each of the atoms in the molecules can determined. The formal charge is an indication of the electron density at the atoms in a particular structure. It is determined as:

formal charge = valence e^{-} - lone pair e^{-} - $\frac{1}{2}$ shared e^{-} Nitrogen is in Group V, and has 5 valence electrons. Oxygen is in Group VI, with 6 valence electrons.

Assign formal charges to all of the atoms in the three resonance structures of N₂O.
a. : N: N: :: O: N → N → O
b. N: :: N: : Ö
c. : N: :: N: : Ö:

- d. Based on the chemical nature of these elements, which is the worst structure and why?
- e. Which is the best structure, and why?

h	e atom	ic Lewis structures of C, N and O are $\cdot \dot{C} \cdot \cdot \dot{N} \cdot \cdot \dot{O} \cdot$
	7.	What is the most stable bonding pattern for <u>bonds</u> <u>lone pair e</u>
		a. C
		b. N

c. O