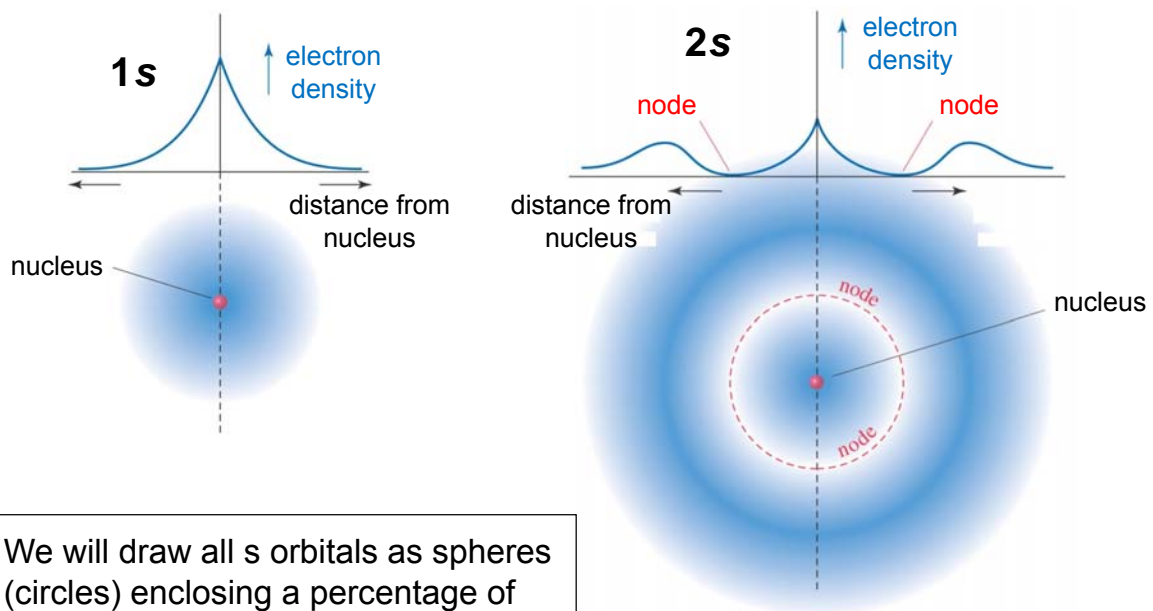


# Electrons and Bonds in Space

- Lewis-dot structures are great for *accounting*, but don't say much about three-dimensional arrangement of atoms and bonds
- For that, need *molecular orbitals*, built from atomic orbitals.

## 1s, 2s Atomic Orbitals



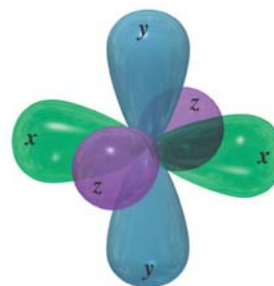
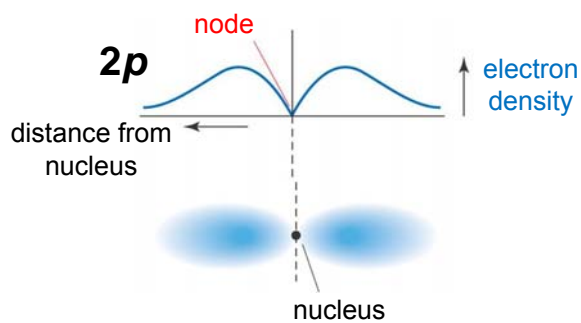
We will draw all s orbitals as spheres (circles) enclosing a percentage of electron density.

H[1s]:



Reminder:  $n - l = \# \text{ of nodes}$

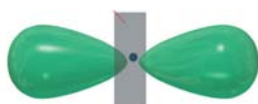
## 2p Atomic Orbitals



$2p_x$ ,  $2p_y$ , and  $2p_z$  orbitals

Again, draw as solid lobes enclosing a percentage of electron density.

C[ $2p_x$ ]:



## Molecular Orbitals

- Bonds, electrons between atoms in molecules are described by molecular orbitals.
- These are built from atomic orbitals via *orbital mixing*.

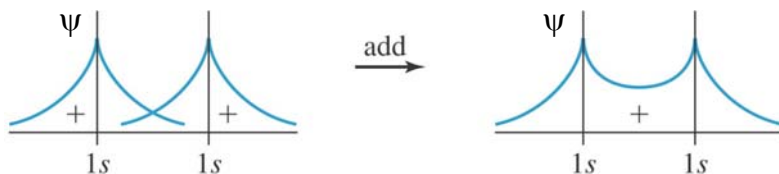
### Rules of orbital mixing:

- Product (mixed) orbitals look like constructive and destructive combinations of starting orbitals, with some distortions.
- You end with the same number of orbitals you started with.
- Degree of mixing depends on orbital overlap, match between orbital energies.

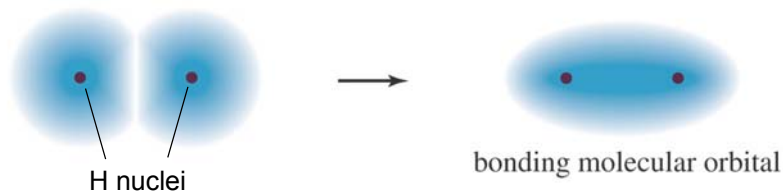
# The Single Bond in H<sub>2</sub>

Constructive mixing yields a "bonding" orbital.

$\psi = (e^- \text{ density}) \cdot \text{phase } (+/-)$



Start with linear combination of atomic orbitals (LCAO)



represented by:

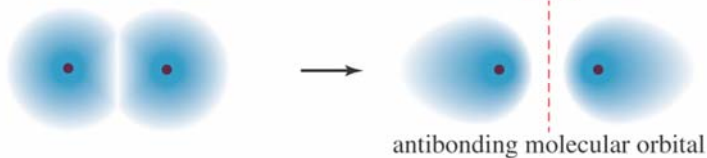
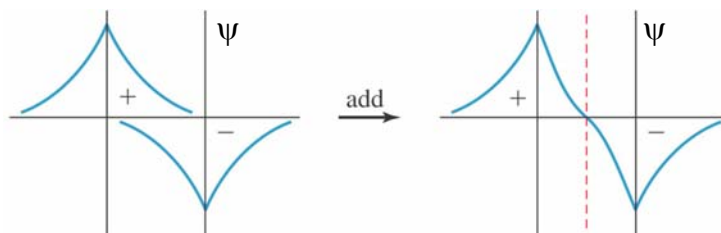


...but we need to make another one...

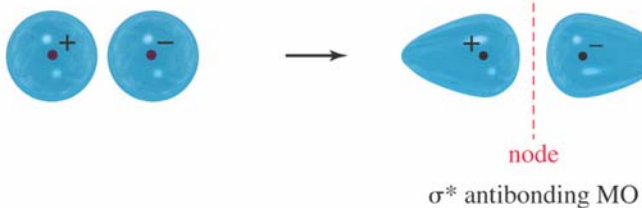
# The Single Bond in H<sub>2</sub>

Destructive mixing yields an "anti-bonding" orbital.

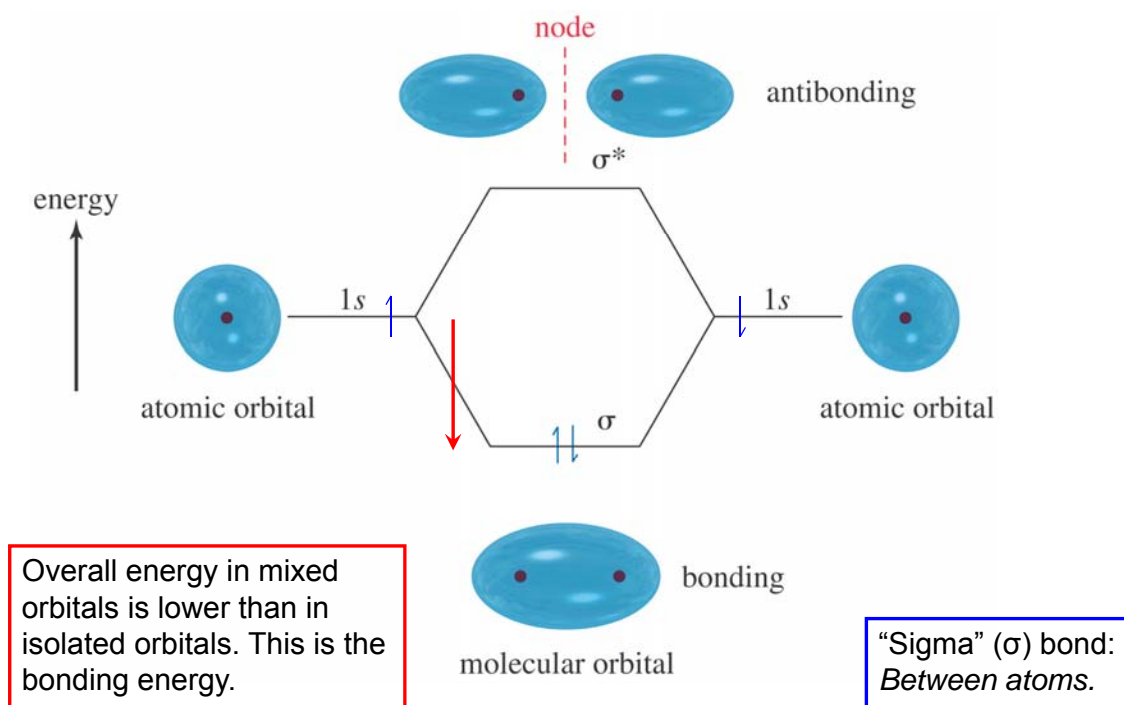
$\psi = (e^- \text{ density}) \cdot \text{phase } (+/-)$



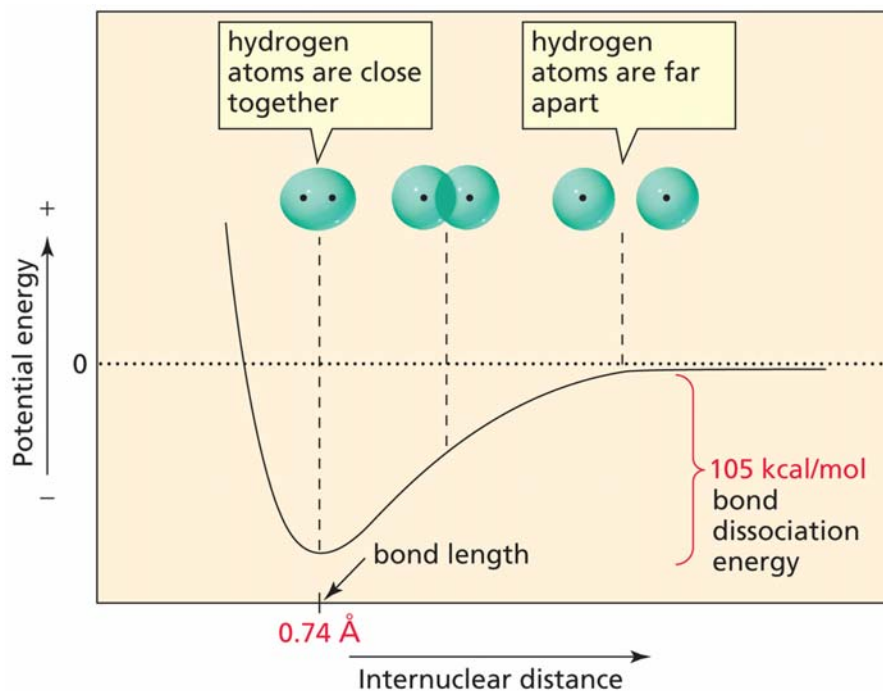
represented by:



# Mixing s Orbitals

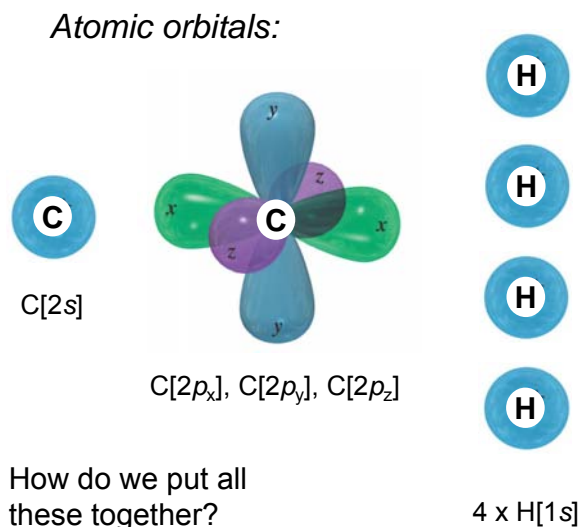
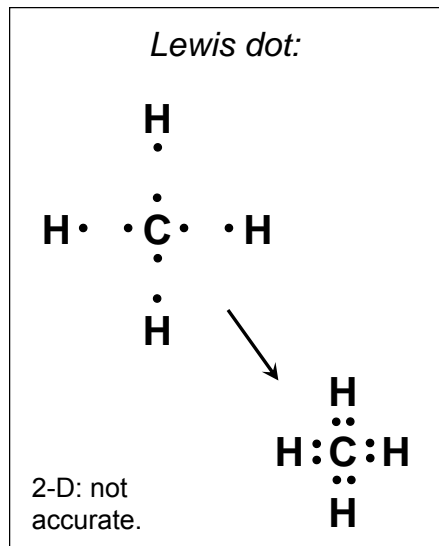


# Orbital Mixing is Responsible for Bond Strength



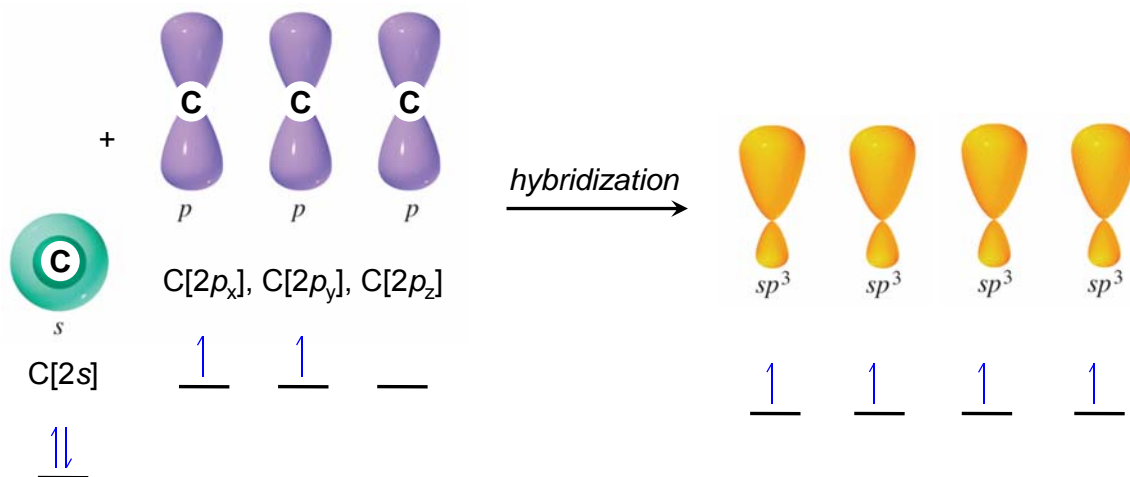
# How Do We Use Orbital Mixing to Describe Molecular Structures?

What does methane ( $\text{CH}_4$ ) look like?



## Hybrid Atomic Orbitals

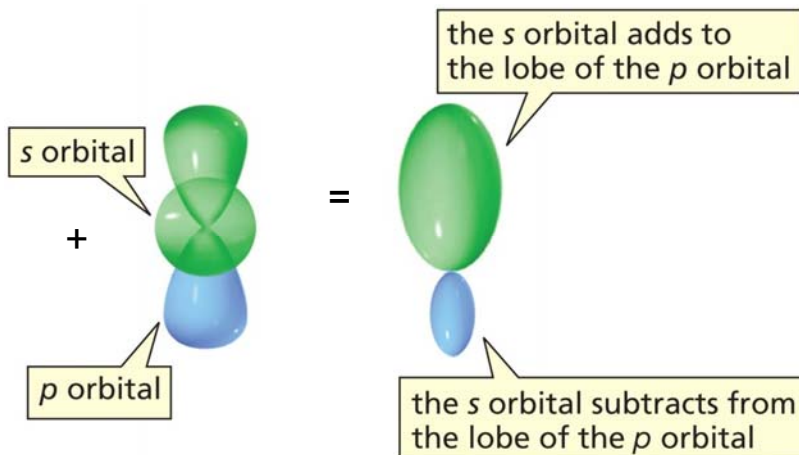
First: “Hybrid orbitals” are formed from  $s$  and  $p$  orbitals (on one atom) to create all  $\sigma$  bonds and lone pairs.



- Called “ $sp^3$ ” because they come from one  $s$  and three  $p$ 's;
- Four of them, because we started with four atomic orbitals to make them.

# Hybrid Atomic Orbitals

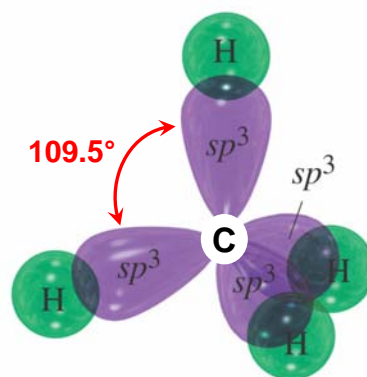
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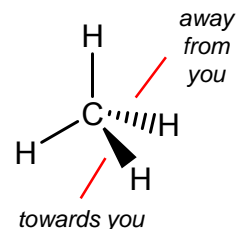
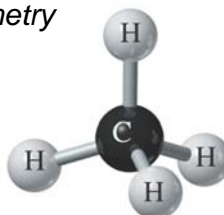
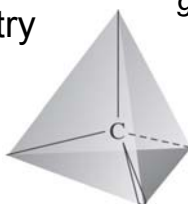
## Valence-Shell Electron Pair Repulsion (VSEPR) Theory Determines Geometry

Methane ( $\text{CH}_4$ ):

- Each  $\sigma$  bonding and non-bonding electron pair repels each other (by electron-electron repulsion);
- These electron pairs organize themselves to maximize distance from each other;
- For  $\text{CH}_4$ , that geometry is tetrahedral.



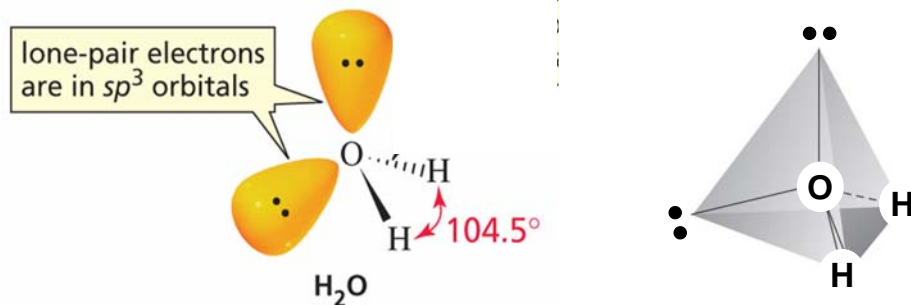
*tetrahedral geometry*



# Valence-Shell Electron Pair Repulsion (VSEPR) Theory Determines Geometry

$\sigma$  bonds (but not  $\pi$  bonds), electron pairs repel each other.

H<sub>2</sub>O:



Actually, lone pairs repel *slightly* better than  $\sigma$  bonds.  
So, H-O-H angle is  $< 109.5^\circ$ .