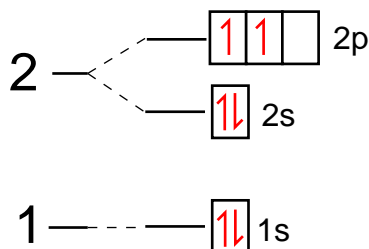
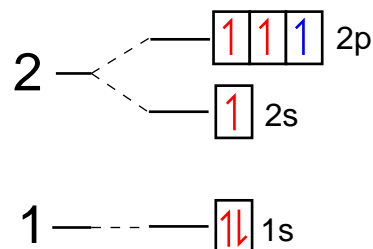


## HYBRIDISATION IN CARBON

What happens?



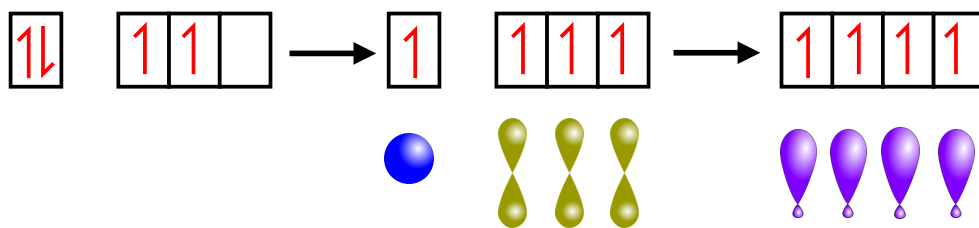
The electronic configuration of a carbon atom is  $1s^2 2s^2 2p^2$



If you provide energy you can promote (lift) one of the s electrons into a p orbital. The configuration is now  $1s^2 2s^1 2p^3$

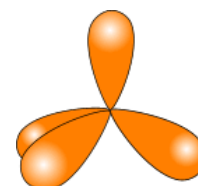
The process is **favourable** because of the arrangement of electrons; four unpaired means **less repulsion** and therefore **more stability**.

$sp^3$

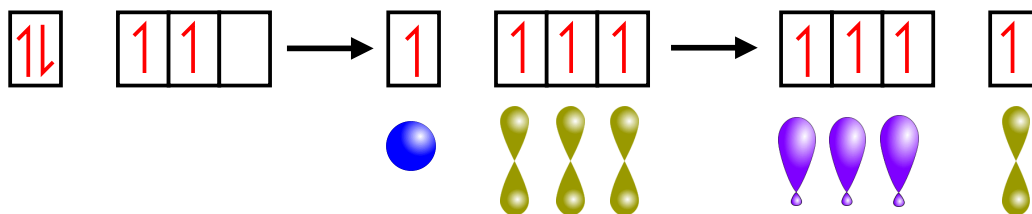


The four orbitals (an s and three p's) combine or HYBRIDISE to give four new orbitals. All four orbitals are equivalent.

In ALKANES, the four  $sp^3$  orbitals repel each other into a tetrahedral arrangement.

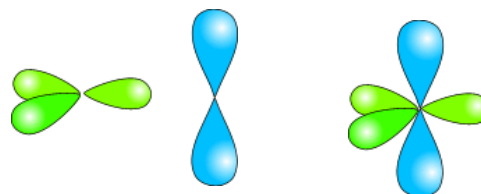


$sp^2$



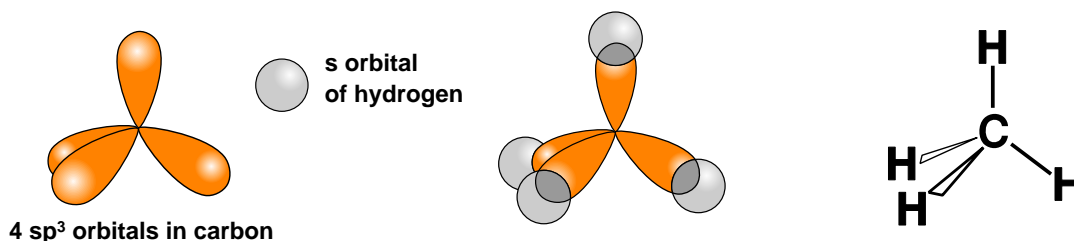
Only three orbitals (an s and two p's) HYBRIDISE to give three new orbitals. All three orbitals are equivalent. The remaining 2p orbital is unchanged.

In ALKENES, the three  $sp^2$  orbitals repel each other into a planar arrangement and the 2p orbital lies at right angles to them



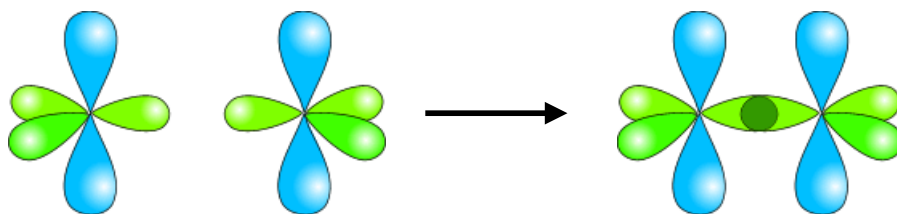
### Bond formation in alkanes

- Covalent bonds are formed by overlap of orbitals
- Each  $sp^3$  orbital from carbon overlaps with a hydrogen s orbital
- The resulting bond C-C bond is called a SIGMA ( $\sigma$ ) bond
- The four  $\sigma$  bonds repel each other to give a tetrahedral shape

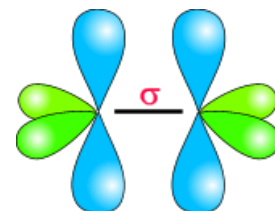


### Bond formation in alkenes

- An  $sp^2$  orbital from each carbon overlaps to form a single C-C bond

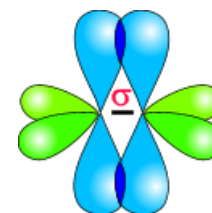


The resulting bond is called a SIGMA ( $\sigma$ ) bond.

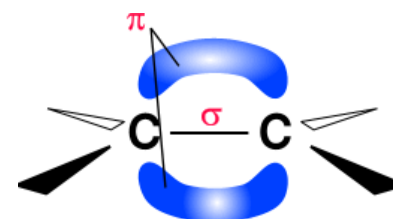


The two 2p orbitals also overlap to form a second bond. This is known as a PI ( $\pi$ ) bond.

For maximum overlap and hence the strongest bond, the 2p orbitals are in line.



This gives rise to the **PLANAR** arrangement around C=C bonds.  
This gives **RESTRICTED ROTATION**.



To complete the structure of ethene, the remaining  $sp^2$  orbitals overlap with four hydrogen 1s orbitals to form four C-H bonds.

