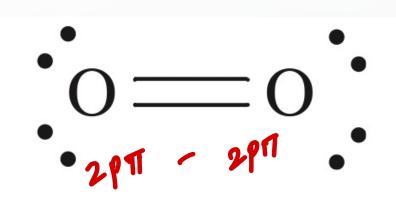
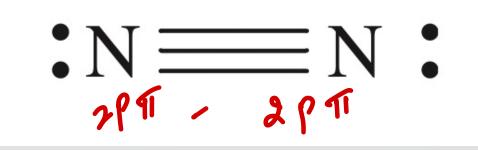
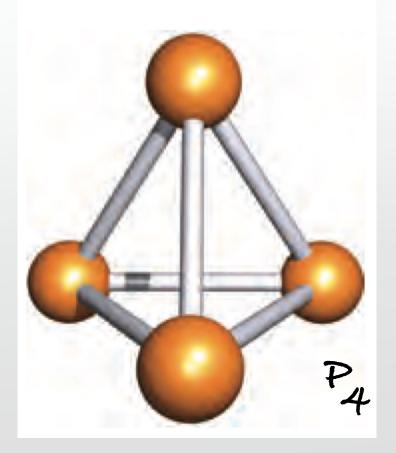
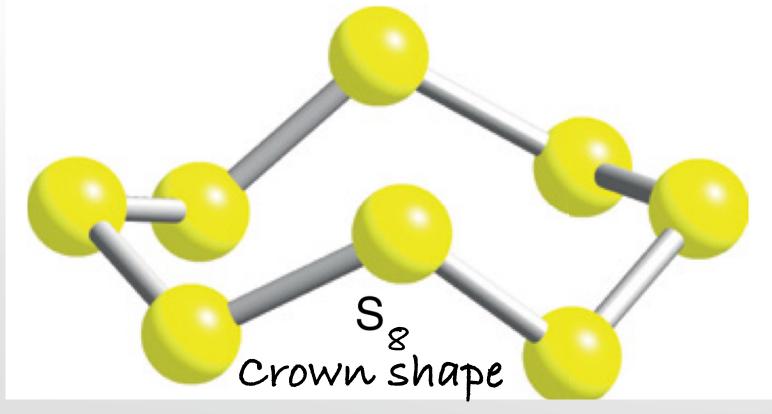


(Q). why oxygen (O2) and Nitrogen (N2) exist in diatomic from while Phosphorus (P_4) and sulphur (S_8) exist in poly atomic form.

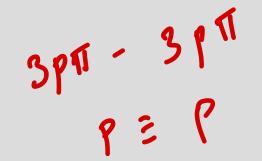


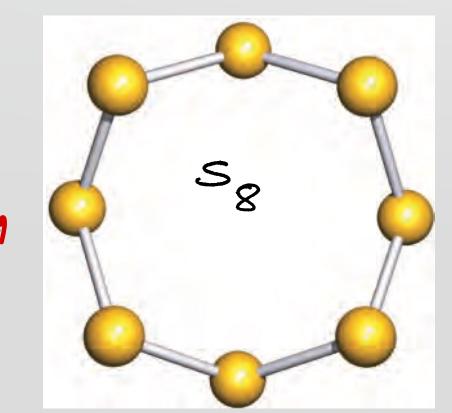




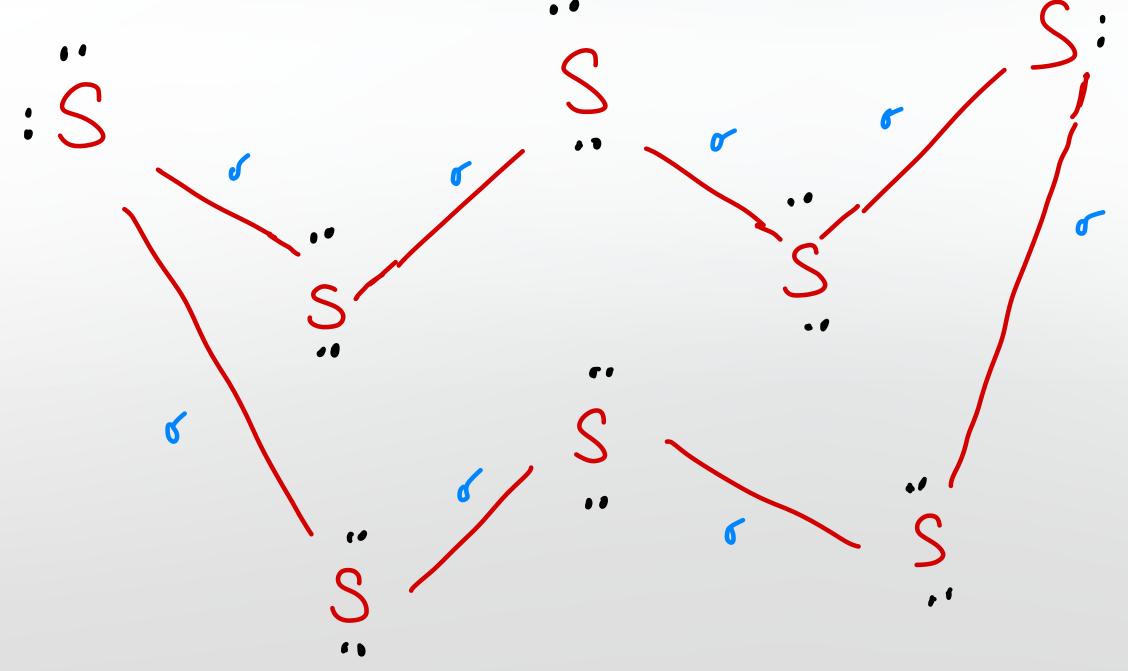


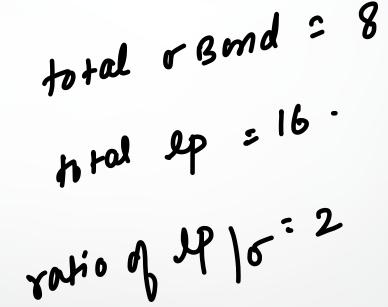
Ans: increases strength of P-P sigma bond in compare to strength of P-P Pi bond (musily) from















(Q) Why Co2 donot form giant structure while from Sio2 form giant structure although carbon ans silicon belongs to same family.?

$$: O = C = O:$$

$$2PII - 2PII \quad Bond \quad formed$$

Ans:)

$$:0 = Si = 0:$$
 | very weak
 $2p\pi - 3p\pi$ | 3 and formed

- 1. tendency of polymerisation decreases with increase in strength of pie bond.
- 2. weak pie bond tends to break them and convert into more stable Structure (polymer)



$$\ddot{o}=s\dot{i}=\ddot{o}$$
 \longrightarrow $\dot{o}=s\dot{i}-o \longrightarrow$ $-\ddot{o}-s\dot{i}-\ddot{o}-$

Sío2 form giant covalent network.

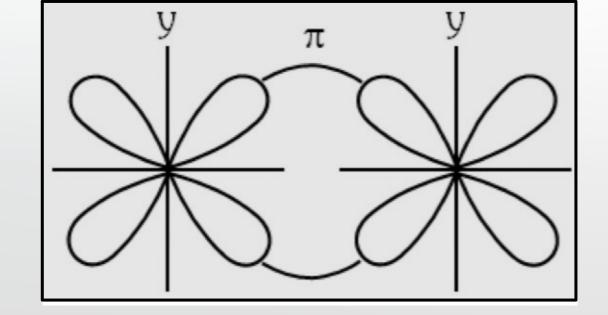
All giant molecule are covalent solids. E.g Sio2 Dry ice (Co2) molecular solid: held by vanderwaals attraction.

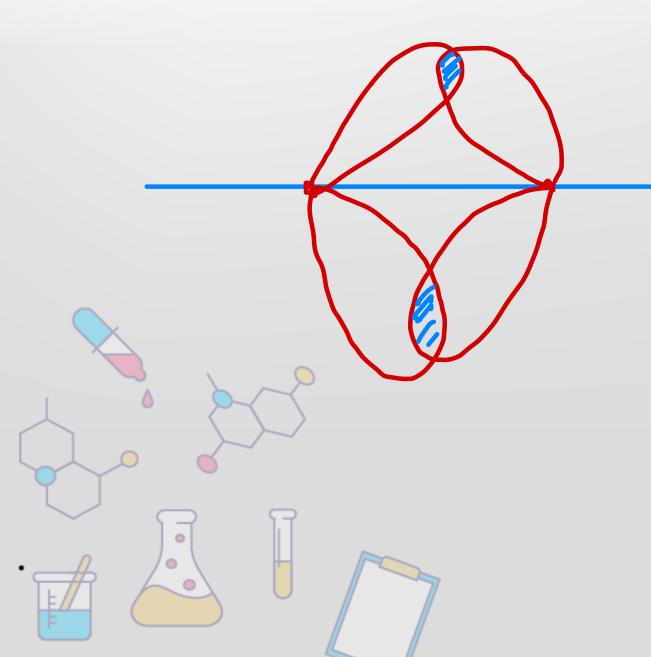


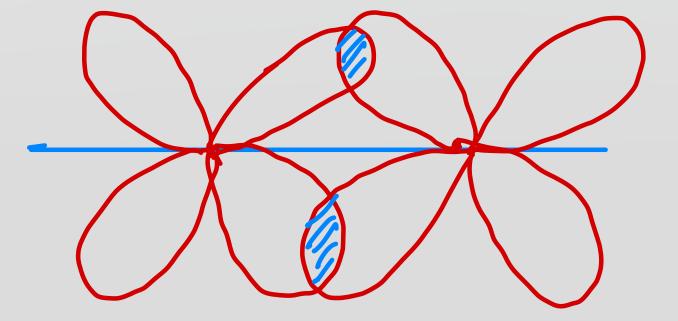


Compare bond strength

**since d has inclined lobe it can form pie bond easily



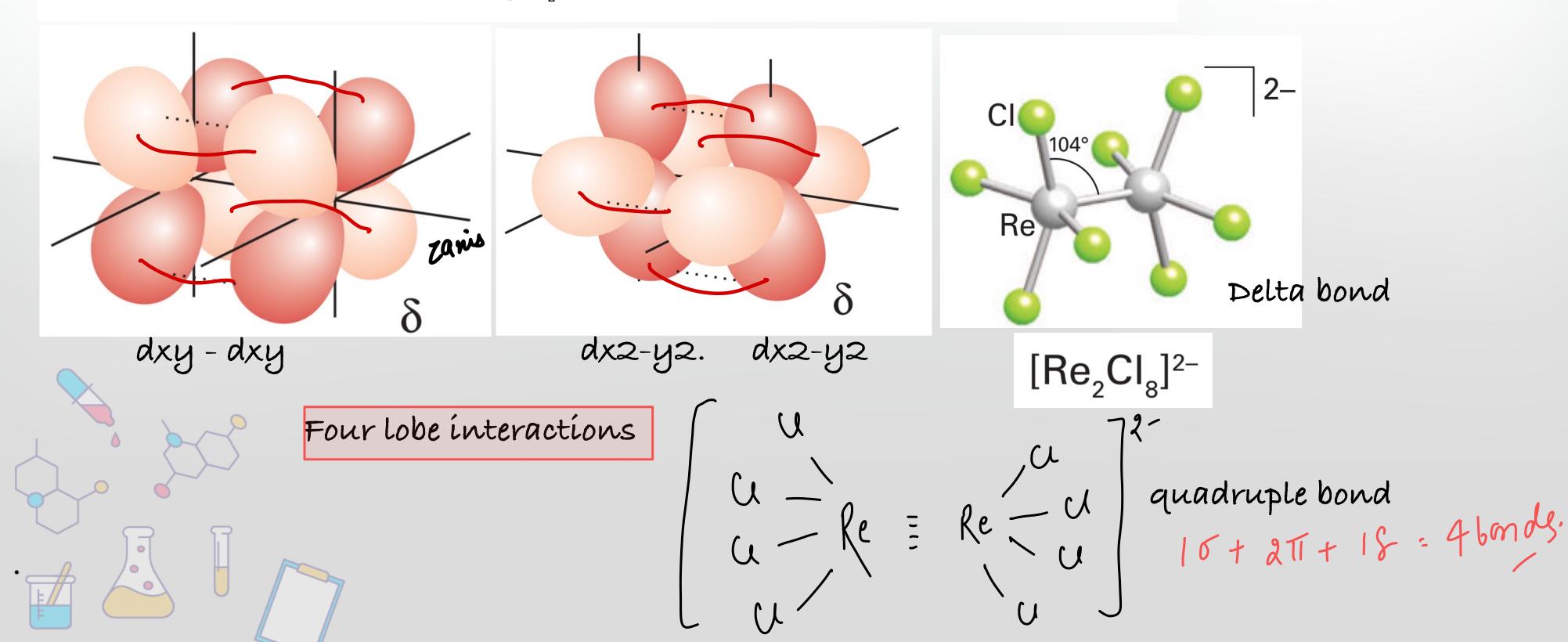






Delta. (8) Bond

delta (δ) bond: are the covalent bonds where four lobes of d-orbital of one atom overlap with four lobes of the similar d-orbital of other atom. Except d_z^2 all d orbitals form δ bond.





Diffenence in σ and π bonds :	
Sigma (σ) bond	Pi (π) bond
It results from the end to end overlapping of two s orbitals or two p—orbitals or one s and one p—orbital.	It result from the sidewise (lateral) overlapping of two p orbitals.
Its bonded orbital consists of a single electron cloud symmetrical about internuclear Axis	Its bonded orbital consists of two electron clouds one above and the other below the plane of symimetry
3. Strong	Weak
4. Bond energy 80 Kcals	65 Kcals Approximately
5. More stable	Less Stable
6. Less reactive	More reactive
7. Can independently exis	Always exists along with a σ bond hybridsation

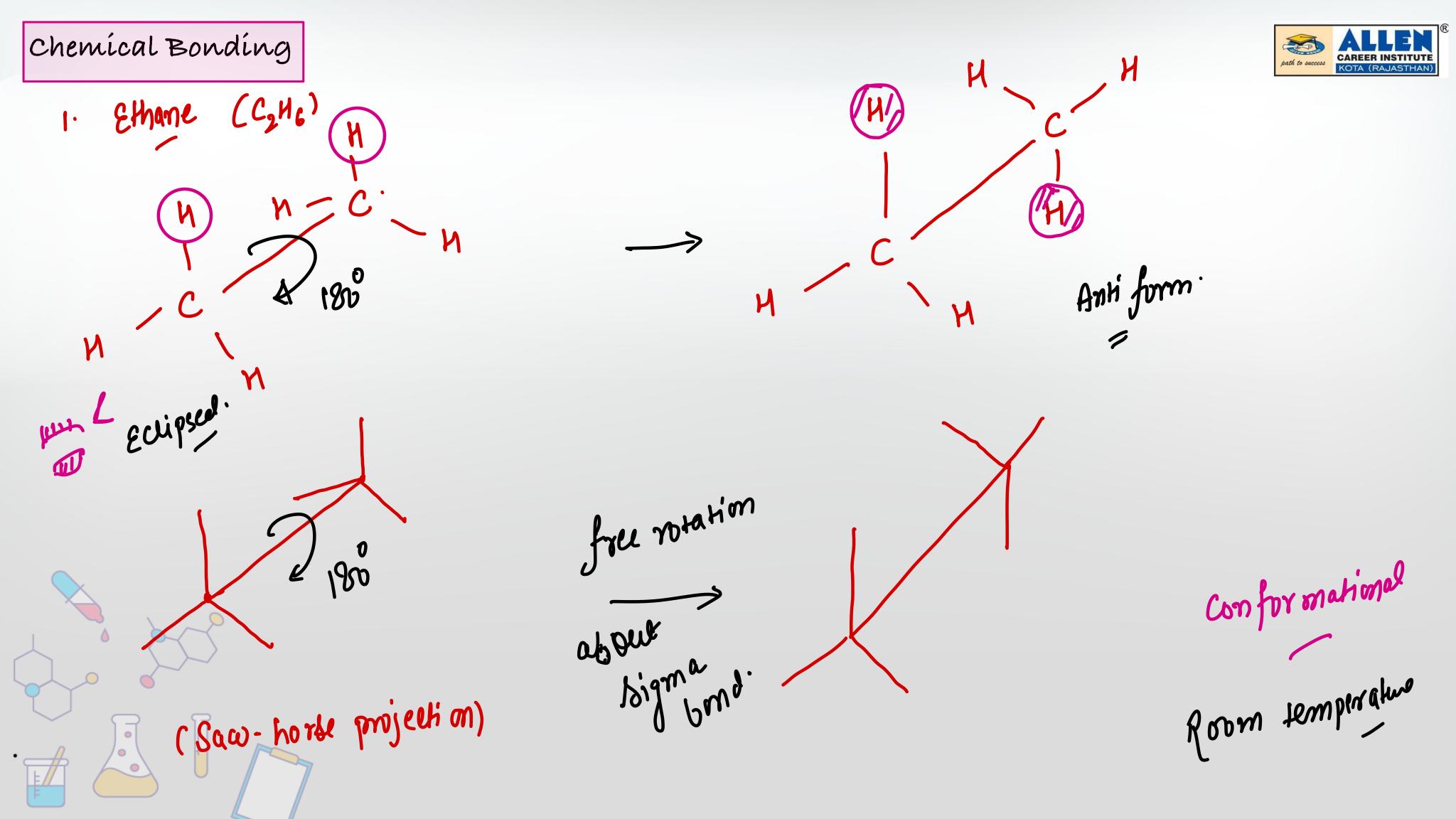
Alkame C2H6 Alkame C2H4.



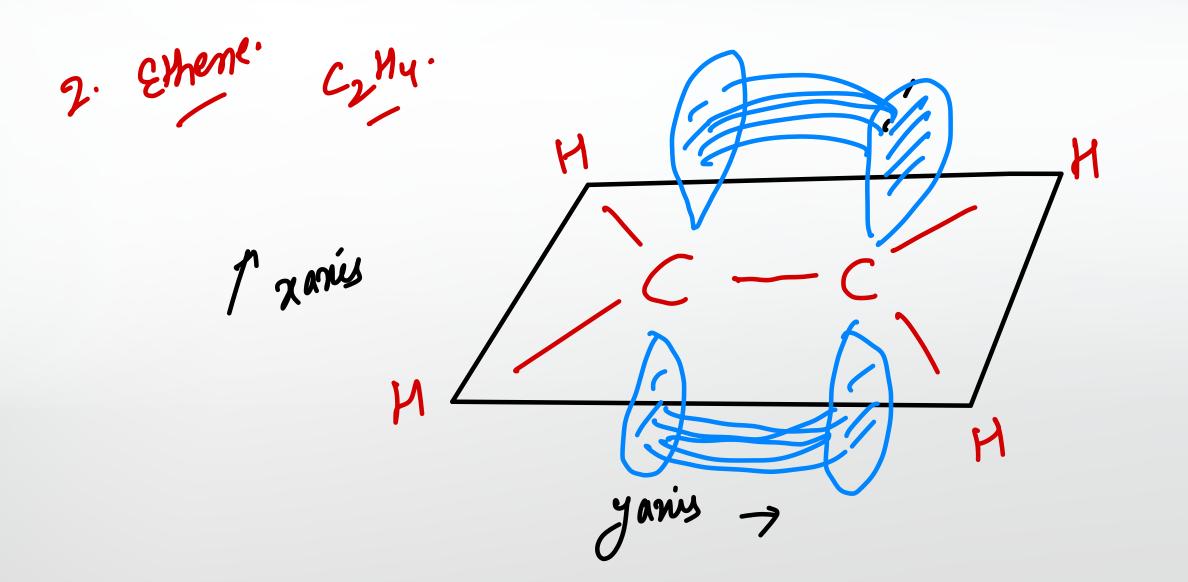


8. Hybridization depends upon σ bond	doesn't depend on σ bond
9. The groups or atoms can undergo bond rotation about single sigma (σ) bonds	Due to resistance to rotation around the π bond the groups attached to it are not free to rotate.
10. The σ electrons are referred as localised	in π bond the electrons are held less firmly bond thus can be easily dissociated or polarised by an external charge and hence the π e are referred as mobile electrons.
11. Shape of the molecule is determined by the σ bonds present in the molecule	$\boldsymbol{\pi}$ bonding does not affect the shape of the molecule









Resmicted rotation about about





Factor affecting bond strength.

1. On the basis of size of overlapping orbital. (n)



2.



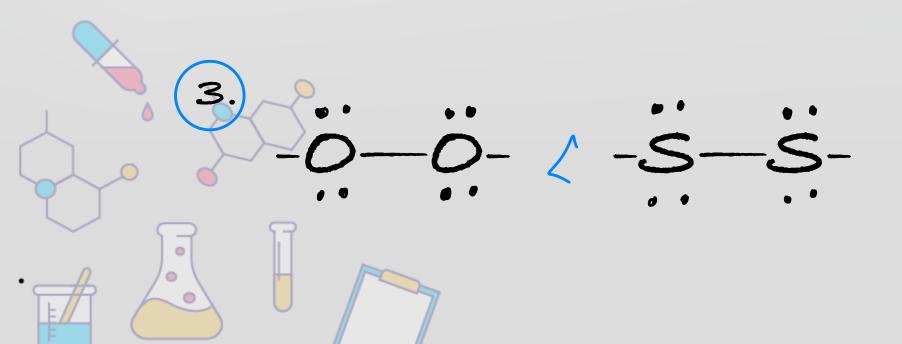


2. On the basis of lp-lp repulsion

Bond strength
$$\propto \frac{1}{|p-p|}$$

1.)
$$-C-C-$$
 > $-N--N-$ > $-O--O-$ On the basis of number of lone pair

On the basis of orbital in which lone pair is present





Experimentally bond dissociation energy order in Halogen is found to be





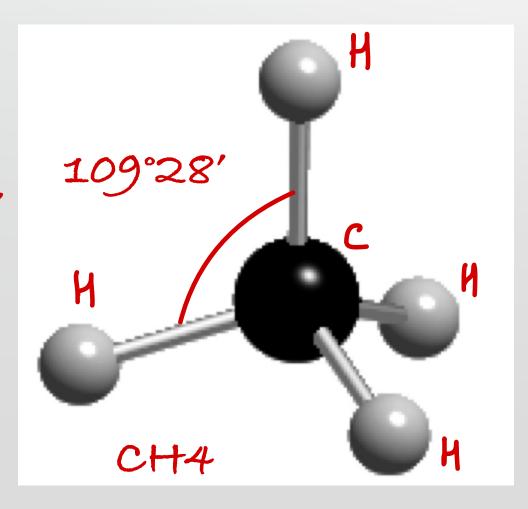
Draw backs of valence bond theory (VBT)

* valence bond theory fails to explain the structure and bonding of molecules which are poly atomic. (molecule having more than two atom.)

Let us consider the CH4 molecule.

For the CH4 molecule following observations are made.

- 1. All four 'C-H' bonds are identical
- 2. All bond are of equal strength.
- 3. All bond angle 'H-C-H' are equal and equal to 109°28'







According to VBT the CH4 molecule can be explained as follows:

