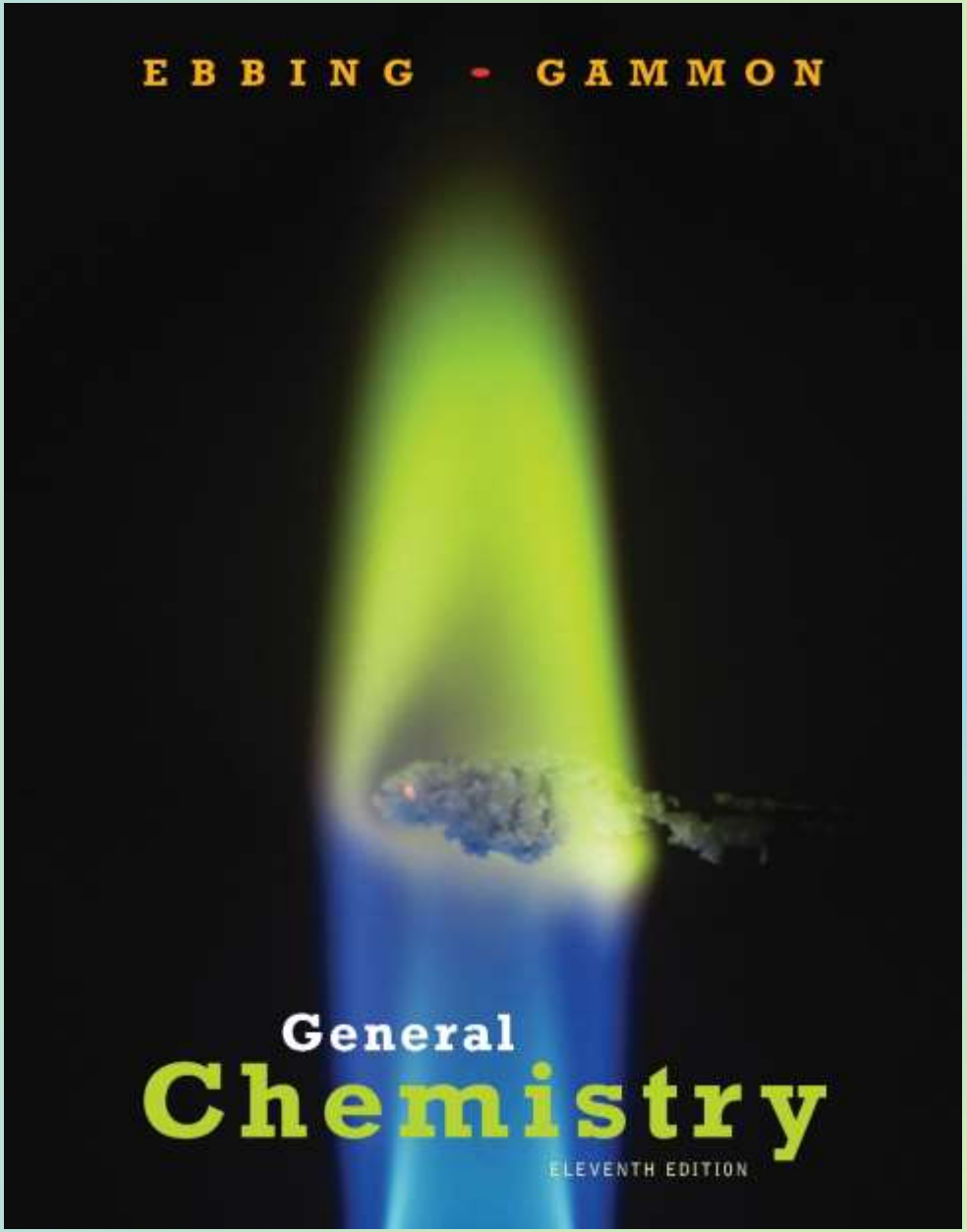
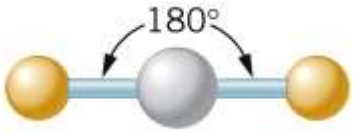
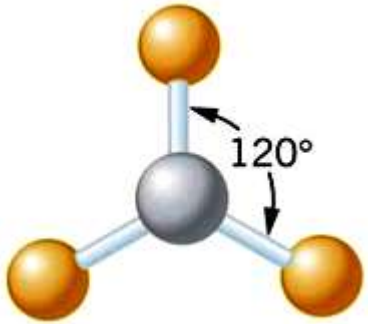
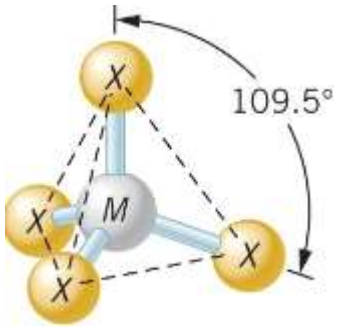
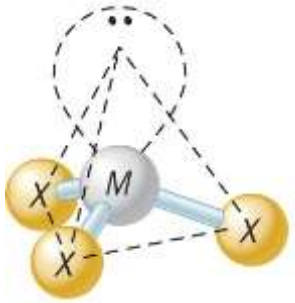
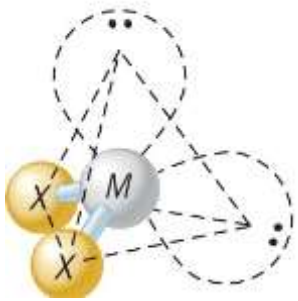
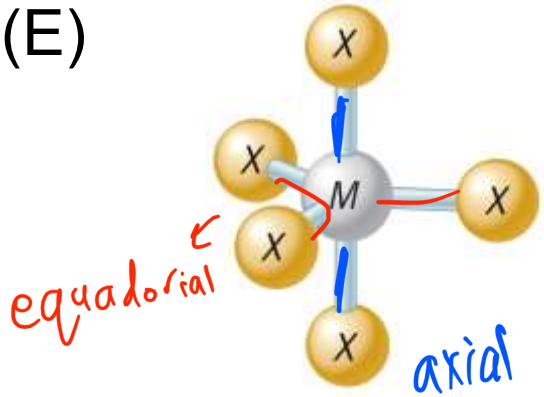
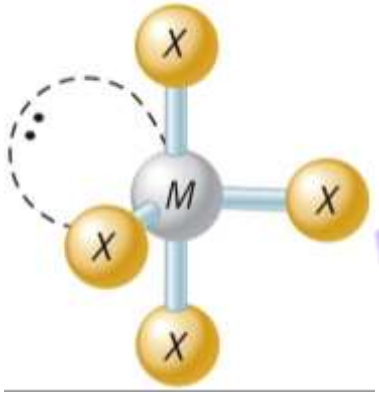


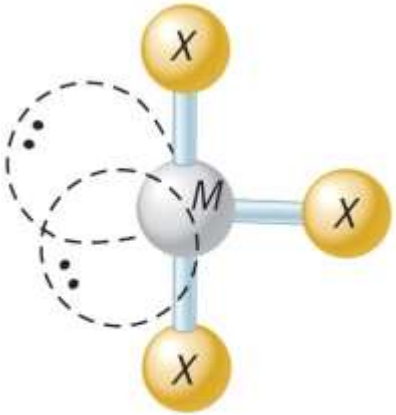
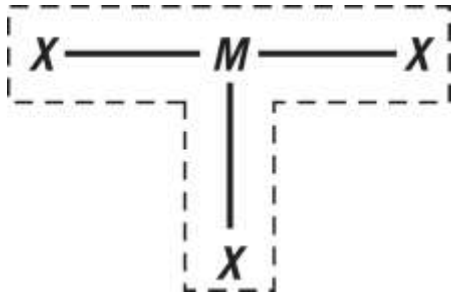
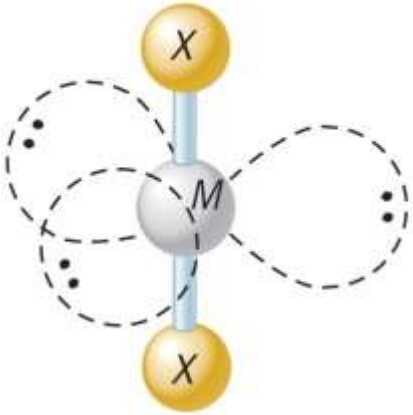
Molecular Geometry and Chemical Bonding Theory

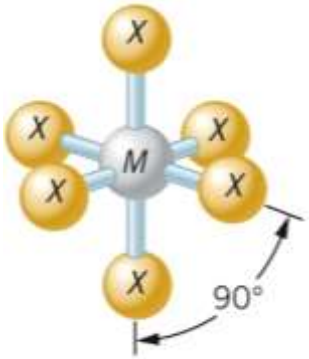
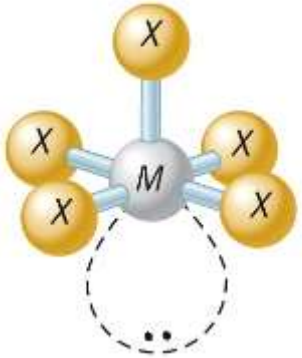
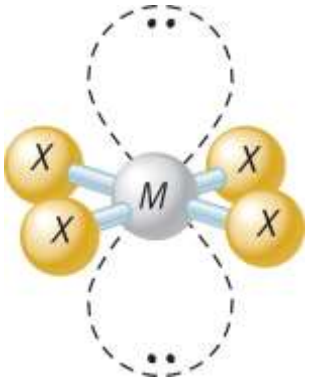
10.1 Valence-Shell Electron-Pair Repulsion (VSEPR) Model

Number of Bonding Pairs	Number of Non-bonding Pairs (E)	Molecular Geometry	Molecular Shape
2	0		AX_2 Linear
3	0		AX_3 Trigonal Planar (e.g., BCl_3) All bond angles 120°
2	1		AX_2E Bent (e.g., SO_2) Bond <120°

Number of Bonding Pairs	Number of Nonbonding Pairs (E)	Molecular Geometry	Molecular Shape
4	0		**AX_4 Tetrahedral** (**e.g.,** CH_4) All bond angles 109.5°
3	1		**AX_3E Trigonal pyramidal** (**e.g.,** NH_3) Bond angle **less than** 109.5°
2	2		**AX_2E_2 bent** (**e.g.,** H_2O) Bond angle less than109.5°

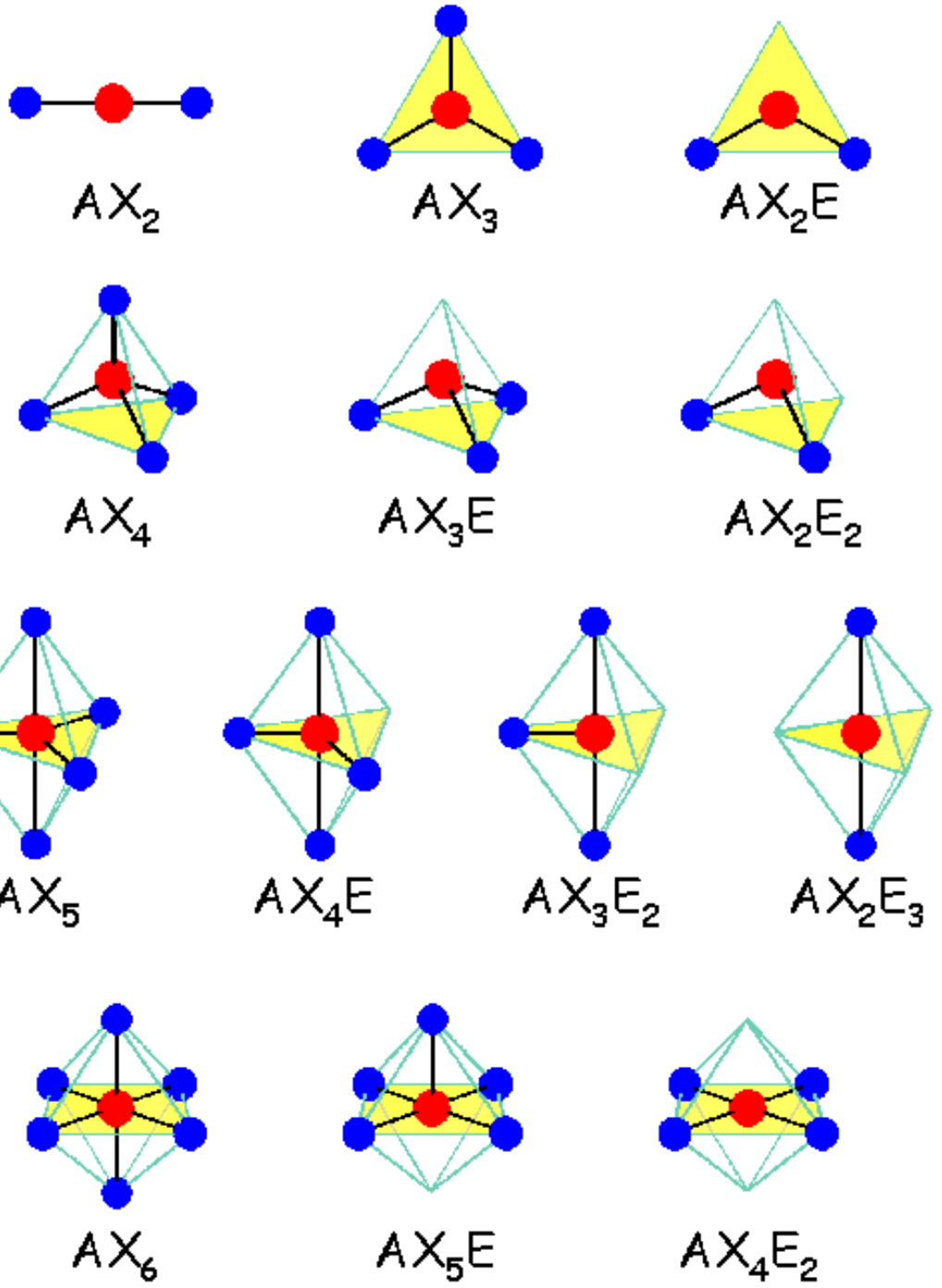
Number of Bonding Pairs	Number of Nonbonding Pairs (E)	Molecular Geometry	Molecular Shape
5	0		**AX_5** **Trigonal bipyramid** (**e.g.,** PF_5) **axial-equatorial** bond ax-eq angles 90° eq-eq 120° ax-ax 180°
4	1		**AX_4E** **Distorted Tetrahedron or Seesaw** (**e.g.,** SF_4) ax-eq bond angles < 90° ax-ax 180° eq-eq < 120°

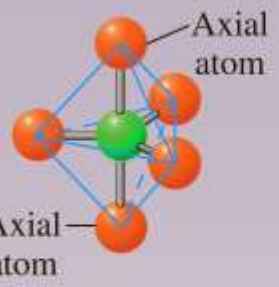
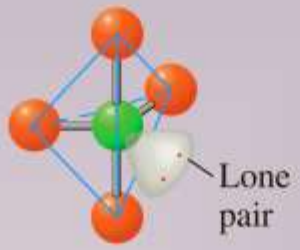
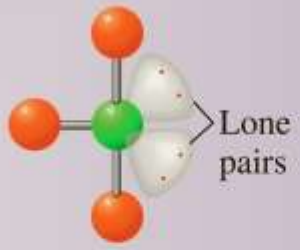
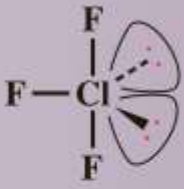
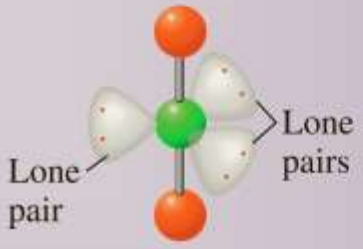
Number of Bonding Pairs	Number of Nonbonding Pairs (E)	Molecular Geometry	Molecular Shape
3	2		**AX_3E_2 T-shape** (**e.g.,** ClF_3) Bond angles 90° 
2	3		**AX_2E_3 Linear** (**e.g.,** I_3^-) Bond angles 180°

Number of Bonding Pairs	Number of Nonbonding Pairs (E)	Molecular Geometry	Molecular Shape
6	0		**AX_6 Octahedral** (**e.g.,** SF_6) Bond angles 180°,90°
5	1		**AX_5E Square Pyramidal** (**e.g.,** BrF_5) Bond angles 90°
4	2		**AX_4E_2 Square planar** (**e.g.,** XeF_4) Bond angles 90°, 180°

Summary of Molecular Geometries

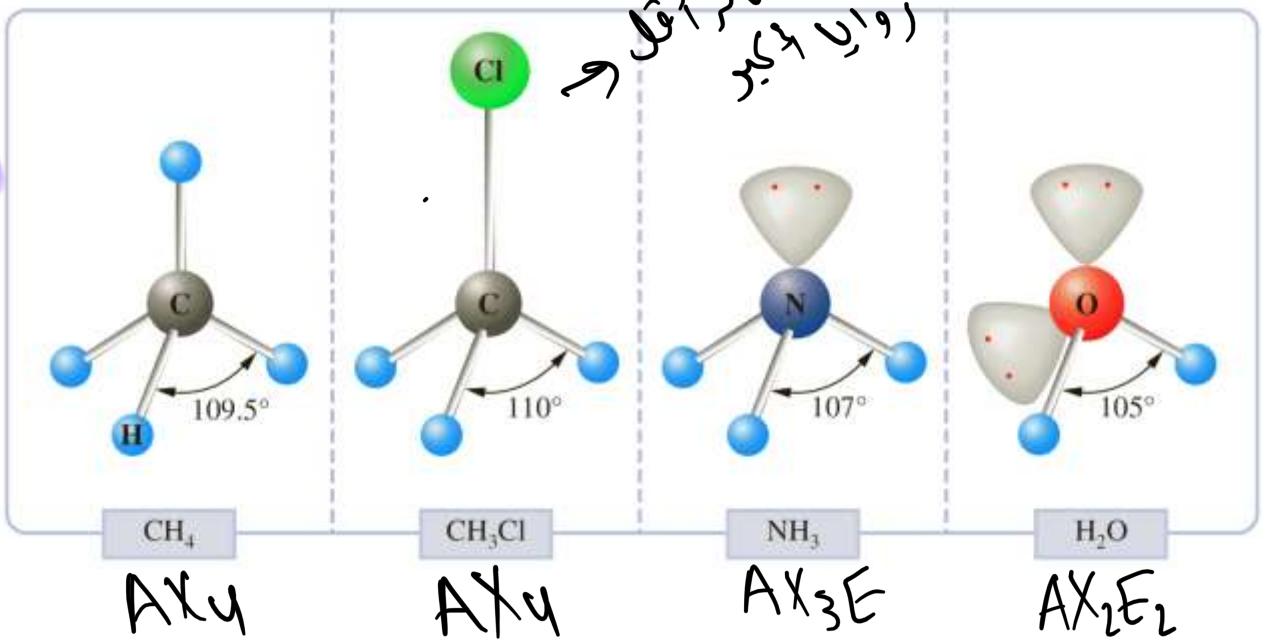
- ✓ **VSEPR is based on minimizing electron repulsion in the molecule**
- ✓ The direction in space of the bonding pairs gives the molecular geometry



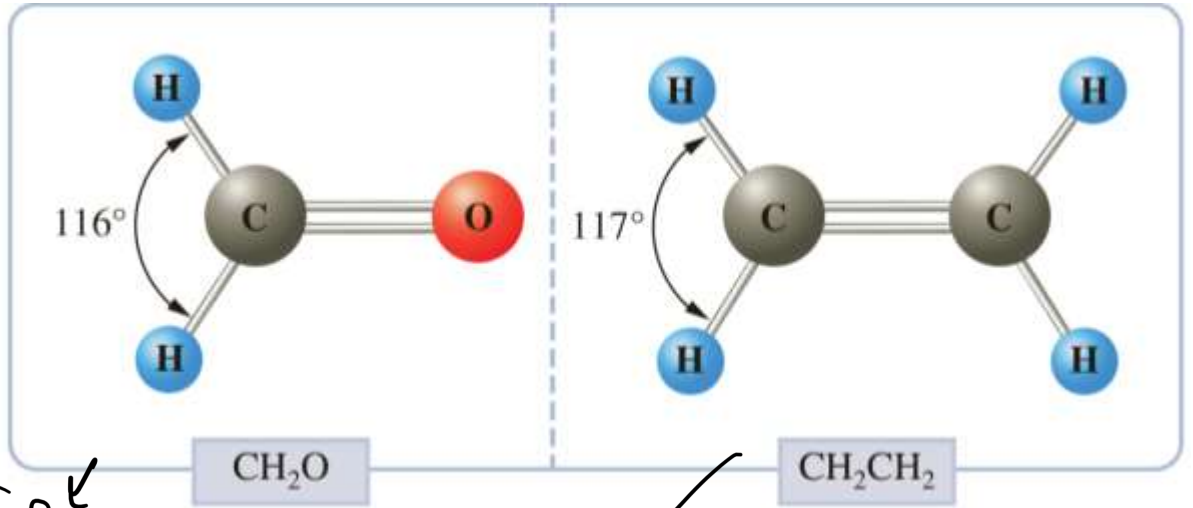
Electron Pairs			Arrangement of Pairs	Molecular Geometry		Example	
Total	Bonding	Lone					
5	5	0	Trigonal bipyramidal	Trigonal bipyramidal AX_5		PCl_5	
	4	1		Seesaw (or distorted tetrahedron) AX_4		SF_4	
	3	2		T-shaped AX_3		ClF_3	
	2	3		Linear AX_2		XeF_2	

Bond Angles and the Effect of Lone Pairs

- ✓ *A lone pair require more space than a bonding pair.*



- ✓ *Multiple bonds require more space than single bonds because of the greater number of electrons.*

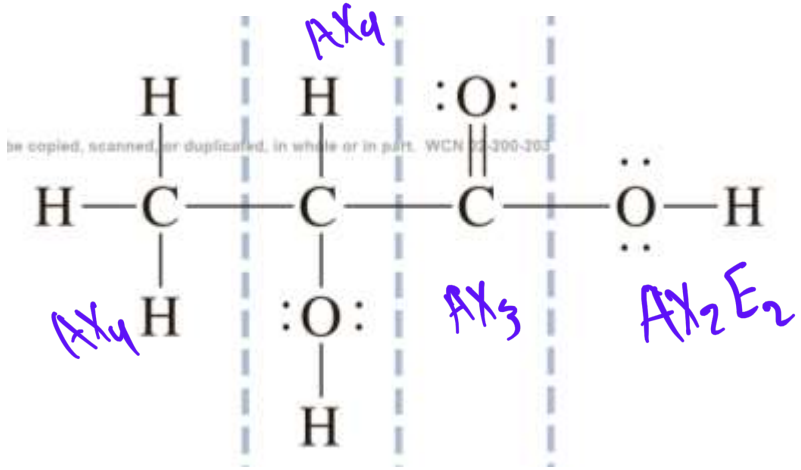


(Q) Predict the geometry of the following molecules or ions, using the VSEPR method:

a. $BeCl_2$ b. NO_2^- c. $SiCl_4$ d. ClO_3^- e. OF_2

f. $TeCl_4$ g. ICl_3

➢ Applying the VSEPR Model to Larger Molecules

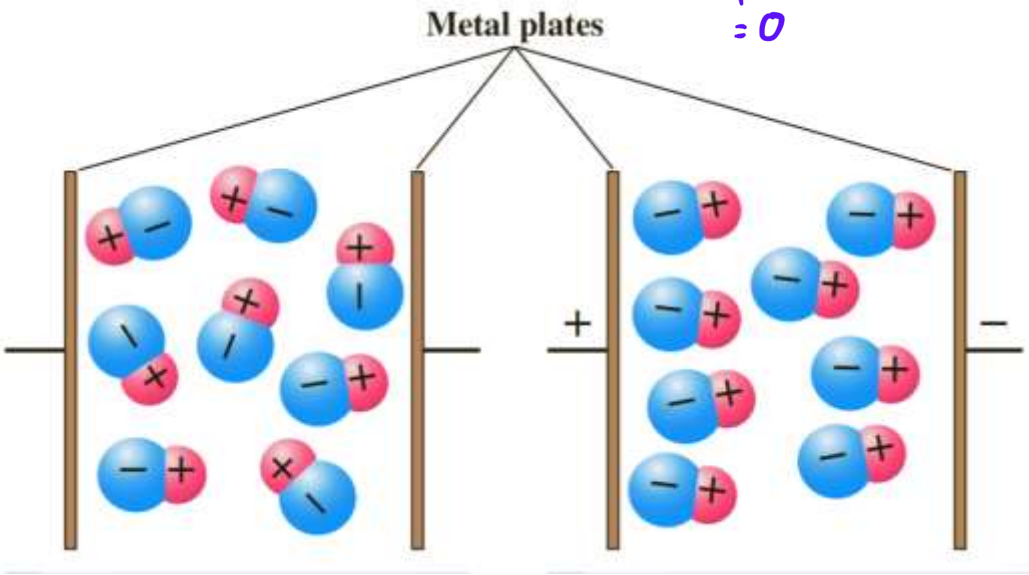


1 IA												13 IIIA	14 IVA	15 VA	16 VIA	17 VIIA	18 VIIIA
1 H Hydrogen 1.008	2 IIA																2 He Helium 4.002602
3 Li Lithium 6.94	4 Be Beryllium 9.0121831											5 B Boron 10.81	6 C Carbon 12.011	7 N Nitrogen 14.007	8 O Oxygen 15.999	9 F Fluorine 18.998403163	10 Ne Neon 20.1797
11 Na Sodium 22.98976928	12 Mg Magnesium 24.305	3 IIIB	4 IVB	5 VB	6 VIB	7 VIIB	8 VIIIB	9 VIIIB	10 VIIIB	11 IB	12 IIB	13 Al Aluminium 26.9815385	14 Si Silicon 28.085	15 P Phosphorus 30.973761998	16 S Sulfur 32.06	17 Cl Chlorine 35.45	18 Ar Argon 39.948
19 K Potassium 39.0983	20 Ca Calcium 40.078	21 Sc Scandium 44.955908	22 Ti Titanium 47.867	23 V Vanadium 50.9415	24 Cr Chromium 51.9961	25 Mn Manganese 54.938044	26 Fe Iron 55.845	27 Co Cobalt 58.933194	28 Ni Nickel 58.6934	29 Cu Copper 63.546	30 Zn Zinc 65.38	31 Ga Gallium 69.723	32 Ge Germanium 72.630	33 As Arsenic 74.921595	34 Se Selenium 78.971	35 Br Bromine 79.904	36 Kr Krypton 83.798
37 Rb Rubidium 85.4678	38 Sr Strontium 87.62	39 Y Yttrium 88.90584	40 Zr Zirconium 91.224	41 Nb Niobium 92.90637	42 Mo Molybdenum 95.95	43 Tc Technetium (98)	44 Ru Ruthenium 101.07	45 Rh Rhodium 102.90550	46 Pd Palladium 106.42	47 Ag Silver 107.8682	48 Cd Cadmium 112.414	49 In Indium 114.818	50 Sn Tin 118.710	51 Sb Antimony 121.760	52 Te Tellurium 127.60	53 I Iodine 126.90447	54 Xe Xenon 131.293

10.2 Dipole Moment and Molecular Geometry

ممكن الرابطة تكون Polar والمركب Dipole moment = 0 معناه Polar

- ✓ Alignment of polar molecules by an electric field



- ✓ **dipole moment** is *a quantitative measure of the degree of charge separation in* ***a molecule*** *and is therefore an indicator of the polarity of the molecule*

$$\mu = q \times d$$

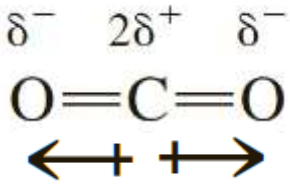
q = positive charge

$-q$ = negative charge

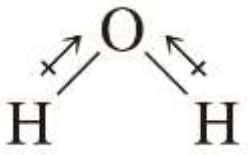
d = distance

10.43 AsF_3 has a dipole moment of 2.59 D. Which of the following geometries are possible: trigonal planar, trigonal pyramidal, or T-shaped?

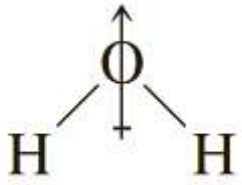
- ✓ dipole moment of HCl is 1.08 D.
- ✓ SI units: coulomb x meter (C·m)
- ✓ 1 D = 3.34 x 10^{-30} C·m



bond dipole



→



- ✓ (linear, trigonal planar, and tetrahedral) give molecules of zero dipole moment; that is, the molecules are ***nonpolar***

Table 10.1 Relationship Between Molecular Geometry and Dipole Moment

Formula	Molecular Geometry	Dipole Moment*
AX	Linear	Can be nonzero
AX_2	Linear	Zero
	Bent	Can be nonzero
AX_3	Trigonal planar	Zero
	Trigonal pyramidal	Can be nonzero
	T-shaped	Can be nonzero
AX_4	Tetrahedral	Zero
	Square planar	Zero
	Seesaw	Can be nonzero
AX_5	Trigonal bipyramidal	Zero
	Square pyramidal	Can be nonzero
AX_6	Octahedral	Zero

Exercise 10.4 Which of the following would be expected to have a dipole moment of zero? Explain

a. $SOCl_2$ b. SiF_4 c. OF_2

(Q) Explain why the dipole moment of NF_3 = 0.2 D, while that of NH_3 =1.47 D

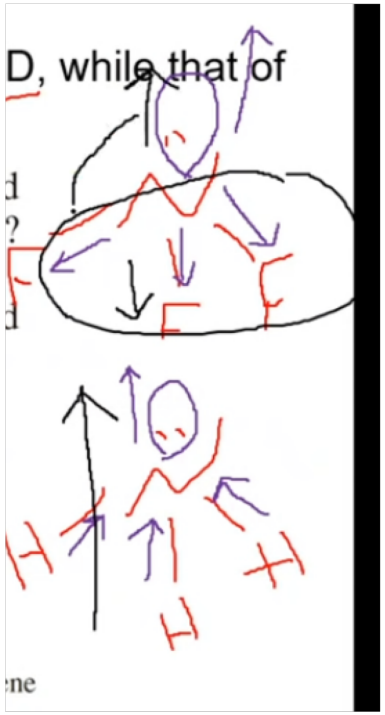
N–H electronegativity

10.45 Which of the following molecules would be expected to have zero dipole moment on the basis of their geometry?

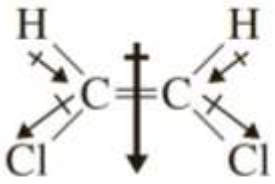
a CS_2 b TeF_2 c $SeCl_4$ d XeF_4

10.46 Which of the following molecules would be expected to have a dipole moment of zero because of symmetry?

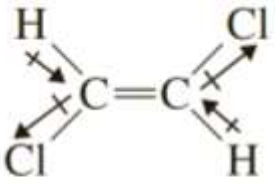
a $BeBr_2$ b H_2Se c AsF_3 d SeF_6



- **Effect of Polarity on Molecular Properties**



cis-1,2-Dichloroethene



trans-1,2-Dichloroethene

Dipole moment:	1.9 D	0 D
B.P (°C) boiling point	60.2	48.5

10.3 Valence Bond Theory

➢ Basic Theory

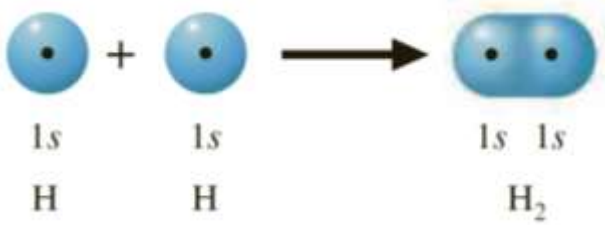
A bond forms between two atoms when the following conditions are met:

تداخل

1. The orbitals containing the electrons *overlap*.
2. The sum of the electrons in both orbitals is no more than two.

$H + H \rightarrow H_2$

$1s^1 \quad 1s^1 \rightarrow$ Total 2e in the newly formed H_2 orbital



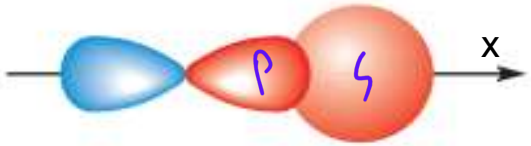
$He + He \rightarrow He_2$ (does not occur)

$1s^2 \quad 1s^2 \rightarrow$ Total 4e (XXXXX)

X

✓ The strength of bonding depends on orbital overlap.

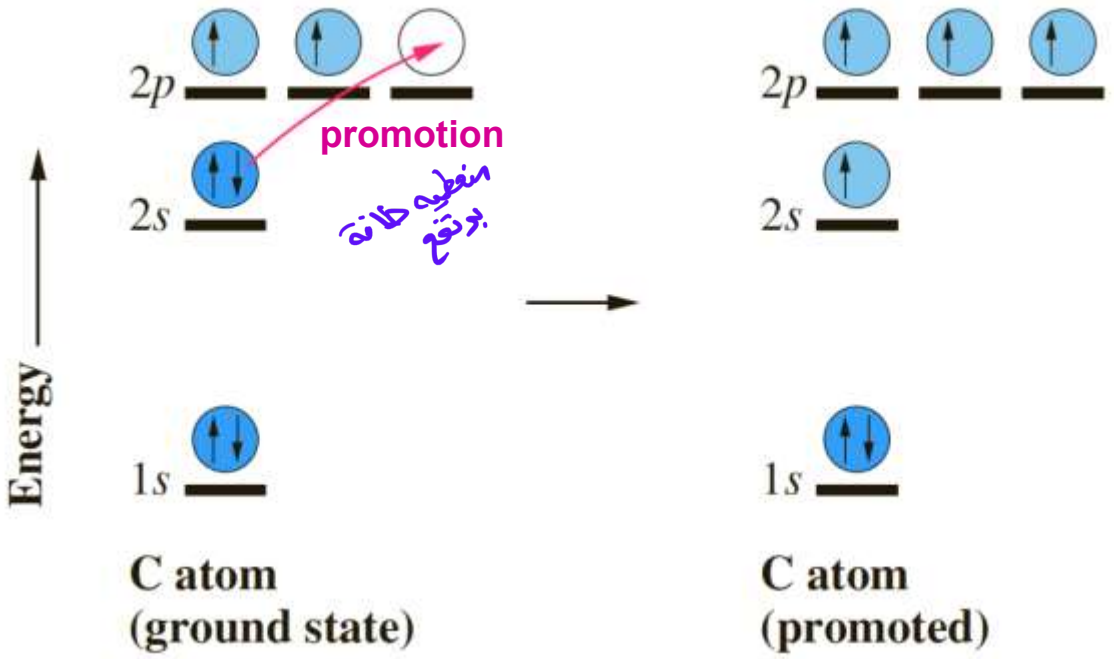
✓ *To obtain maximum overlap,* orbitals other than *s* bond only in given directions.



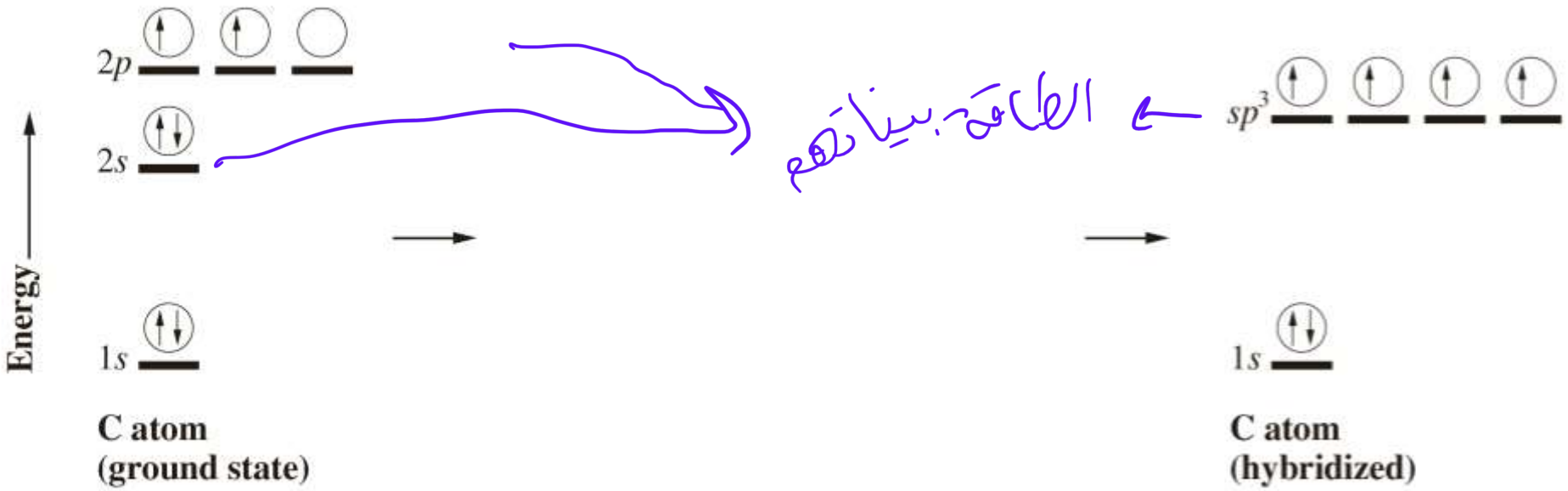
➢ Bonding in HCl

H: $1s^1$ Cl: $1s^2 2s^2 2p^6 3s^2 3p^5$

- **Hybrid Orbitals**
- ✓ **Bonding in CH_4**



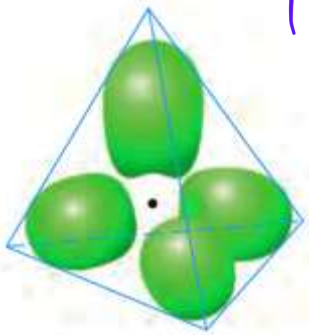
- ✓ Experiment shows that the four C-H bonds in CH_4 are identical. This implies that the carbon orbitals involved in bonding are also equivalent.
- ✓ → **Hybrid orbitals are used**



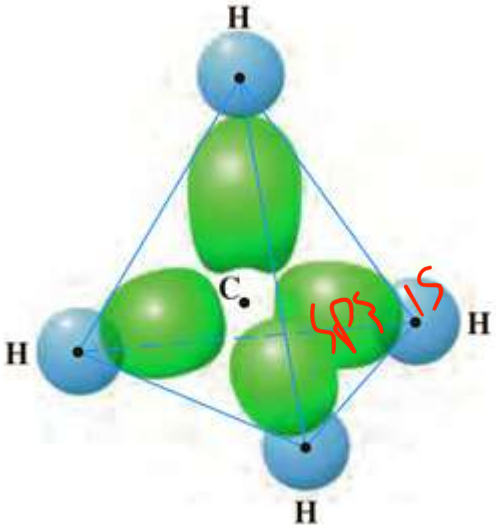
- ✓ *The number of hybrid orbitals formed always equals the number of atomic orbitals used.*

- ✓ Shape of a single sp^3 hybrid orbital.
- ✓ Each orbital consists of two lobes. One lobe is small, but dense, and concentrated near the nucleus.
- ✓ The other lobe is large, but diffuse.
- ✓ Bonding occurs with the large lobe, since it extends farther from the nucleus.

offers high overlap



- ✓ Four hybrid orbitals are arranged tetrahedrally in space. (Small lobes are omitted here for clarity, and large lobes are stylized and greatly narrowed for ease in depicting the directional bonding.)



- ✓ Bonding in CH_4.
- ✓ Each C—H bond is formed by the overlap of:
- ✓ 1s orbital from a hydrogen atom **+** one sp^3 hybrid orbital of the carbon atom.

- ✓ **Number of hybrid orbitals formed = number of atomic orbitals used.**

same energy

$s + p \rightarrow 2 \times sp$

$s + 2p \rightarrow 3 \times sp^2$

$s + 3p \rightarrow 4 \times sp^3$

$s + 3p \times d \rightarrow 5\ sp^3d$

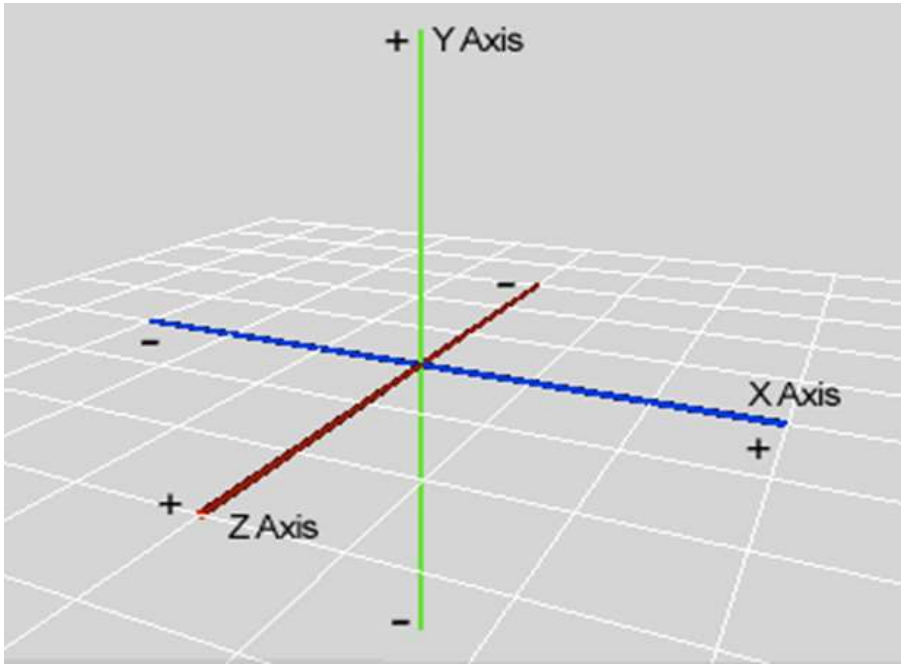
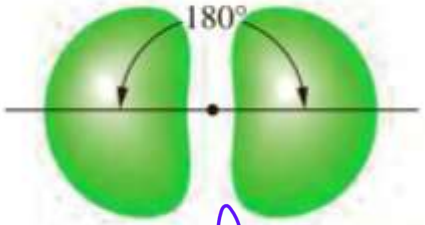
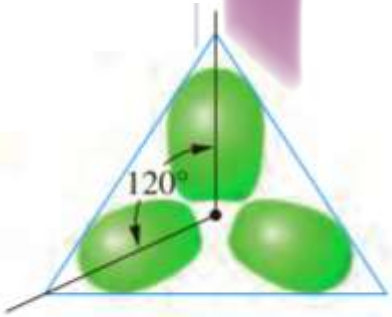


Table 10.2 Kinds of Hybrid Orbitals

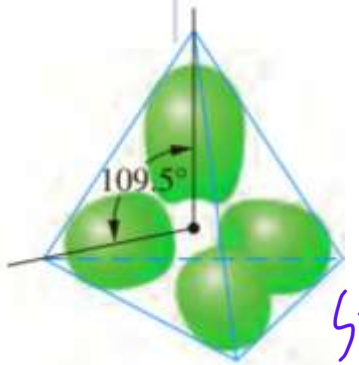
Hybrid Orbitals	Geometric Arrangement	Number of Orbitals	Example
sp	Linear	2	Be in BeF_2
sp^2	Trigonal planar	3	B in BF_3
sp^3	Tetrahedral	4	C in CH_4



sp

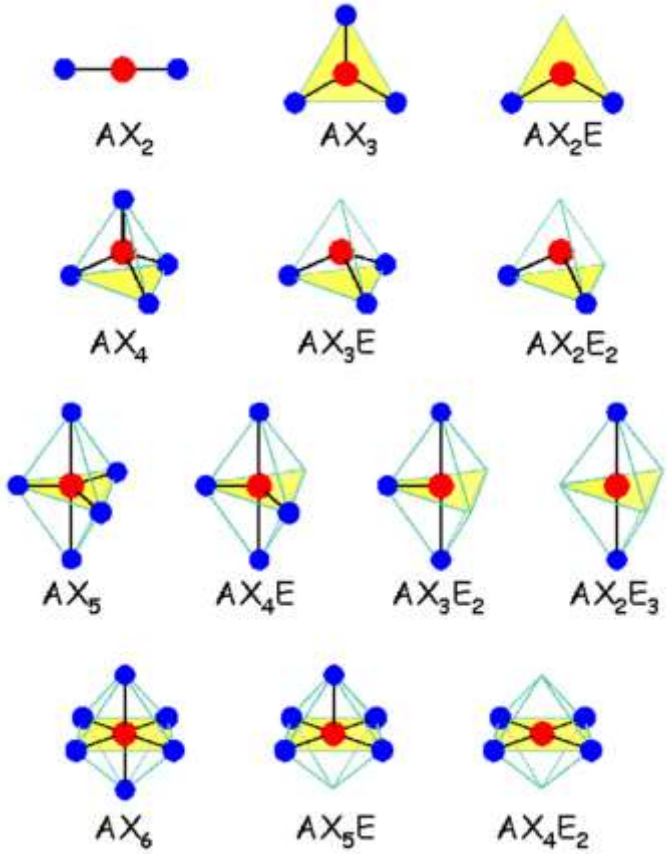


sp^2



sp^3

How to figure out the hybridization via Lewis structures !



what is the hybridization for central atom in BeH_2

$2+2=4$

H — Be — H

AX_2 ⇒ linear ⇒ SP

2 objects SP

H_2O SP^3

HCN SP

- **More Than Eight Electrons About a Central Atom**

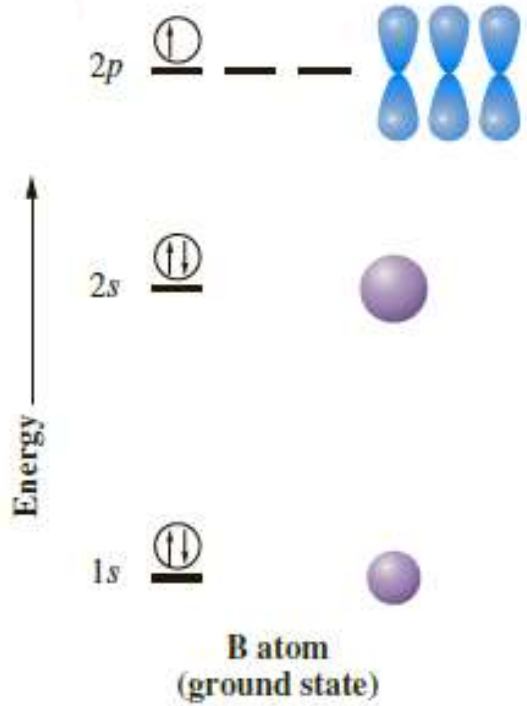
PF_5 → sp^3d

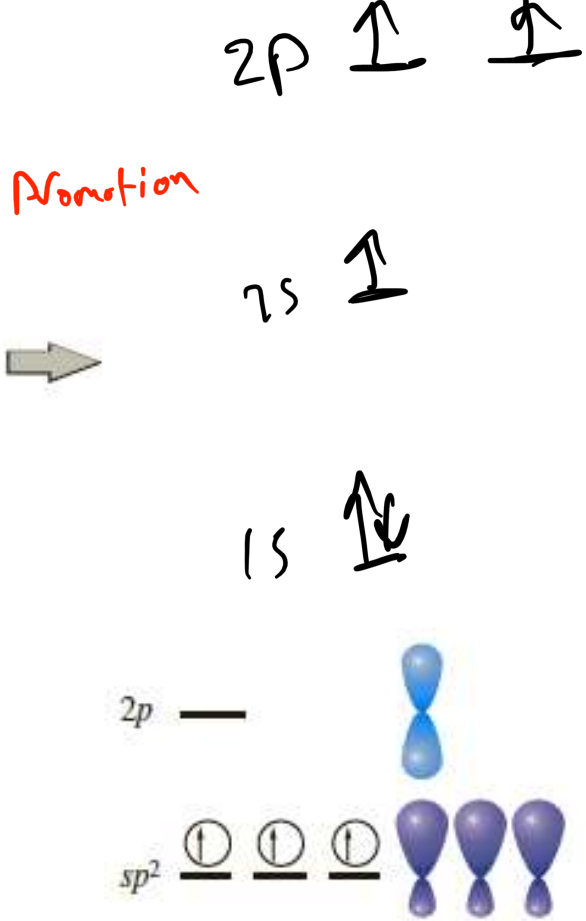
SF_6 → sp^3d^2

SF_6

sp^3d^2

- Hybridization in BF_3





Promotion

hybridized

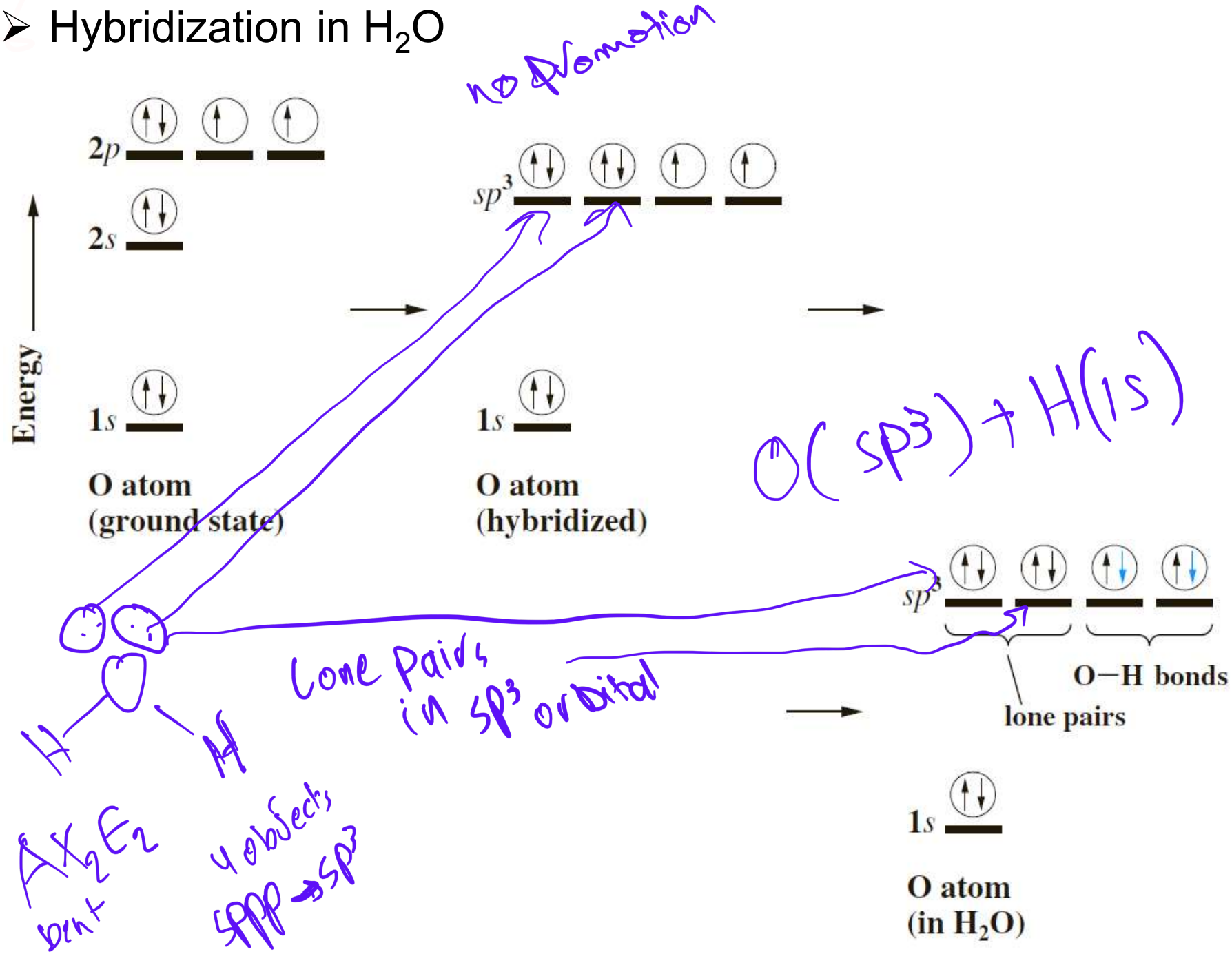
overlap

for 3F

$F: 1s^2 2s^2 2p^5$

$B(sp^2) - F(2p)$

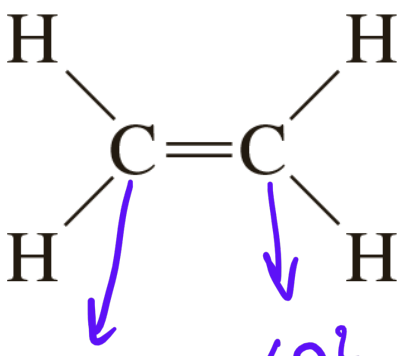
- Hybridization in H_2O

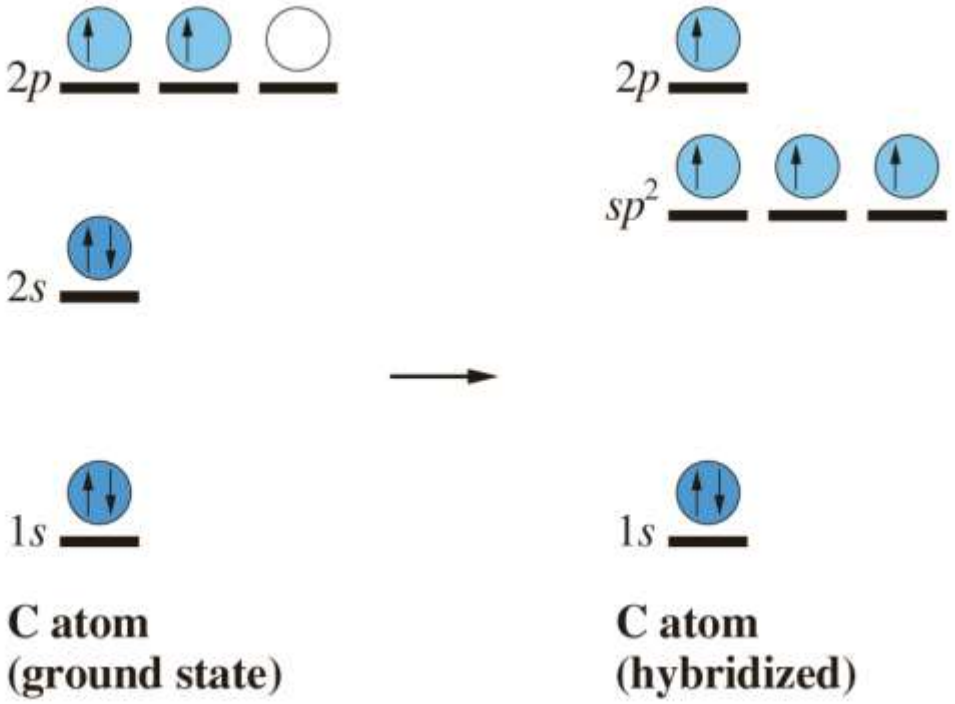


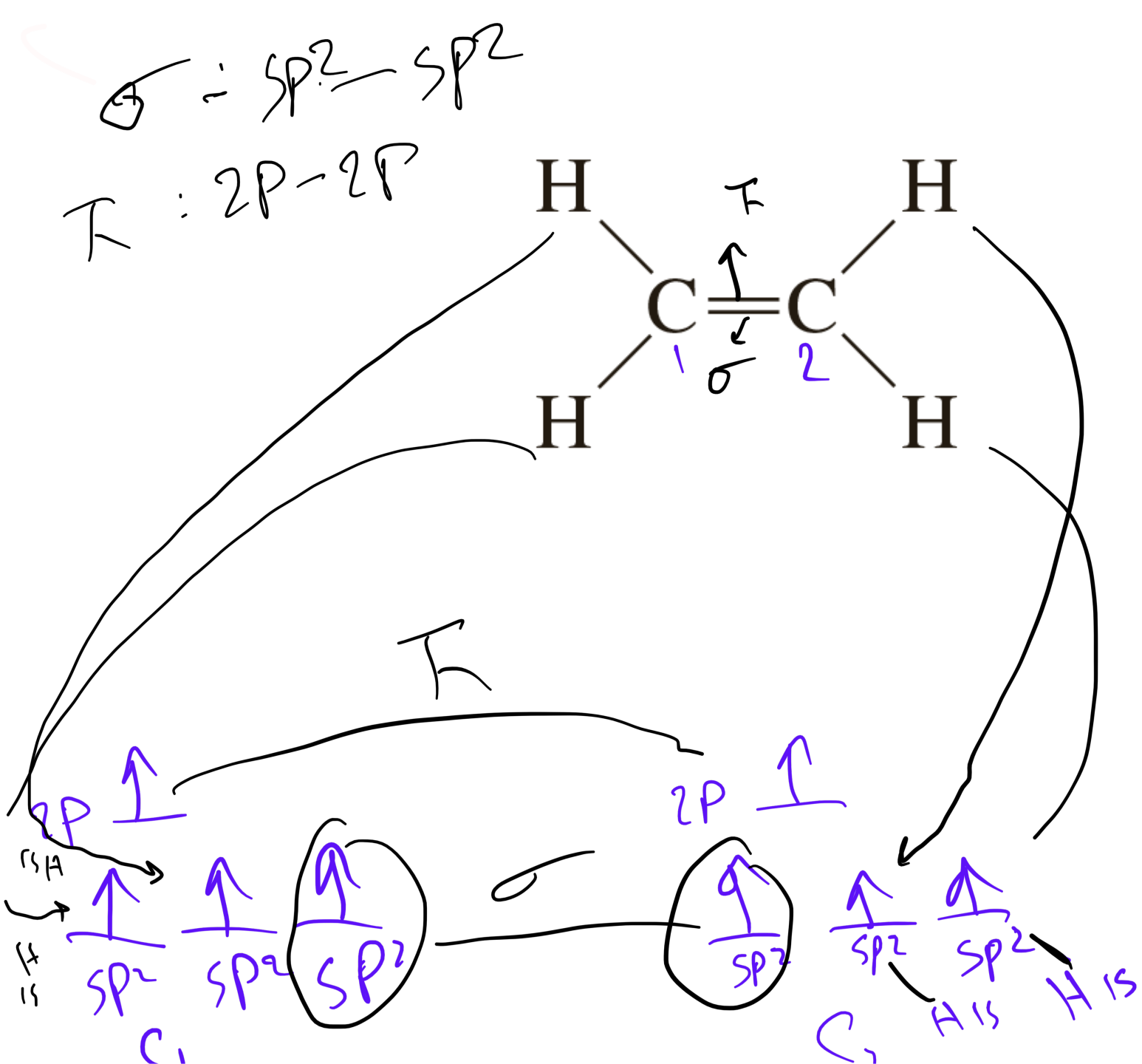
10.4 Description of Multiple Bonding

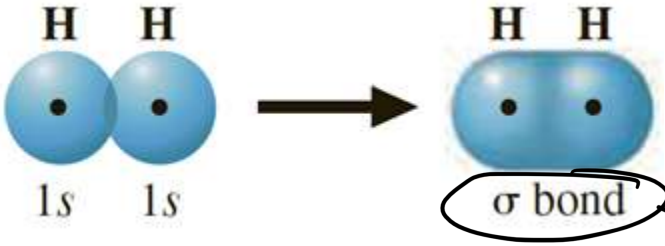
- ✓ More than one orbital from each bonding atom might overlap
- ✓ One hybrid orbital is needed for each bond (whether a single or a multiple bond) and for each lone pair

➢ Bonding in ethylene $H_2C=CH_2$



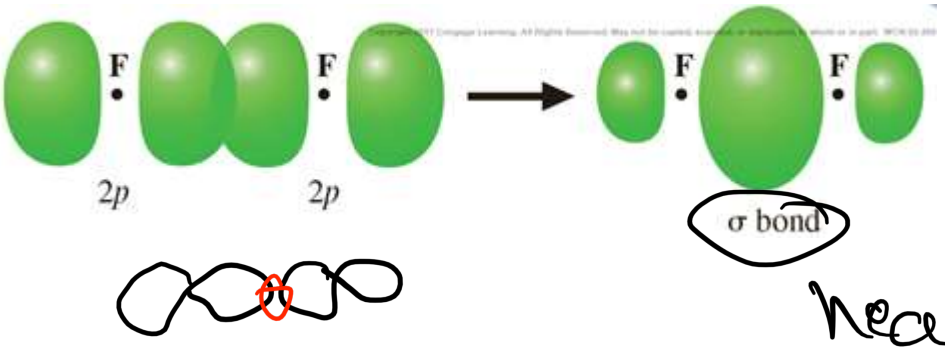



σ : sp² — sp²
π : 2p — 2p
H
H
C
C
π
σ
1
2
H
H
π
2p
2p
sp²
sp²
sp²
sp²
sp²
sp²
σ
C1
C2
H 1s
H 1s



The formation of a σ bond by the overlap of two *s* orbitals.

head to head

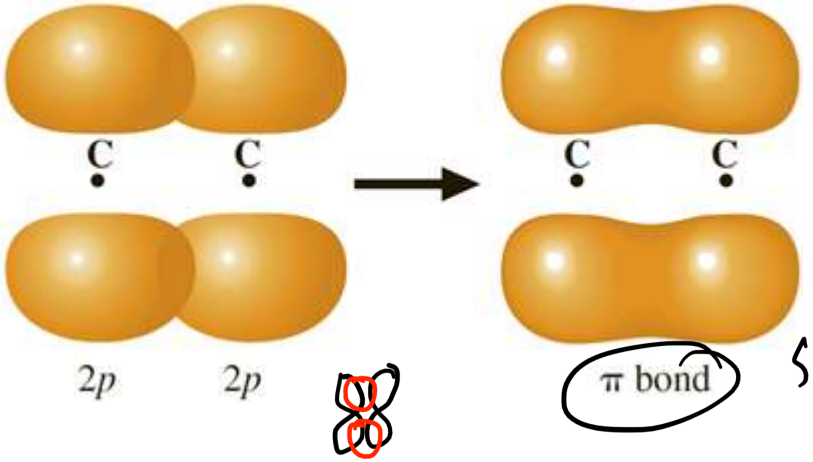


A σ bond can also be formed by the overlap of two *p* orbitals along their axes.

head to head

p or s

head to head

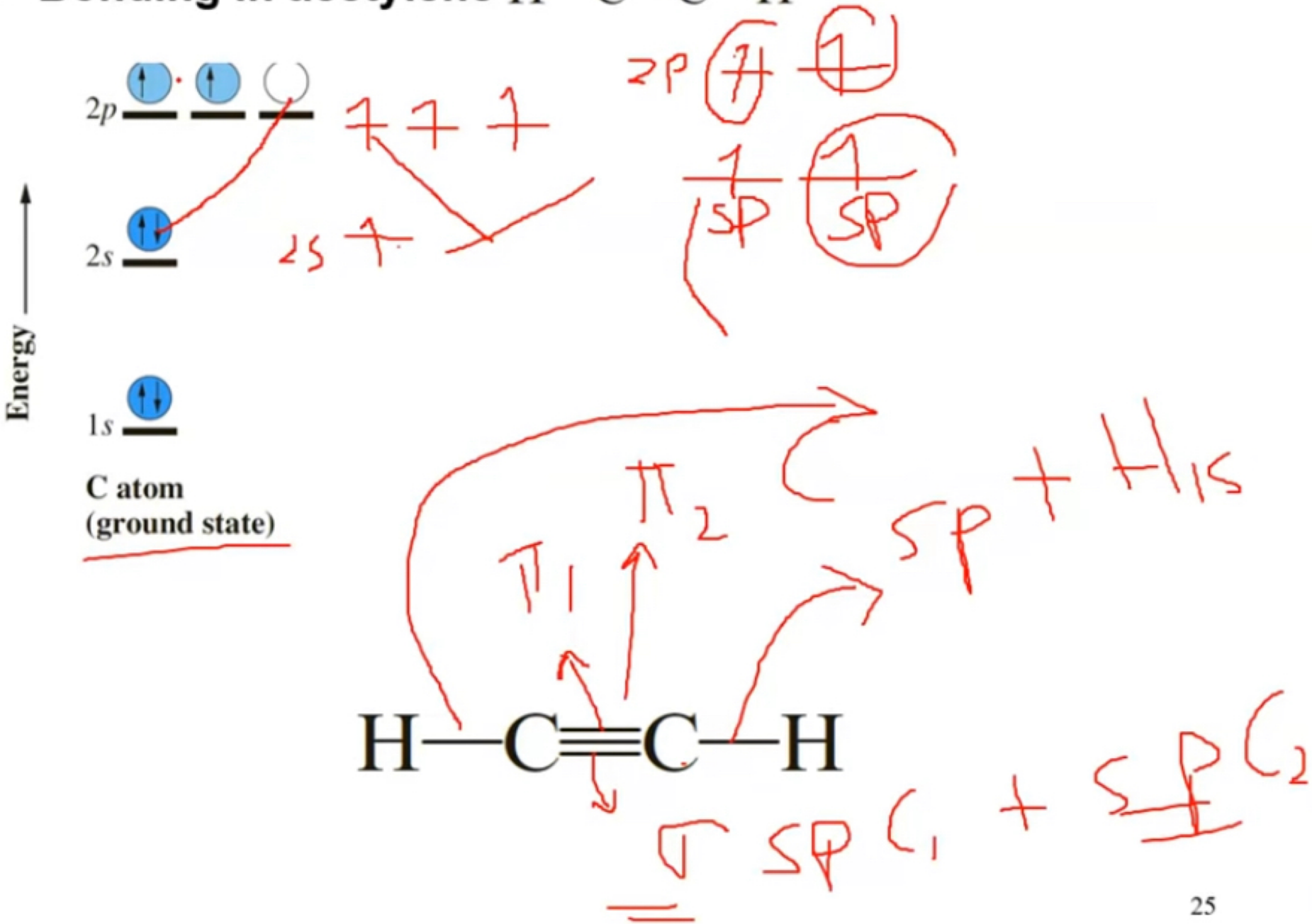


When two *p* orbitals overlap sideways, a π bond is formed.

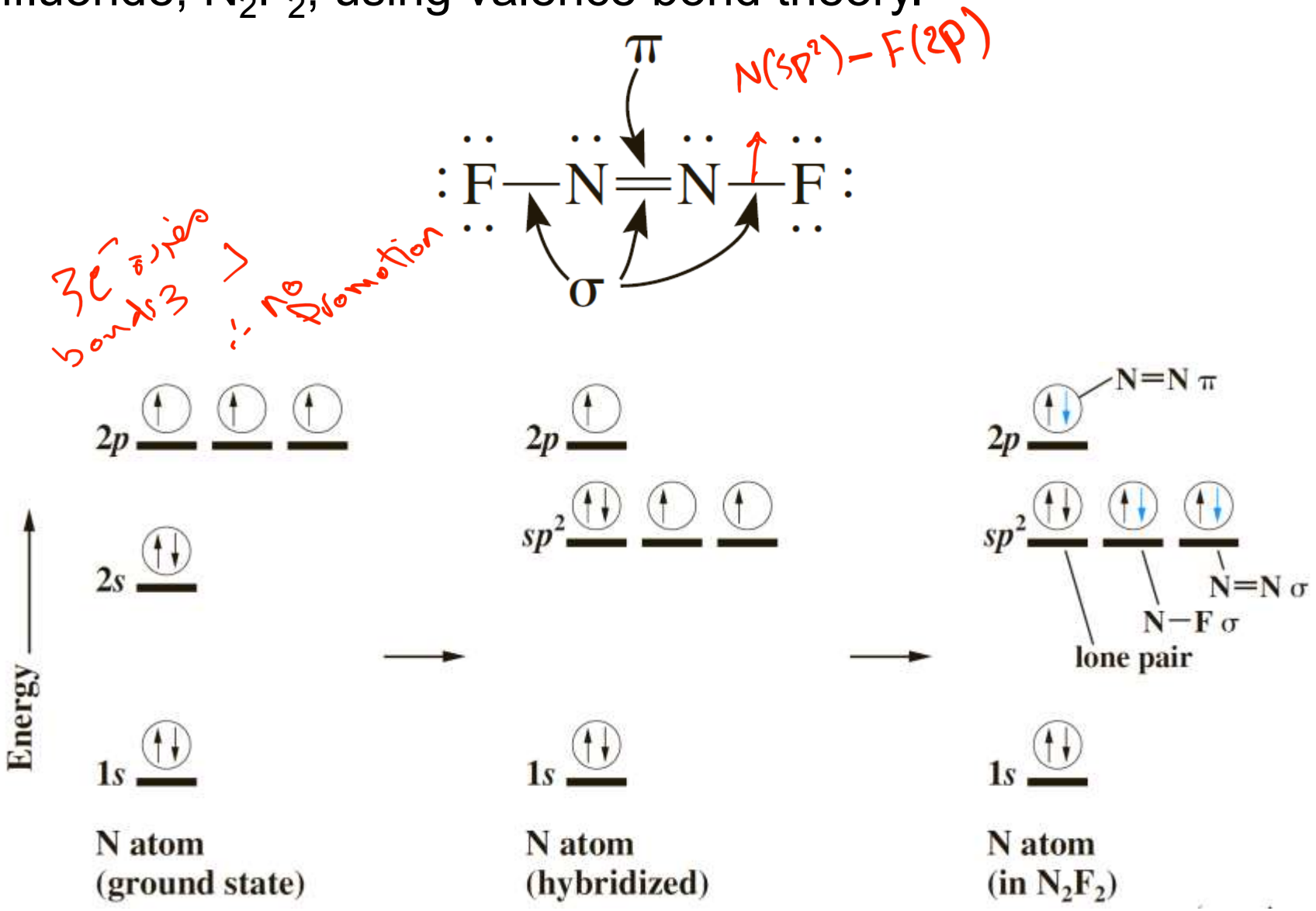
sideway

in π overlap is less than σ so its weaker

➢ Bonding in acetylene H—C≡C—H



(Q) Describe the bonding on a given N atom in dinitrogen difluoride, N_2F_2, using valence bond theory.

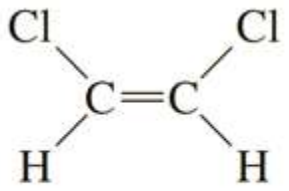


10.53 **a** Carbonyl fluoride, COF_2, is an extremely poisonous gas used in organofluorine synthesis. Give the valence bond description of the carbonyl fluoride molecule. (Both fluorine atoms are attached to the carbon atom.) **b** Nitrogen, N_2, makes up about 80% of the earth's atmosphere. Give the valence bond description of this molecule.

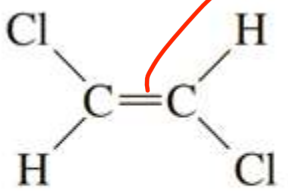
10.54 HN=NH

10.55 HCN

✓ *Isomers* are compounds of the same molecular formula but with different arrangements of the atoms.



cis-1,2-Dichloroethene

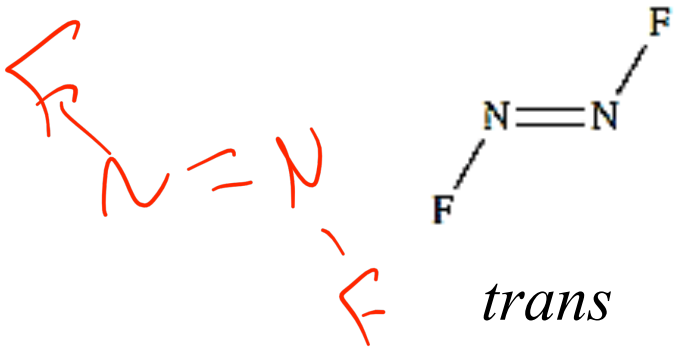


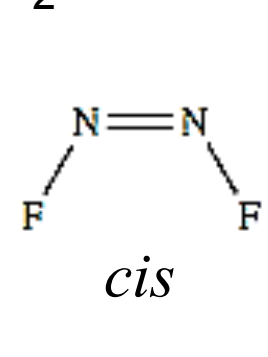
trans-1,2-Dichloroethene

Dipole moment:	1.9 D	0 D
B.P (°C)	60.2	48.5

no free rotation (need High energy)

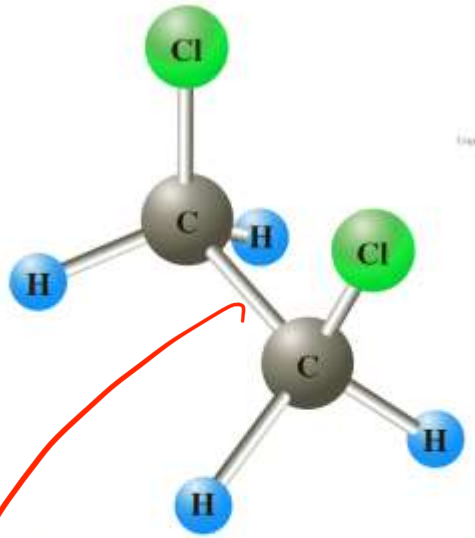
✓ *cis* and *trans* isomers of N_2F_2



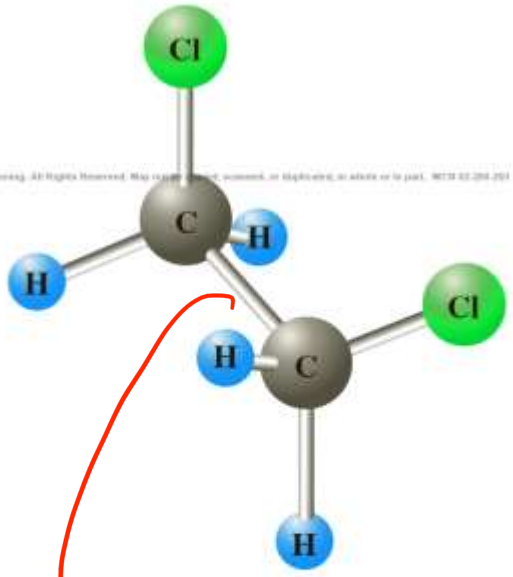


or

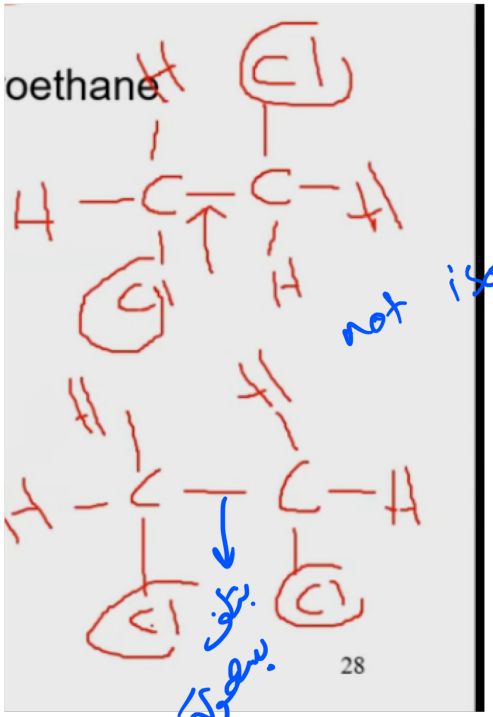
✓ Lack of geometric isomers in 1,2-dichloroethane



a Because of rotation about the carbon–carbon bond in 1,2-dichloroethane, geometric isomers are not possible.



b Note that the molecule pictured at the left can be twisted easily to give the molecule pictured here.



Free rotation
(low energy)

cis
Isomers