## Chapter – 4

# Chemical Bonding and Molecular Structure

## 2marks:

1. Write the favourable factors for the formation of ionic bond.

#### **Answer:**

An ionic bond is formed by the transfer of one or more electrons from one atom to another. Hence, the formation of ionic bonds depends upon the ease with which neutral atoms can lose or gain electrons.

Bond formation also depends upon the lattice energy of the compound formed.

Hence, favourable factors for ionic bond formation are as follows:

- (i) Low ionization enthalpy of metal atom.
- (ii) High electron gain enthalpy ( $\Delta$ eg H) of a non-metal atom.
- (iii) High lattice energy of the compound formed.

### 2. How do you express the bond strength in terms of bond order?

#### **Answer:**

Bond strength represents the extent of bonding between two atoms

forming a molecule. The larger the bond energy, the stronger is the bond and the greater is the bond order.

### 3. Define the bond length.

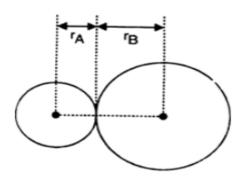
#### **Answer:**

Bond length is defined as the equilibrium distance between the nuclei of two bonded atoms in a molecule.

Bond lengths are expressed in terms of Angstrom (10–10 m) or picometer

(10–12 m) and are measured by spectroscopic X-ray diffractions and electron-diffraction techniques.

In an ionic compound, the bond length is the sum of the ionic radii of the constituting atoms ( $d = r_+ + r_-$ ). In a covalent compound, it is the sum of their covalent radii ( $d = r_A + r_B$ ).



4. Explain the important aspects of resonance with reference to the  $CO_3^{2-}$  ion.

#### **Answer:**

According to experimental findings, all carbon to oxygen bonds in  $^{\text{CO}_3^{2^-}}$  are equivalent. Hence, it is inadequate to represent ion by a single Lewis structure having two single bonds and one double bond.

Therefore, carbonate ion is described as a resonance hybrid of the following structures:

5. H<sub>3</sub>PO<sub>3</sub> can be represented by structures 1 and 2 shown below. Can these two structures be taken as the canonical forms of the resonance hybrid representing H<sub>3</sub>PO<sub>3</sub>? If not, give reasons for the same.

#### **Answer:**

The given structures cannot be taken as the canonical forms of the resonance hybrid of H<sub>3</sub>PO<sub>3</sub> because the positions of the atoms have changed.

6. Arrange the bonds in order of increasing ionic character in the molecules: LiF,  $K_2O$ ,  $N_2$ ,  $SO_2$  and  $ClF_3$ .

#### **Answer:**

The ionic character in a molecule is dependent upon the electronegativity difference between the constituting atoms. The greater the difference, the greater will be the ionic character of the molecule.

On this basis, the order of increasing ionic character in the given molecules is

$$N_2 < SO_2 < ClF_3 < K_2O < LiF.$$

7. The skeletal structure of CH<sub>3</sub>COOH as shown below is correct, but some of the bonds are shown incorrectly. Write the correct Lewis structure for acetic acid.

#### **Answer:**

The correct Lewis structure for acetic acid is as follows:

## 8. Explain why BeH<sub>2</sub> molecule has a zero dipole moment although the Be–H bonds are polar.

#### **Answer:**

The Lewis structure for BeH<sub>2</sub> is as follows:

There is no lone pair at the central atom (Be) and there are two bond pairs. Hence, BeH<sub>2</sub> is of the type AB<sub>2</sub>. It has a linear structure.

$$H \xrightarrow{+-} Be \xrightarrow{--} H$$

Dipole moments of each H–Be bond are equal and are in opposite directions. Therefore, they nullify each other. Hence, BeH2 molecule has zero dipole moment.

## 9. Is there any change in the hybridisation of B and N atoms as a result of the following reaction?

$$BF_3 + NH_3 \rightarrow F_3B.NH_3$$

#### **Answer:**

Boron atom in BF<sub>3</sub> is  $s^{p2}$  hybridized. The orbital picture of boron in the excited state can be shown as:

$$\begin{array}{c|c}
\uparrow & \uparrow \uparrow \\
2s & 2p_x 2p_y 2p_z
\end{array}$$

Nitrogen atom in NH<sub>3</sub> is sp<sup>3</sup> hybridized. The orbital picture of nitrogen can be represented as:

$$\begin{array}{ccc}
\uparrow \downarrow & & \uparrow & \uparrow & \uparrow \\
2s & & 2p_x & 2p_y & 2p_z
\end{array}$$

After the reaction has occurred, an adduct F<sub>3</sub>B·NH<sub>3</sub> is formed as hybridization of 'B' changes to sp3. However, the hybridization of 'N' remains intact.

## 10. What is the total number of sigma and pi bonds in the following molecules?

#### **Answer:**

A single bond is a result of the axial overlap of bonding orbitals.

Hence, it contributes a sigma bond. A multiple bond (double or triple bond) is always formed as a result of the sidewise overlap of orbitals. A pi-bond is always present in it. A triple bond is a combination of two pi-bonds and one sigma bond.

Structure of C<sub>2</sub>H<sub>2</sub> can be represented as:

$$H \xrightarrow{\sigma} C \xrightarrow{\pi} C \xrightarrow{\sigma} H$$

Hence, there are three sigma and two pi-bonds in  $C_2H_2$ .

The structure of  $C_2H_4$  can be represented as:

Hence, there are five sigma bonds and one pi-bond in C<sub>2</sub>H<sub>4</sub>.

11. Considering x-axis as the internuclear axis which out of the following will not form a sigma bond and why? (a) 1s and 1s (b)

1s and 2px (c) 2py and 2py (d) 1s and 2s.

#### **Answer:**

2py and 2py orbitals will not a form a sigma bond. Taking x-axis as the internuclear axis, 2py and 2py orbitals will undergo lateral overlapping, thereby forming a pi  $(\pi)$  bond.

12. What do you understand by bond pairs and lone pairs of electrons? Illustrate by giving one example of each type.

#### **Answer:**

When two atoms combine by sharing their one or more valence electrons, a covalent bond is formed between them.

The shared pairs of electrons present between the bonded atoms are called bond pairs. All valence electrons may not participate in bonding. The electron pairs that do not participate in bonding are called lone pairs of electrons.

For example, in  $C_2H_6$  (ethane), there are seven bond pairs but no lone pair present.

$$egin{array}{cccc} \mathbf{H} & \mathbf{H} \\ & | & | \\ \mathbf{H} - \mathbf{C} - \mathbf{C} - \mathbf{H} \\ & | & | \\ \mathbf{H} & \mathbf{H} \end{array}$$

In H<sub>2</sub>O, there are two bond pairs and two lone pairs on the central atom (oxygen).



13. Write the important conditions required for the linear combination of atomic orbitals to form molecular orbitals.

#### **Answer:**

The given conditions should be satisfied by atomic orbitals to form molecular orbitals:

- (a) The combining atomic orbitals must have the same or nearly the same energy. This means that in a homonuclear molecule, the 1s-atomic orbital of an atom can combine with the 1s-atomic orbital of another atom, and not with the 2s-orbital.
- (b) The combining atomic orbitals must have proper orientations to ensure that the overlap is maximum.
- (c) The extent of overlapping should be large.

## 14. Use molecular orbital theory to explain why the Be<sub>2</sub> molecule does not exist.

**Answer:** The electronic configuration of Beryllium is 1s<sup>2</sup> 2s<sup>2</sup>

The molecular orbital electronic configuration for Be<sub>2</sub> molecule can be written as:

$$\sigma_{1s}^2$$
  $\sigma_{1s}^{\bullet 2}$   $\sigma_{2s}^2$   $\sigma_{2s}^{\bullet 2}$ 

Hence, the bond order for Be2 is 1/2(Nb - Na)

Where,

Nb = Number of electrons in bonding orbitals

Na = Number of electrons in anti-bonding orbitals

∴ Bond order of Be2 1/2(4 - 4) = 0

A negative or zero bond order means that the molecule is unstable. Hence, Be2 molecule does not exist.

## 15. Write the significance of a plus and a minus sign shown in representing the orbitals.

#### **Answer:**

Molecular orbitals are represented by wave functions. A plus sign in an orbital indicates a positive wave function while a minus sign in an orbital represents a negative wave function.

## 4marks:

## 1. Explain the formation of a chemical bond.

#### **Answer:**

A chemical bond is defined as an attractive force that holds the constituents (atoms, ions etc.) together in a chemical species.

Various theories have been suggested for the formation of chemical bonds such as the electronic theory, valence shell electron pair repulsion theory, valence bond theory, and molecular orbital theory.

A chemical bond formation is attributed to the tendency of a system to attain stability. It was observed that the inertness of noble gases was because of their fully filled outermost orbitals. Hence, it was postulated that the elements having incomplete outermost shells are unstable (reactive). Atoms, therefore, combine with each other and complete their respective octets or duplets to attain the stable configuration of the nearest noble gases. This combination can occur either by sharing of electrons or by transferring one or more electrons from one atom to another. The chemical bond formed as a result of sharing of electrons between atoms is called a covalent bond. An ionic bond is formed as a result of the transference of electrons from one atom to another.

2. Write Lewis dot symbols for atoms of the following elements: Mg, Na, B, O, N, Br.

### **Answer:**

Mg: There are two valence electrons in Mg atom. Hence, the Lewis dot symbol for Mg is:  $^{12}$ Mg = 2, 8, 2  $\therefore$  Lewis symbol =  $\dot{M}$ g.

Na: There is only one valence electron in an atom of sodium. Hence, the Lewis dot structure is:  $_{11}$ Na = 2, 8, 1  $\therefore$  Lewis symbol =  $\dot{N}$ a

**B:** There are 3 valence electrons in Boron atom. Hence, the Lewis dot structure is:  ${}_{5}B = 2$ , 3  $\therefore$  Lewis symbol =  $\cdot \dot{B} \cdot$ 

O: There are six valence electrons in an atom of oxygen. Hence, the Lewis dot structure is:  ${}_{8}O = 2$ , 6  $\therefore$  Lewis symbol =  $:\ddot{O}$ :

N: There are five valence electrons in an atom of nitrogen. Hence, the Lewis dot structure is:  $_{7}N = 2,5$   $\therefore$  Lewis symbol =  $:\dot{N}$ 

**Br:** There are seven valence electrons in bromine. Hence, the Lewis dot structure is:  ${}_{35}Br = 2, 8, 18, 7$  :: Lewis symbol = :  $\ddot{B}r$ .

3. Write Lewis symbols for the following atoms and ions:

S and S<sup>2-</sup>; Al and Al<sup>3+</sup>; H and H<sup>-</sup>

**Answer:** (i) S and S<sup>2-</sup>

The number of valence electrons in sulphur is 6.

The Lewis dot symbol of sulphur (S) is :S:.

The di negative charge infers that there will be two electrons more in addition to the six valence electrons. Hence, the Lewis dot symbol of  $S^{2-}$  is  $[:\ddot{S}:]^{2-}$ .

The number of valence electrons in aluminium is 3.

The Lewis dot symbol of aluminium (Al) is.AI

The tri positive charge on a species infers that it has donated its three electrons. Hence, the Lewis dot symbol is [Al]<sup>+3</sup>.

The number of valence electrons in hydrogen is 1.

The Lewis dot symbol of hydrogen (H) is H.

The uni negative charge infers that there will be one electron more in addition to the one valence electron. Hence, the Lewis dot symbol is [H]<sup>+1</sup>.

### 4. Draw the Lewis structures for the following molecules and ions:

## H<sub>2</sub>S, SiCl<sub>4</sub>, BeF2,CO<sub>3</sub><sup>2-</sup>, HCOOH

#### **Answer:**

 H2S H has 1 electron in its valence shell & sulphur has 6 electron its valence shell. A planar structure will be formed

 Si has 4 electrons its valence shell & CI need only 1 electron to complete its octet, therefore 4 CI will make 4 bonds with Si.A tetrahedral shape will be formed

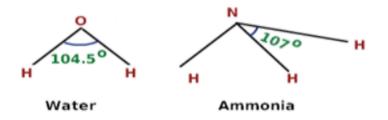
Be has 2 electrons in its valence shell & Fe has 2 electrons in its valence shell, so they make a planar structure

4) here C has 4 electron its valence shell, therefore it can make 4 bond with other atoms, whereas O has 6 electrons in its valence shell, it can accept 2 electron easily to complete its octet, therefore one O will make a double bond with C, while other two O will make 2 single bond with C.

5. Although geometries of NH<sub>3</sub> and H<sub>2</sub>O molecules are distorted tetrahedral, bond angle in water is less than that of ammonia. Discuss.

#### **Answer:**

The molecular geometry of NH<sub>3</sub> and H<sub>2</sub>O can be shown as:



The central atom (N) in NH3 has one lone pair and there are three bond pairs. In H<sub>2</sub>O, there are two lone pairs and two bond pairs.

The two lone pairs present in the oxygen atom of  $H_2O$  molecule repels the two bond pairs. This repulsion is stronger than the repulsion between the lone pair and the three bond pairs on the nitrogen atom.

Since the repulsions on the bond pairs in H<sub>2</sub>O molecule are greater than that in NH<sub>3</sub>, the bond angle in water is less than that of ammonia.

6. Write the resonance structures for SO<sub>3</sub>, NO<sub>2</sub> and NO<sub>3</sub><sup>-</sup>.

#### **Answer:**

The resonance structures are:

(a) SO3:

(b) **NO**2

(c)  $NO_3^-$ 

7. Although both  $CO_2$  and  $H_2O$  are triatomic molecules, the shape of  $H_2O$  molecule is bent while that of  $CO_2$  is linear. Explain this on the basis of dipole moment.

### **Answer:**

According to experimental results, the dipole moment of carbon dioxide is zero. This is possible only if the molecule is linear so that

the dipole moments of C–O bonds are equal and opposite to nullify each other.

## Resultant $\mu = 0$ D

H<sub>2</sub>O, on the other hand, has a dipole moment value of 1.84 D (though it is a triatomic molecule as CO<sub>2</sub>). The value of the dipole moment suggests that the structure of H<sub>2</sub>O molecule is bent where the dipole moment of O–H bonds are unequal.



## 8. Define electronegativity. How does it differ from electron gain enthalpy?

#### **Answer:**

Electronegativity is the ability of an atom in a chemical compound to attract a bond pair of electrons towards itself.

Electronegativity of any given element is not constant. It varies

according to the element to which it is bound. It is not a measurable quantity. It is only a relative number.

On the other hand, electron gain enthalpy is the enthalpy change that takes place when an electron is added to a neutral gaseous atom to form an anion. It can be negative or positive depending upon whether the electron is added or removed. An element has a constant value of the electron gain enthalpy that can be measured experimentally.

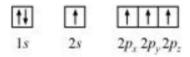
9. Apart from tetrahedral geometry, another possible geometry for CH<sub>4</sub> is square planar with the four H atoms at the corners of the square and the C atom at its centre. Explain why CH<sub>4</sub> is not square planar?

### **Answer:**

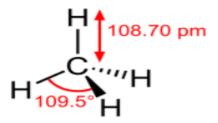
Electronic configuration of carbon atom:

6C: 
$$1s^2 2s^2 2p^2$$

In the excited state, the orbital picture of carbon can be represented as:



Hence, carbon atom undergoes sp<sup>3</sup> hybridization in CH<sub>4</sub> molecule and takes a tetrahedral shape.



For a square planar shape, the hybridization of the central atom has to be dsp<sup>2</sup>. However, an atom of carbon does not have d-orbit alsto undergo dsp<sup>2</sup> hybridization. Hence, the structure of CH<sub>4</sub> cannot be square planar.

Moreover, with a bond angle of 90° in square planar, the stability of CH<sub>4</sub> will be very less because of the repulsion existing between the bond pairs. Hence, VSEPR theory also supports a tetrahedral structure for CH<sub>4</sub>.

## 10. Which out of NH<sub>3</sub> and NF<sub>3</sub> has higher dipole moment and why?

#### **Answer:**

In both molecules i.e., NH<sub>3</sub> and NF<sub>3</sub>, the central atom (N) has a lone pair electron and there are three bond pairs. Hence, both molecules have a pyramidal shape. Since fluorine is more electronegative than hydrogen, it is expected that the net dipole moment of NF<sub>3</sub> is greater

than  $NH_3$ . However, the net dipole moment of  $NH_3$  (1.46 D) is greater than that of  $NF_3$  (0.24 D).

This can be explained on the basis of the directions of the dipole moments of each individual bond in NF<sub>3</sub> and NH<sub>3</sub>. These directions can be shown as:

$$F = \begin{cases} O \downarrow \\ N \downarrow \\ F \end{cases}$$
Resultant of  $H = \begin{cases} O \downarrow \\ N \downarrow \\ H \end{cases}$ 
Resultant of  $H = \begin{cases} O \downarrow \\ N \downarrow \\ N \downarrow \\ H \end{cases}$ 
Resultant of  $H = \begin{cases} O \downarrow \\ N \downarrow \\ N$ 

Thus, the resultant moment of the N-H bonds add up to the bond moment of the lone pair (the two being in the same direction), whereas that of the three N-F bonds partly cancels the moment of the lone pair.

Hence, the net dipole moment of NF3 is less than that of NH3.

## 11. Describe the change in hybridisation (if any) of the Al atom in the following reaction.

$$AlCl_3 + Cl^- \rightarrow AlCl_4$$

#### **Answer:**

The valence orbital picture of aluminium in the ground state can be

represented as:

The orbital picture of aluminium in the excited state can be represented as:

$$\begin{array}{c|c}
\uparrow & \uparrow & \uparrow \\
3s & 3p_x 3p_y 3p_z
\end{array}$$

Hence, it undergoes sp<sup>2</sup> hybridization to give a trigonal planar arrangement (in AlCl3).

To form AlCl<sub>4</sub><sup>-</sup>, the empty 3pz orbital also gets involved and the hybridization changes from sp<sup>2</sup> to sp<sup>3</sup>. As a result, the shape gets changed to tetrahedral.

## 12. Distinguish between a sigma and a pi bond.

### **Solution:**

Sigma (σ) Bond	Pi (π) Bond
(a) It is formed by the end to end	It is formed by the lateral overlap
overlap of orbitals.	of orbitals.
(b) The orbitals involved in the	These bonds are formed by the
overlapping are s–s, s–p, or p–p.	overlap of p-p orbitals only.
(c) It is a strong bond.	It is weak bond.

(d) The electron cloud is symmetrical about the line joining the two nuclei.	The electron cloud is not symmetrical.
(e) It consists of one electron	There are two electron clouds
cloud, which is symmetrical about	lying above and below the plane
the internuclear axis.	of the atomic nuclei.
(f) Free rotation about σ bonds is	Rotation is restricted in case of
possible.	pi-bonds.

The following are the differences between sigma and pi-bonds:

## 13. Explain the formation of $H_2$ molecule on the basis of valence bond theory.

#### **Answer:**

Let us assume that two hydrogen atoms (A and B) with nuclei (NA and NB) and electrons (eA and eB) are taken to undergo a reaction to form a hydrogen molecule.

When A and B are at a large distance, there is no interaction between them. As they begin to approach each other, the attractive and repulsive forces start operating.

#### Attractive force arises between:

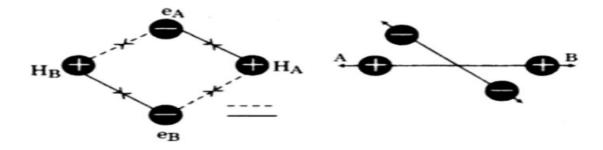
(a) Nucleus of one atom and its own electron i.e., NA – eA and NB – eB.

(b) Nucleus of one atom and electron of another atom i.e., NA - eB and NB - eA.

Repulsive force arises between:

- (a) Electrons of two atoms i.e., eA eB.
- (b) Nuclei of two atoms i.e., NA NB.

The force of attraction brings the two atoms together, whereas the force of repulsion tends to push them apart.

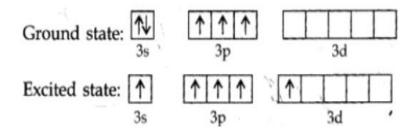


The magnitude of the attractive forces is more than that of the repulsive forces. Hence, the two atoms approach each other. As a result, the potential energy decreases. Finally, a state is reached when the attractive forces balance the repulsive forces and the system acquires minimum energy. This leads to the formation of a dihydrogen molecule.

## 14. Describe the hybridisation in case of PCl5. Why are the axial bonds longer as compared to equatorial bonds?

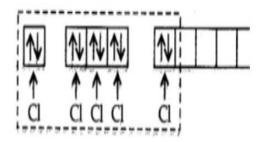
Answer: The ground state and excited state outer electronic

configurations of phosphorus (Z = 15) are:

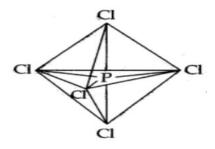


Phosphorus atom is sp3d hybridized in the excited state. These orbitals are filled by the electron pairs donated by five Cl atoms as:

 $PCl_5$ 



The five sp<sup>3</sup>d hybrid orbitals are directed towards the five corners of the trigonal bi pyramidals. Hence, the geometry of PCl<sub>5</sub> can be represented as:



There are five P–Cl sigma bonds in PCl<sub>5</sub>. Three P–Cl bonds lie in one plane and make an angle of 120° with each other. These bonds are called equatorial bonds.

The remaining two P–Cl bonds lie above and below the equatorial plane and make an angle of 90° with the plane. These bonds are called axial bonds.

As the axial bond pairs suffer more repulsion from the equatorial bond pairs, axial bonds are slightly longer than equatorial bonds.

## 15. Define hydrogen bond. Is it weaker or stronger than the van der Waals forces?

#### **Answer:**

A hydrogen bond is defined as an attractive force acting between the hydrogen attached to an electronegative atom of one molecule and an electronegative atom of a different molecule (may be of the same kind).

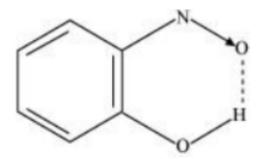
Due to a difference between electronegativities, the bond pair between hydrogen and the electronegative atom gets drifted far away from the hydrogen atom. As a result, a hydrogen atom becomes electropositive with respect to the other atom and acquires a positive charge.

$$4^g - X^{g^-} \dots H^{g^+} - X^{g^-} \dots H^{g^+} - X^{g^-}$$

The magnitude of H-bonding is maximum in the solid state and minimum in the gaseous state.

There are two types of H-bonds:

- (i) Intermolecular H-bond e.g., HF, H<sub>2</sub>O etc.
- (ii) Intramolecular H-bond e.g., o-nitrophenol



Hydrogen bonds are stronger than Van der Walls forces since hydrogen bonds are regarded as an extreme form of dipole-dipole interaction.

## 7marks:

## 1. Define octet rule. Write its significance and limitations.

#### **Answer:**

The octet rule or the electronic theory of chemical bonding was developed by Kossel and Lewis. According to this rule, atoms can combine either by transfer of valence electrons from one atom to another or by sharing their valence electrons in order to attain the nearest noble gas configuration by having an octet in their valence shell.

The octet rule successfully explained the formation of chemical bonds depending upon the nature of the element.

Limitations of the octet theory:

The following are the limitations of the octet rule:

- (a) The rule failed to predict the shape and relative stability of molecules.
- (b) It is based upon the inert nature of noble gases. However, some noble gases like xenon and krypton form compounds such as XeF2, KrF2 etc.

(c) The octet rule cannot be applied to the elements in and beyond the third period of the periodic table. The elements present in these periods have more than eight valence electrons around the central atom. For example: PF<sub>5</sub>, SF<sub>6</sub>, etc.



(d) The octet rule is not satisfied for all atoms in a molecule having an odd number of electrons. For example, NO and NO<sub>2</sub> do not satisfy the octet rule.

$$N = O$$
  $O = N - O$ 

(e) This rule cannot be applied to those compounds in which the number of electrons surrounding the central atom is less than eight. For example, LiCl, BeH<sub>2</sub>, AlCl<sub>3</sub> etc. do not obey the octet rule.

## 2. Discuss the shape of the following molecules using the VSEPR model:

BeCl<sub>2</sub>, BCl<sub>3</sub>, SiCl<sub>4</sub>, AsF<sub>5</sub>, H<sub>2</sub>S, PH<sub>3</sub>

#### **Answer:**

BeCl<sub>2</sub>:

Cl: Be: Cl

The central atom has no lone pair and there are two bond pairs. i.e., BeCl<sub>2</sub> is of the type AB<sub>2</sub>. Hence, it has a linear shape.

BCl<sub>3</sub>:

The central atom has no lone pair and there are three bond pairs. Hence, it is of the type AB<sub>3</sub>. Hence, it is trigonal planar.



SiCl<sub>4</sub>:

The central atom has no lone pair and there are four bond pairs.

Hence, the shape of SiCl4 is tetrahedral being the AB4 type molecule.

### AsF<sub>5</sub>:



The central atom has no lone pair and there are five bond pairs. Hence, AsF<sub>5</sub> is of the type AB<sub>5</sub>. Therefore, the shape is trigonal bipyramidal.

 $H_2S$ :

The central atom has one lone pair and there are two bond pairs. Hence, H<sub>2</sub>S is of the type AB<sub>2</sub>E. The shape is Bent.

PH<sub>3</sub>:

The central atom has one lone pair and there are three bond pairs. Hence, PH<sub>3</sub> is of the AB<sub>3</sub>E type. Therefore, the shape is trigonal pyramidal.

3. Use Lewis symbols to show electron transfer between the following atoms to form cations and anions: (a) K and S (b) Ca and O (c) Al and N.

#### **Answer:**

## (a) K and S:

The electronic configurations of K and S are as follows:

Sulphur (S) requires 2 more electrons to complete its octet. Potassium (K) requires one electron more than the nearest noble gas i.e., Argon. Hence, the electron transfer can be shown as:

### (b) Ca and O:

The electronic configurations of Ca and O are as follows:

Ca: 2, 8, 8, 2

0:2,6

Oxygen requires two electrons more to complete its octet, whereas calcium has two electrons more than the nearest noble gas i.e., Argon. Hence, the electron transfer takes place as:

Ca: 
$$O: Ca^{2+} O: Ca$$

### (c) Al and N:

The electronic configurations of Al and N are as follows:

Al: 2, 8, 3

N: 2, 5

Nitrogen is three electrons short of the nearest noble gas (Neon), whereas aluminium has three electrons more than Neon. Hence, the electron transference can be shown as:

$$Al$$
 $N$ :
 $Al^{3+}$ 
 $N$ :
 $2.8 \quad 2.8$ 
 $2.8 \quad 2.8$ 
 $Al^{3+}$ 
 $N$ :
 $Al^{3+}$ 
 $Al^{3+}$ 

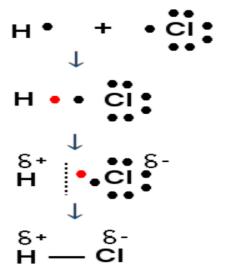
### 4. Explain with the help of suitable example polar covalent bond.

#### **Answer:**

When two dissimilar atoms having different electronegativities combine to form a covalent bond, the bond pair of electrons is not shared equally. The bond pair shifts towards the nucleus of the atom having greater electronegativity. As a result, electron distribution gets distorted and the electron cloud is displaced towards the electronegative atom.

As a result, the electronegative atom becomes slightly negatively charged while the other atom becomes slightly positively charged. Thus, opposite poles are developed in the molecule and this type of a bond is called a polar covalent bond.

HCl, for example, contains a polar covalent bond. Chlorine atom is more electronegative than hydrogen atom. Hence, the bond pair lies towards chlorine and therefore, it acquires a partial negative charge.



5. What is meant by hybridisation of atomic orbitals? Describe the shapes of sp,  $sp^2$ ,  $sp^3$  hybrid orbitals.

#### **Answer:**

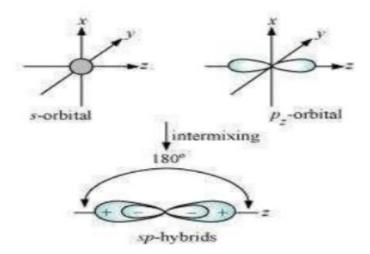
Hybridization is defined as an intermixing of a set of atomic orbitals of slightly different energies, thereby forming a new set of orbitals having equivalent energies and shapes.

For example, one 2s-orbital hybridizes with two 2p-orbitals of carbon to form three new sp<sup>2</sup> hybrid orbitals.

These hybrid orbitals have minimum repulsion between their electron pairs and thus, are more stable. Hybridization helps indicate the geometry of the molecule.

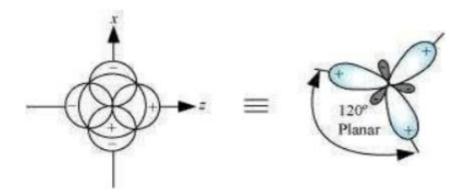
Shape of sp hybrid orbitals: sp hybrid orbitals have a linear shape.

They are formed by the intermixing of s and p orbitals as:



Shape of sp<sup>2</sup> hybrid orbitals:

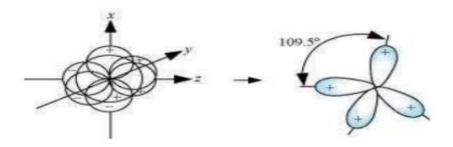
sp<sup>2</sup> hybrid orbitals are formed as a result of the intermixing of one sorbital and two 2p-orbitals. The hybrid orbitals are oriented in a trigonal planar arrangement as:



Shape of sp<sup>3</sup> hybrid orbitals:

Four sp<sup>3</sup> hybrid orbitals are formed by intermixing one s-orbital with three p-orbitals.

The four sp<sup>3</sup> hybrid orbitals are arranged in the form of a tetrahedron as:



6. Draw diagrams showing the formation of a double bond and a triple bond between carbon atoms in  $C_2H_4$  and  $C_2H_2$  molecules.

#### **Answer:**

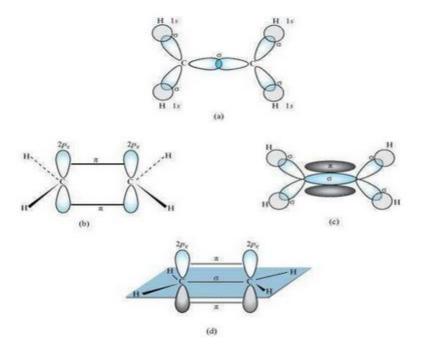
### $C_2H_4$ :

The electronic configuration of C-atom in the excited state is:

$$_{6}C = 1s^{2}2s^{1}2p_{x}^{1}2p_{y}^{1}2p_{z}^{1}$$

In the formation of an ethane molecule ( $C_2H_4$ ), one sp2 hybrid orbital of carbon overlaps a sp2 hybridized orbital of another carbon atom, thereby forming a C-C sigma bond.

The remaining two sp<sup>2</sup> orbitals of each carbon atom form a sp<sup>2</sup>-s sigma bond with two hydrogen atoms. The unhybridized orbital of one carbon atom undergoes sidewise overlap with the orbital of a similar kind present on another carbon atom to form a weak  $\pi$ -bond.

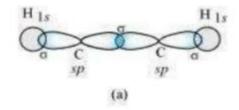


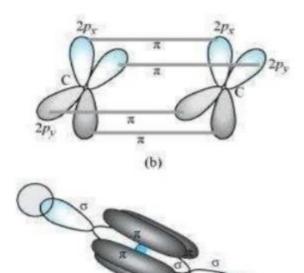
# $C_2H_2$ :

In the formation of  $C_2H_2$  molecule, each C-atom is sp hybridized with two 2p-orbitals in an unhybridized state.

One sp orbital of each carbon atom overlaps with the other along the internuclear axis forming a C–C sigma bond. The second sp orbital of each C–atom overlaps a half-filled 1s-orbital to form a  $\sigma$  bond.

The two unhybridized 2p-orbitals of the first carbon undergo sidewise overlap with the 2p orbital of another carbon atom, thereby forming two pi  $(\pi)$  bonds between carbon atoms. Hence, the triple bond between two carbon atoms is made up of one sigma and two  $\pi$ -bonds.

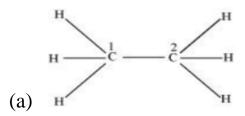




# 7. Which hybrid orbitals are used by carbon atoms in the following molecules?

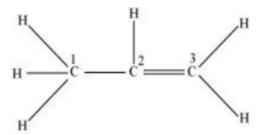
CH<sub>3</sub>-CH<sub>3</sub>; (b) CH<sub>3</sub>-CH=CH<sub>2</sub>; (c) CH<sub>3</sub>-CH<sub>2</sub>-OH; (d) CH<sub>3</sub>-CHO (e) CH<sub>3</sub>COOH

# **Answer:**



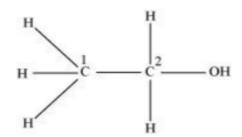
Both  $C_1$  and  $C_2$  are  $sp_3$  hybridized.

(b)



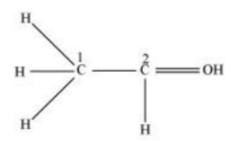
 $C_1$  is  $sp^3$  hybridized, while  $C_2$  and  $C_3$  are  $sp^2$  hybridized.

(c)



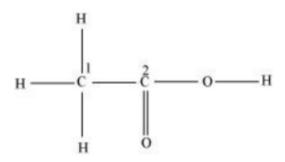
Both  $C_1$  and  $C_2$  are  $sp^3$  hybridized.

(d)



 $C_1$  is  $sp^3$  hybridized and  $C_2$  is  $sp^2$  hybridized.

(e)



 $C_1$  is  $sp^3$  hybridized and  $C_2$  is  $sp^2$  hybridized.

8. Compare the relative stability of the following species and indicate their magnetic properties;

$$O_2,O_2^+,O_2^-$$
 (superoxide),  $O_2^{2-}$  (peroxide)

## **Answer:**

There are 16 electrons in a molecule of dioxygen, 8 from each oxygen atom. The electronic configuration of oxygen molecule can be written as:

$${[\sigma - (1\mathsf{s})]^2}{\left[\sigma^{^*}(1\mathsf{s})\right]^2}{[\sigma(2\mathsf{s})]^2}{\left[\sigma(2\mathsf{p}_z)\right]^2}{\left[\pi(2\mathsf{p}_x)\right]^2}{\left[\pi(2\mathsf{p}_x)\right]^2}{\left[\pi(2\mathsf{p}_x)\right]^2}{\left[\pi^{^*}(2\mathsf{p}_x)\right]^1}$$

Since the 1s orbital of each oxygen atom is not involved in boding, the number of bonding electrons = 8 = Nb and the number of antibonding orbitals = 4 = Na.

Bond order 1/2(Nb - Na)

$$1/2(8-4)=2$$

Similarly, the electronic configuration of  $O_2^+$  can be written as:

$$\mathsf{KK}[\sigma(2\mathsf{s})]^2 \Big[\sigma^{^{\star}}(2\mathsf{s})\Big]^2 \big[\sigma\big(2\mathsf{p}_{_{\mathsf{Z}}}\big)\big]^2 \big[\pi(2\mathsf{p}_{_{\mathsf{Z}}})\big]^2 \big[\pi\big(2\mathsf{p}_{_{\mathsf{Z}}}\big)\big]^2 \Big[\pi^{^{\star}}(2\mathsf{p}_{_{\mathsf{Z}}})\Big]^1$$

Nb = 8

Na = 3

Bond order of  $O_2^+$  = 1/2 (8 - 3)

= 2.5

Electronic configuration of  $\frac{O_2^-}{2}$  ion will be:

$$KK[\sigma(2\mathsf{s})]^2 \Big[\sigma^{^*}(2\mathsf{s})\Big]^2 \Big[\sigma\big(2\mathsf{p}_{_{\mathbb{Z}}}\big)\Big]^2 [\pi(2\mathsf{p}_{_{\mathbb{Z}}})]^2 \Big[\pi\big(2\mathsf{p}_{_{\mathbb{Z}}}\big)\Big]^2 \Big[\pi^{^*}(2\mathsf{p}_{_{\mathbb{Z}}})\Big]^2 \Big[\pi^{^*}(2\mathsf{$$

Nb = 8

Na = 5

Bond order of  $O_2^- = 1/2(8 - 5)$ 

= 1.5

Electronic configuration of  $O_2^{2-}$  ion will be:

$$\mathsf{KK}[\sigma(2\mathsf{s})]^2 \Big[\sigma^*(2\mathsf{s})\Big]^2 \big[\sigma(2\mathsf{p}_{\mathsf{z}})\big]^2 [\pi(2\mathsf{p}_{\mathsf{x}})]^2 \big[\pi(2\mathsf{p}_{\mathsf{y}})\big]^2 \Big[\pi^*(2\mathsf{p}_{\mathsf{y}})\Big]^2 \Big[\pi^*(2\mathsf{p}_{\mathsf{y}})\Big]^2$$

$$Nb = 8$$

$$Na = 6$$

Bond order of 
$$O_2^{2-} = 1/2(8 - 6)$$

$$= 1$$

Bond dissociation energy is directly proportional to bond order. Thus, the higher the bond order, the greater will be the stability. On this basis, the order of stability is

$$\mathrm{O}_2^+ > \mathrm{O}_2 > \mathrm{O}_2^- > \mathrm{O}_2^{2-}$$

9. What is meant by the term bond order? Calculate the bond order of:  $N_2$ ,  $O_2$ ,  $O_2^+$  and  $O_2^-$ .

## **Answer:**

Bond order is defined as one half of the difference between the number of electrons present in the bonding and anti-bonding orbitals of a molecule.

If Na is equal to the number of electrons in an anti-bonding orbital, then Nb is equal to the number of electrons in a bonding orbital.

Bond order = [no. of electrons in bonding MO - no. of electrons in antibonding MO]/2

If Nb > Na, then the molecule is said be stable. However, if Nb  $\leq$  Na, then the molecule is considered to be unstable.

Bond order of  $N_2$  can be calculated from its electronic configuration as:

$$[\sigma(1s)]^2[\sigma^*(1s)]^2[\sigma(2s)]^2[\sigma^*(2s)]^2[\pi(2p_x)]^2[\pi(2p_y)]^2[\sigma(2p_z)]^2$$

Number of bonding electrons, Nb = 10

Number of anti-bonding electrons, Na = 4

Bond order of nitrogen molecule 1/2(10 - 4) = 3

There are 16 electrons in a dioxygen molecule, 8 from each oxygen atom. The electronic configuration of oxygen molecule can be written as:

$$[\sigma - (1s)]^2 [\sigma^*(1s)]^2 [\sigma(2s)]^2 [\sigma^*(2s)]^2 [\sigma(1p_z)]^2 [\pi(2p_x)]^2 [\pi(2p_y)]^2 [\pi^*(2p_x)]^1 [\pi^*(2p_y)]^1$$

Since the 1s orbital of each oxygen atom is not involved in boding, the number of bonding electrons = 8 = Nb and the number of antibonding electrons = 4 = Na.

Bond order 1/2(8-4)

$$=2$$

Hence, the bond order of oxygen molecule is 2.

Similarly, the electronic configuration of  $O_2^+$  can be written as:

$$\mathsf{KK}[\sigma(2\mathsf{s})]^2 \Big[\sigma^{^{\star}}(2\mathsf{s})\Big]^2 \big[\sigma\big(2\mathsf{p}_{_{\mathsf{Z}}}\big)\big]^2 \big[\pi(2\mathsf{p}_{_{\mathsf{Z}}})\big]^2 \big[\pi\big(2\mathsf{p}_{_{\mathsf{Z}}}\big)\big]^2 \Big[\pi^{^{\star}}(2\mathsf{p}_{_{\mathsf{Z}}})\Big]^1$$

$$Nb = 8$$

$$Na = 3$$

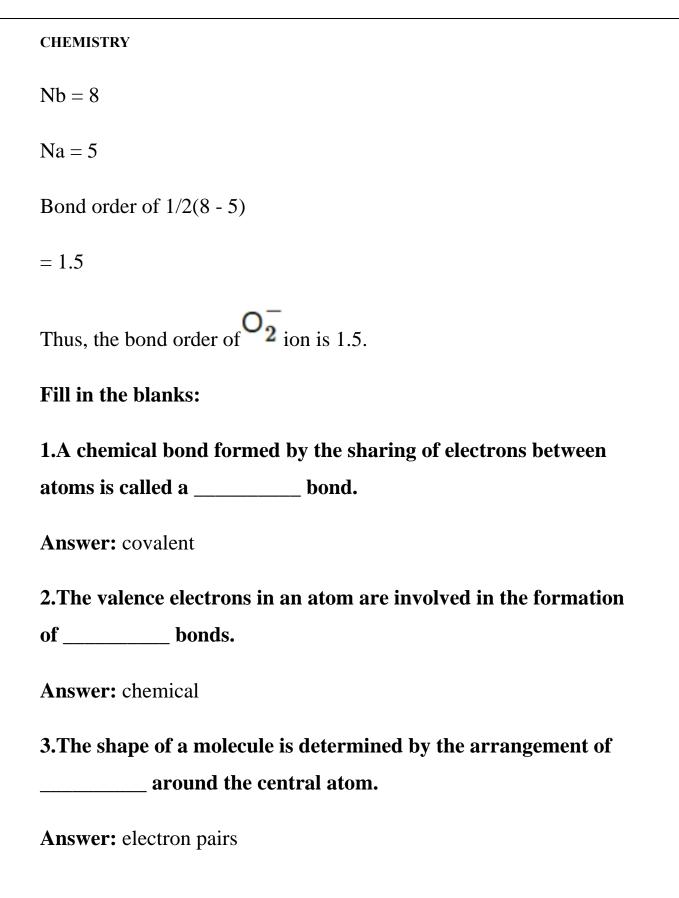
Bond order of 1/2(8 - 5)

$$= 2.5$$

Thus, the bond order of  $O_2^+$  is 2.5.

The electronic configuration of  $O_2^-$  ion will be:

$$KK[\sigma(2\mathsf{s})]^2 \Big[\sigma^*\,(2\mathsf{s})\Big]^2 \big[\sigma\big(2\mathsf{p}_{\mathsf{z}}\big)\big]^2 [\pi(2\mathsf{p}_{\mathsf{z}})]^2 \big[\pi\big(2\mathsf{p}_{\mathsf{y}}\big)\big]^2 \Big[\pi^*\,(2\mathsf{p}_{\mathsf{y}}\big)\Big]^2 \Big[\pi^*\,(2\mathsf{p}_{\mathsf{y}}\big)\Big]^1$$



4. The sharing of one pair of electrons between two atoms results
in a bond.
Answer: single
5.The electrostatic force of attraction between oppositely charged
ions is the basis of a bond.
Answer: ionic
6.The concept of resonance is used to explain the delocalization of in a molecule.
Answer: electrons
7.A bond formed by the transfer of electrons from one atom to another is known as an bond.
Answer: ionic
8.The bond angle in a water molecule (H2O) is approximately degrees.
<b>Answer:</b> 104.5
9.A molecule with a trigonal planar shape has electron pairs around the central atom.
Answer: three

10.In a polar covalent bond, the shared electrons are pulled closes to the atom with the higher
Answer: electronegativity
Multiple choice:
1. What type of bond is formed by the transfer of electrons from one atom to another?
a) Covalent bond
b) Polar bond
c) Ionic bond
d) Metallic bond
Answer:
c) Ionic bond
2.In a covalent bond, electrons are:
a) Shared
b) Transferred
c) Donated

CHEMISTRY
d) Lost
Answer:
a) Shared
3. Which of the following is a diatomic molecule?
a) CO
b) H <sub>2</sub> O
c) O <sub>2</sub>
d) CH <sub>4</sub>
Answer:
c) O <sub>2</sub>
4.The VSEPR theory is used to predict:
a) Molecular weight
b) Molecular shape
c) Bond strength
d) Electron affinity
Answer:
b) Molecular shape

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5.A sigma ( $\sigma$ ) bond is formed by the overlap of:
a) p orbitals
b) s orbitals
c) d orbitals
d) $\pi$ orbitals
Answer:
b) s orbitals
6. The bond angle in a tetrahedral molecule is approximately:
a) 90 degrees
b) 109.5 degrees
c) 120 degrees
d) 180 degrees
Answer:
b) 109.5 degrees

7.In a polar covalent bond, the electronegativity difference

between atoms is:

a) Zero

b) Small
c) Large
d) Constant
Answer:
c) Large
8. Which molecule exhibits hydrogen bonding?
a) CH <sub>4</sub>
b) H <sub>2</sub> O
c) CO <sub>2</sub>
d) $N_2$
Answer:
b) H <sub>2</sub> O
9.The concept of resonance is used to describe:
a) Electron cloud distribution
b) Multiple bond formation
c) Molecular weight

d) HybridizationAnswer:b) Multiple bond formation

10. What is the shape of a molecule with the formula AB<sub>4</sub>E?

a) Linear

**CHEMISTRY** 

b) Trigonal planar

c) Tetrahedral

d) Bent

**Answer:** 

d) Bent

# **Summary:**

The chapter on "Chemical Bonding and Molecular Structure" explores the fundamental principles governing the formation of chemical bonds and the three-dimensional arrangement of atoms in molecules.

The chapter begins by introducing the concept of chemical bonding, emphasizing the role of valence electrons. It delves into various types of chemical bonds, such as covalent and ionic bonds. In a covalent bond, electrons are shared between atoms, leading to the formation of molecules. On the other hand, ionic bonds result from the transfer of electrons between atoms, creating ions that are held together by electrostatic forces.

The VSEPR (Valence Shell Electron Pair Repulsion) theory is introduced to explain the molecular geometry based on the repulsion between electron pairs around the central atom. This theory helps predict the shapes of molecules and molecular ions.

The chapter explores the different types of hybridization, such as sp<sup>3</sup>, sp<sup>2</sup>, and sp, which play a crucial role in determining the shapes of molecules. Molecular orbital theory is also introduced to explain the formation of molecular orbitals through the overlap of atomic orbitals.