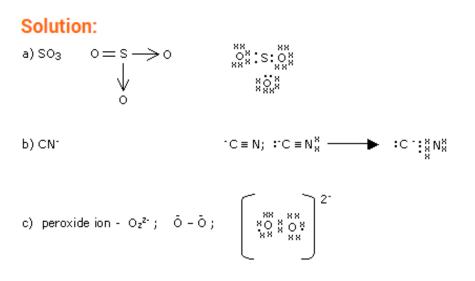


Chapter 4. Chemical Bonding and Molecular Structure

Question-1

Write Lewis dot structure of SO₃, CN⁻, peroxide ion.



Question-2

Discuss the shapes of the following molecules using the VSEPR model: (a) SF_6 , (b) XeF_2 , (c) NH_4^+ , (d) ICl_4^- , (e) BrF_3 .

Solution:

(a) SF₆: Number of electrons around S

= (Valence electron of S+ electrons donated by Fluoride)

= 6 + 6 = 12

Number of bond pairs = 12/2 = 6

To have minimum repulsion between bond pairs structure of SF_6 is octahedran. Shape: Octahedral, Hybridisation= Sp^3d^2 .

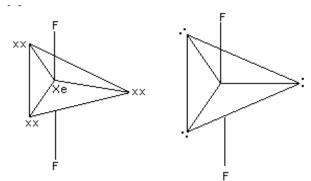
(b) XeF_2 : Number of bond pairs = (8+2)/2=5 pairs

Bond pairs=2

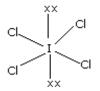
Lone pairs=3

Hence molecule has trigonal pyramidal structure.

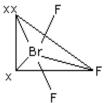
..Shape: trigonal bipyramidal



(c) NH₄⁺: Electrons (5+4-1)=8; Bond pairs=8/2=4 Hence tetra hedral:



(d) ICl₄: (7+4+1)=12/2=6 pairs: octahedral structure only four bond pairs: (i.e.) shape=square planar
(e) BrF₃: 7+3=10 electrons; 5 pairs: 3 bond pairs; 2 lone pairs. 5 pairs: trigonal bipyramidal structure: Sp³d hybridization. Shape (To have minimum repulsion, it has bent T shape)



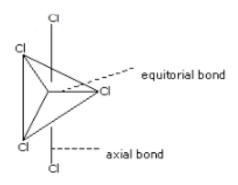
To get structure or hybridization, bonds formed by bond pairs with central atoms and the lone pairs should be considered.

Discuss the shape of following molecule through valence bond theory. PCI₅.

Solution:

(PCI₅) Electronic configuration of P in ground state = 1s²2s²3s².3p³ In exited state = 1s²2s²3s¹.3p³3d¹.

Five orbitals, one 3s, three 3orbitals, one 3d hybridizes to give sp³d orbitals. Five Cl atoms, each gives one electron to the sp³d orbitals and forms a covalent bond. sp³d orbitals are directed towards a corner of a trigonal bipyramidal structure.



Axial bonds are longer than equatorial bonds to have minimum repulsion between axial and equatorial bonds.

Question-4

Write order of reactivity of HF, HBr, HCl and HI.

Solution:

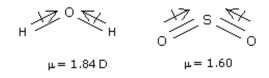
Greater the difference in electro negativity between two atoms, stronger will be the bond strength. Difference in electro negativity is as per following order

HF > HBr > HCl > HI Hence, lesser the difference in electro negativity, greater will be bond strength reactivity is of the following order HI > HCl > HBr > HF.

H₂O and SO₂ have dipole moments. Explain the shape.

Solution:

A molecule having two bonds, has certain dipole moment indicates that the two bond dipoles do not cancel each other but add up to give a net dipole moment to the molecule. Such molecule is unsymmetric. It has bent structures.

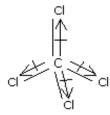


Question-6

Why does CH₄, CCl₄, SiF₄ and SnCl₄ have zero dipole moments?

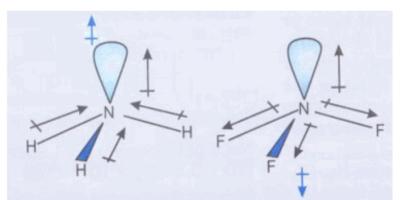
Solution:

All of the above molecules have polar bonds. Due to symmetrical structure, dipole moment of one bond is cancelled by the opposite dipole moment of the other bonds.



Why NF₃ has less dipole moment than NH₃?

Solution:



In N-H bond direction of dipole moment is towards hydrogen. Resultant dipole moment is in the same direction of dipole moment due to lone pair of N: But in NF₃, resultant dipole moment is in opposition direction to that of dipole moment due to lone pair of N. Hence dipole moment of NF₃ has less dipole moment than NH₃.

Question-8

Why AlCl₃ is largely covalent while aluminium fluoride is largely ionic?

Solution:

Fluoride is more electronegative than chlorine. Hence the difference in electro negativity of Al and Fluorine is more (2.31), above 2. Hence H - F bond is ionic. In AlCl₃, the difference in electro negativity between Al and chlorine is 1.55, which is less than two, and then bond will be more covalent than ionic.

Why is Pi bond weaker than a sigma bond?

Solution:

Pi bonds between atoms are formed due to lateral overlap of atomic orbitals. Which involves effective overlap results in the formation of weaker bond.

Sigma bonds are formed due to axial overlap of atomic orbitals, which involves effective overlapping. Hence sigma bonds are stronger.

Question-10

Why compounds with C = C double bonds have geometrical isomerism?

Solution:

Double bonds consist of one sigma and one Pi bond. Free rotation about a sigma bond is possible because the electron densities lies along the inter nuclear axis. But free rotation about a Pi bond is hindered as the two carbon atoms are connected above and below an axis. Thus positions of groups in carbon compounds contain double bonds are fixed. This leads to geometrical isomerism.

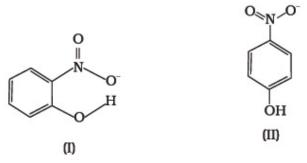


Rotation above π - bond is not possible because it would break the π - bond.

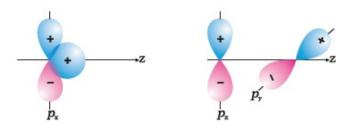
Chemical Bonding & Molecular Structure

Short Answer Type Questions

- 1. Explain the non linear shape of H2S and non planar shape of PCl₃ using valence shell electron pair repulsion theory.
- 2. Using molecular orbital theory, compare the bond energy and magnetic character of O_2^+ and O_2^- species.
- Explain the shape of BrF₅.
- 4. Structures of molecules of two compounds are given below :



- (a) Which of the two compounds will have intermolecular hydrogen bonding and which compound is expected to show intramolecular hydrogen bonding.
- (b) The melting point of a compound depends on, among other things, the extent of hydrogen bonding. On this basis explain which of the above two compounds will show higher melting point.
- (c) Solubility of compounds in water depends on power to form hydrogen bonds with water. Which of the above compounds will form hydrogen bond with water easily and be more soluble in it.
- 5. Why does type of overlap given in the following figure not result in bond formation?



- 6. Explain why PCl_5 is trigonal bipyramidal whereas IF_5 is square pyramidal.
- In both water and dimethyl ether (CH₃ Ö CH₃), oxygen atom is central atom, and has the same hybridisation, yet they have different bond angles.
 Which one has greater bond angle? Give reason.
- 8. Write Lewis structure of the following compounds and show formal charge on each atom. HNO_3 , NO_2 , H_2SO_4
- 9. The energy of $\sigma 2p_z$ molecular orbital is greater than $\pi 2p_x$ and $\pi 2p_y$ molecular orbitals in nitrogen molecule. Write the complete sequence of energy levels in the increasing order of energy in the molecule. Compare the relative stability and the magnetic behaviour of the following species :
 - N_2 , $N_2^{+>/sup>}$, N_2^{-} , N_2^{2+}
- 10. What is the effect of the following processes on the bond order in N_2 and O_2 ?
 - \circ (i) N₂ \rightarrow N₂⁺ + e⁻
 - \circ (ii) $O_2 \rightarrow O_2^+ + e^-$
- 11. Give reasons for the following :
 - (i) Covalent bonds are directional bonds while ionic bonds are nondirectional.
 - (ii) Water molecule has bent structure whereas carbon dioxide molecule is linear.
 - (iii) Ethyne molecule is linear.
- 12. What is an ionic bond? With two suitable examples explain the difference between an ionic and a covalent bond?
- 13. Arrange the following bonds in order of increasing ionic character giving reason. N–H, F–H, C–H and O–H

- 14. Explain why CO₃²⁻ ion cannot be represented by a single Lewis structure. How can it be best represented?
- 15. Predict the hybridisation of each carbon in the molecule of organic compound given below. Also indicate the total number of sigma and pi bonds in this molecule.

$$CH = C - C - CH_2 - C OH$$

16. Group the following as linear and non-linear molecules :

H₂O, HOCl, BeCl₂, Cl₂O

- 17. Elements X, Y and Z have 4, 5 and 7 valence electrons respectively.
 - (i) Write the molecular formula of the compounds formed by these elements individually with hydrogen.
 - (ii) Which of these compounds will have the highest dipole moment?
- 18. Draw the resonating structure of
 - (i) Ozone molecule
 - (ii) Nitrate ion
- 19. Predict the shapes of the following molecules on the basis of hybridisation. BCl_3 , CH_4 , CO_2 , NH_3
- 20. All the C–O bonds in carbonate ion (CO_3^{2-}) are equal in length. Explain.
- 21. What is meant by the term average bond enthalpy? Why is there difference in bond enthalpy of O-H bond in ethanol (C_2H_5OH) and water?

Matching Type Questions

1. Match the species in Column I with the type of hybrid orbitals in Column II.

	Column I	Column II	
(i)	SF ₄	(a)	sp^3d^2
(ii)	IF ₅	(b)	d^2sp^3
(iii)	NO_2^+	(c)	$sp^{3}d$
(iv)	NH_4^+	(d)	sp^3
		(e)	sp

2. Match the species in Column I with the geometry/shape in Column II.

	Column I	Column II	
(i)	H_3O^{\dagger}	(a)	Linear
(ii)	$HC \equiv CH$	(b)	Angular
(iii)	ClO ₂	(c)	Tetrahedral
(iv)	NH_4^+	(d)	Trigonal bipyramidal
		(e)	Pyramidal

3. Match the species in Column I with the bond order in Column II.

	Column I	Column II
(i)	NO	(a) 1.5
(ii)	CO	(b) 2.0
(iii)	O_2^-	(c) 2.5
(iv)	O ₂	(d) 3.0

4. Match the items given in Column I with examples given in Column II.

	Column I	Column II
(i)	Hydrogen bond	(a) C
(ii)	Resonance	(b) LiF
(iiii)	Ionic solid	(c) H ₂
(iv)	Covalent solid	(d) HF
		(e) O ₃

5. Match the shape of molecules in Column I with the type of hybridisation in Column II.

	Column I	Column II
(i)	Tetrahedral	(a) <i>sp</i> ²
(ii)	Trigonal	(b) <i>sp</i>
(iii)	Linear	(c) sp ³

Assertion and Reason Type Questions

In the following questions a statement of Assertion (A) followed by a statement of Reason (R) is given. Choose the correct option out of the choices given below each question.

1. Assertion (A) : Sodium chloride formed by the action of chlorine gas on sodium metal is a stable compound.

Reason (R) : This is because sodium and chloride ions acquire octet in sodium chloride formation.

- (i) A and R both are correct, and R is the correct explanation of A.
- (ii) A and R both are correct, but R is not the correct explanation of A.
- (iii) A is true but R is false.
- (iv) A and R both are false.
- 2. Assertion (A) : Though the central atom of both NH_3 and H_2O molecules are sp3 hybridised, yet H-N-H bond angle is greater than that of H-O-H.

Reason (R) : This is because nitrogen atom has one lone pair and oxygen atom has two lone pairs.

- (i) A and R both are correct, and R is the correct explanation of A.
- (ii) A and R both are correct, but R is not the correct explanation of A.
- (iii) A is true but R is false.
- (iv) A and R both are false.
- 3. Assertion (A): Among the two O-H bonds in H2O molecule, the energy required to break

the first O-H bond and the other O-H bond is the same.

Reason (R) : This is because the electronic environment around oxygen is the same even after breakage of one O-H bond.

- (i) A and R both are correct, and R is correct explanation of A.
- (ii) A and R both are correct, but R is not the correct explanation of A.
- (iii) A is true but R is false.
- (iv) A and R both are false.

Long Answer Type Questions

1. i) Discuss the significance/ applications of dipole moment.

- Use the molecular orbital energy level diagram to show that N₂ would be expected to have a triple bond, F₂, a single bond and Ne₂, no bond.
- 3. Briefly describe the valence bond theory of covalent bond formation by taking an example of hydrogen. How can you interpret energy changes taking place in the formation of dihydrogen?
- Describe hybridisation in the case of PCl₅ and SF₆. The axial bonds are longer as compared to equatorial bonds in PCl₅ whereas in SF₆ both axial bonds and equatorial bonds have the same bond length. Explain.
- 5. 64. (i) Discuss the concept of hybridisation. What are its different types in a carbon atom.(ii) What is the type of hybridisation of carbon atoms marked with star.

(a)
$$\stackrel{*}{CH}_{2} = CH - \stackrel{\parallel}{*C} - O - H$$

0

(c)
$$CH_3 - CH_2 - CH_2 - H$$

⁽ii) Represent diagrammatically the bond moments and the resultant dipole moment in CO_2 , NF₃ and CHCl₃.

- (d) $\overset{*}{C}H_3 CH = CH CH_3$
- (e) $CH_3 \mathring{C} \equiv CH$

Comprehension given below is followed by some multiple choice questions. Each question has one correct option. Choose the correct option.

Molecular orbitals are formed by the overlap of atomic orbitals. Two atomic orbitals combine to form two molecular orbitals called bonding molecular orbital (BMO) and anti bonding molecular orbital (ABMO). Energy of anti bonding orbital is raised above the parent atomic orbitals that have combined and the energy of the bonding orbital is lowered than the parent atomic orbitals. Energies of various molecular orbitals for elements hydrogen to nitrogen increase in the order :

$$\begin{split} &\sigma 1s < \sigma^* \ 1s < \sigma 2s < \sigma^* \ 2s < (\pi 2p_x \approx \pi 2p_y) < \sigma 2p_z < (\pi^* \ 2p_x \approx \pi^* \ 2p_y) < \sigma^* \ 2p_z \text{ and} \\ &\text{for oxygen and fluorine order of energy of molecular orbitals is given below :} \\ &\sigma 1s < \sigma^* \ 1s < \sigma 2s < \sigma^* \ 2s < \sigma 2p_z < (\pi 2p_x \approx \pi 2p_y) < (\pi^* 2p_x \approx \pi^* 2p_y) < \sigma^* \ 2p_z \end{split}$$

Different atomic orbitals of one atom combine with those atomic orbitals of the second atom which have comparable energies and proper orientation. Further, if the overlapping is head on, the molecular orbital is called 'Sigma', (σ) and if the overlap is lateral, the molecular orbital is called 'pi', (π). The molecular orbitals are filled with electrons according to the same rules as followed for filling of atomic orbitals. However, the order for filling is not the same for all molecules or their ions. Bond order is one of the most important parameters to compare the strength of bonds.

- 6. Which of the following statements is correct?
 - (i) In the formation of dioxygen from oxygen atoms 10 molecular orbitals will be formed.
 - (ii) All the molecular orbitals in the dioxygen will be completely filled.
 - (iii) Total number of bonding molecular orbitals will not be same as total number of anti bonding orbitals in dioxygen.

- (iv) Number of filled bonding orbitals will be same as number of filled antibonding orbitals.
- 7. Which of the following molecular orbitals has maximum number of nodal planes?
 - (i) σ^{*}1s
 - \circ (ii) $\sigma^* 2p_z$
 - ∘ (ііі) п2р_х
 - ∘ (iv) п^{*}2р_у
- 8. Which of the following pair is expected to have the same bond order?
 - \circ (i) O₂ , N₂ \circ (ii) O₂⁺, N₂⁻ \circ (iii) O₂⁻ , N₂⁺
 - (iv) O₂⁻, N₂⁻
- 9. In which of the following molecules, $\sigma 2p_z$ molecular orbital is filled after $n2p_x$ and $n2p_y$ molecular orbitals?
 - (i) O₂
 - (ii) Ne₂
 - ∘ (iii) N₂
 - (iv) F₂

CBSE Class 11 Chemistry Important Questions Chapter 4 Chemical Bonding and Molecular Structure

1 Marks Questions

1.Define a chemical bond.

Ans. The attractive force which holds various constituents (atoms, ions etc.) together in different chemical species is called a chemical bond.

2. Give the main feature of Lewis approach of chemical bonding.

Ans. Lewis postulated that atoms achieve the stable octet when they are linked by chemical bonds. He assured that atoms are positively charged centre and the outer shell that could accommodate a maximum of eight electrons. These electrons occupy the corners of a cube which surrounds the centre. Lewis introduced simple notations to represent valence electrons in an atom called Lewis symbol

3.Write electron dot structure (Lewis structure) of Na, Ca, B, Br, Xe, As, Ge, N³⁻.

Na[•], Ca[:], $\cdot \dot{B}$ [•], \dot{B} [•], $\cdot \dot{X}_{e}$ [•], $\cdot \dot{A}_{s}$ [•], $\cdot \dot{G}_{e}$ [•], $(:N^{:})^{3^{-}}$ 4. Give the octet rule in

short.

Ans.The atoms tend to adjust the arrangement of their electrons in such a way that they (except H and He) achieve eight electrons in their outermost shell. This is known as the octet rule.

5.Define an ionic bonding. [?]

Ans.An ionic bond (or electrovalent bond) is formed by a complete transfer of one or more of outer most electrons from the atom of a metal to that of a non – metal.

6. Which one of the following has the highest bond order? N_2 , N_2^+ or N_2^- .

Ans. N_2 has the highest bond order.

7.Define bond order.

Ans. Bond order is defined as number of bonds between two atoms in a molecule.

8.What type of bond is formed when atoms have high difference of electornegativity?

Ans. Electrovalent or ionic bond.

9.Define dipole moment.

Ans. Dipole moment is defined as the product of the magnitude of the charge and the distance between the centers of positive and negative charge.

10. Give the mathematical expression of dipole moment.

Ans. Mathematically dipole moment is expressed as dipole moment (M) = charge (Q) x distance of separation (r). Dipole moment is usually expressed in Debye units (D).

11.Why is dipole moment of CO₂, BF₃, CCl₄ is zero?

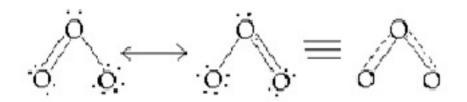
Ans. Because there molecules have symmetrical shapes and thus the dipoles gets cancelled and the net dipole moment is zero.

12.Why is BF₃ non – polar?

Ans. Because BF_3 has symmetrical shape, the net dipole moment is zero and thus it is non – polar.

13. Write the resonating structure of O_3 molecule.

Ans.



14.What is sigma bond?

Ans.A covalent bond formed due to the overlap of orbitals of the two atoms along the line going the two nuclei (orbital axis) is called sigma (s) bond.

15.What is pi – bond?

Ans. A covalent bond formed between the two atoms due to the sideways overlap of their p – orbitals is called a pi (π) bond.

16.How many s – and π - bond are there in a molecule of C₂H₄ (ethene)?

Ans. In a molecule of ethene, there are 5 s - bonds (one between C-C , and four between C-H and one π - bond.

17. How many s - and π - bonds are there in a molecule of CH₂ = CH – CH = CH₂ ?

Ans. There are 9 s - bonds (three between C – C and 6 between C – H) and 2 π - bonds.

18.What type of bond exists in multiple bond (double / triple)?

Ans. pi (p) – bond is always present in molecules containing multiple bond.

19.What type of bond are formed due to orbital overlap?

Ans. Covalent bonds are formed due to the overlap of certain orbitals that are oriented favourably in the space.

20.How do covalent bonds form due to orbital overlapping?

Ans. According to orbital overlap concept, the formation of a covalent bond between two atoms results by pairing of electrons present in the valence shell having opposite spins.

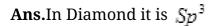
21.Define hybridisation.

Ans. Hybridisation is defined as the process of intermixing of the orbitals of slightly different energies so as to redistribute their energies, resulting in the formation of new set of orbitals of equivalent energies and shape.

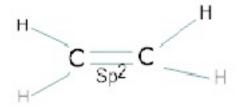
22.State the hybrid orbitals associated with B in BCl_3 and C in $\mathrm{C_2H_4}$

Ans. (i) Sp^2 hybridization (ii) Sp^2 hybridization.

23.What is the state of hybridization of carbon atoms in diamond and graphite?



In graphite it is Sp^2



24.What type of hybridisation takes place in (i) p in PCL_5 and (ii) S in S F_6 ?

Ans. (i) Sp^3d

(ii)
$$Sp^{3}d^{2}$$
.

25.Define bonding molecular orbital.

Ans. The molecular orbital formed by the addition of atomic orbitals is called bonding

molecular orbital.

$$\sigma = \Psi A + \Psi B$$

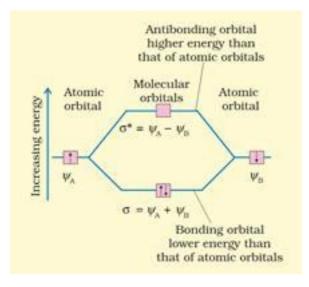
26.Define antibonding molecular orbital.

Ans. The molecular orbital formed by the subtraction of atomic orbitals is called antibonding molecular orbital.

 $\sigma^+ = \Psi A - \Psi B \cdot$

27.Explain diagrammatically the formation of molecular orbital by LCAO.

Ans. The molecular orbital formed by subtraction of atomic orbital is called antibonding molecular orbital.



28. Which one O_2^{-1} and O_2^{2-1} may exhibit paramagnetism?

Ans. O_2 would exhibit paramagnetism because it contains one unpaired electron in its Mo configuration.

29.Why are bonding molecular orbitals more stable than antibonding molecular orbitals?

Ans. Bonding molecular orbital has lower energy and hence greater stability than the

corresponding antibonding molecular orbital.

30.Define bond order.

Ans. Bond order (b.o) is defined as one half the difference between the number of electrons present in the bonding and the antibonding orbitals i.e;

Bond order (b.o) = $\frac{1}{2} (N_b - N_a)$

If $N_b > N_a$, molecule is stable and

If $N_b < N_a$, molecule is unstable.

31.Define hydrogen bonding

Ans. Hydrogen bond can be defined as the attractive force which binds hydrogen atom of one molecule with the electronegative atom (F, O or N) of another molecule.

32.What are the types of H-bonding? Which of them is stronger?

Ans. (i)Inter-molecular H-bonding

(ii) Intra molecular H-bonding. Inter molecular H-bonding is stronger than intra-molecular H-bonding.

33. NH_3 has higher boiling point than PH_3 . Give reason.

Ans.In $N\!H_3$, there is hydrogen bonding whereas in PH_3 there is no hydrogen bonding.

34.Define electrovalent bond.

Ans. The bond formed, as a result of the electrostatic attraction between the positive and negative ions are termed as the electrovalent bond.

CBSE Class 12 Chemistry Important Questions Chapter 4 Chemical Bonding and Molecular Structure

2 Marks Questions

1. Give the main feature of Kossel's explanation of chemical bonding.

Ans. Kossel in relation to chemical bonding drew attention to the following facts -

(i) In the periodic table, the highly electronegative halogens and the highly electropositive alkali metals are separated by the noble gases.

(ii) In the formation of a negative ion from a halogen atom and a positive ion from an alkali metal, atom is associated with a gain and loss of an electron by the respective atoms.

(iii) The negative and positive ions so formed attain stable noble gas electronic configurations. The noble gases have particularly eight electrons, $ns^2 np^6$.

The –ve and +ve ions are stabilized by electrostatic attraction.

2.How can you explain the formation of NaCl according to kossel concept?

Ans. The formation of NaCl from sodium and chlorine can be explained as

Na ® Na⁺ + e⁻

[Ne] 3s¹ ® [Ne]

Cl + e⁻ ® Cl⁻

[Ne] $3s^2 3p^5$. [Ne] $3s^2 3p^6$ or [Ar]

 $Na^{+} + Cl^{-}$ ® $Na^{+} Cl^{-}$ or NaCl.

3.Write the significance of octet rule.

Ans. Octet rule signifies –

(i) It is useful for understanding the structures of most of the organic compounds.

It mainly applies to the second period elements of the periodic table.

4.Write the Lewis structure for CO molecule

Ans. (i) The outer (valence) shell configurations of carbon and oxygen atoms are

Carbon : (6) – $1s^2 2s^2 2p^2$

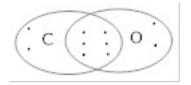
Oxygen : (8) $- 1s^2 2s^2 2p^4$.

The valence electrons (4 + 6 = 10)

:C :O: or :C ---- O:

But it does not complete octet, thus multiple bond is exhibited.

Thus,



 $\circ r : \subset \equiv \circ :$

(ii) N (2s² 2p³), O (2s² 2p⁴)

5 + (2 x 6) + 1 = 18 electrons.

Thus,

$$[0: N: 0:] \text{ or } [0 = N - 0:]$$

or, $[0 - N = 0]$

5. Give the Lewis dot structure of HNO_3

Ans. HNO₃ ®

$$\begin{bmatrix} 0 & \stackrel{+}{N} & 0 & H \\ 0 & \stackrel{-}{O} & \stackrel{-}{\Box} & \stackrel{-}{O} & \stackrel{+}{N} & 0 & - & H \\ 0 & \stackrel{-}{\Box} & \stackrel{-}{\Box} & \stackrel{-}{\Box} & \stackrel{-}{\Box} & \stackrel{-}{\Box} \end{bmatrix}$$

6.What changes are observed in atoms undergoing ionic bonding?

Ans. Due to the electron transfer the following changes occurs -

(i) Both the atoms acquire stable noble gas configuration.

(ii) The atom that loses electrons becomes +vely charged called cation whereas that gains electrons becomes –vely charged called anion.

(iii) Cation and anion are held together by the coulombic forces of attraction to form an ionic bond.

7.Mention the factors that influence the formation of an Ionic bond.

Ans.Ionic bond formation mainly depends upon three factors -

(i) Low ionization energy – elements with low ionization enthalpy have greater tendency to form an ionic bonds.

(ii) High electron gain enthalpy – high negative value of electron gain enthalpy favours ionic bond.

(iii) Lattice energy – high lattice energy value favours ionic bond formation.

8. Give reason why H_2^+ ions are more stable than H_2^- though they have the same bond order.

Ans.In H_2^- ion, one electron is present in anti bonding orbital due to which destabilizing effect is more and thus the stability is less than that of H_2^+ ion.

9. How would the bond lengths vary in the following species? C_2 , $C_2^- C_2^{-2-}$.

Ans.The order of bond lengths in C_2 , C_2^- and C_2^{2-} is $C_2 > C_2^- > C_2^{2-}$.

10.Out of covalent and hydrogen bonds, which is stronger.

Ans. Covalent bond.

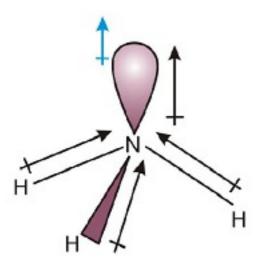
11.Define covalent radius.

Ans. The covalent radius is measured approximately as the radius of an atom's core which is in contact with the core of an adjacent atom in a bonded situation.

eg H \longrightarrow F:

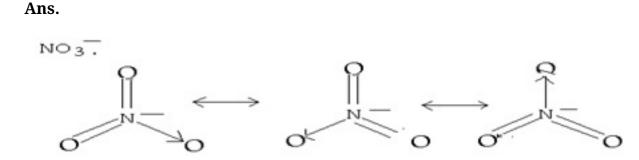
12.Why NH₃ has high dipole moment than NF₃ though both are pyramidal?

Ans. In case of NH_3 the orbital dipole due to lone pair is in the same direction as the resultant dipole moment of the N-H bonds, whereas in NF_3 the orbital dipole is in the direction opposite to the resultant dipole moment of the three N-F



bonds. The orbital dipole become of lone pair decreases, which results in the low dipole moment.

13. Draw the resonating structure of $\mathrm{NO_3}^-$



14.On which factor does dipole moment depend in case of polyatomic molecules.

Ans.The dipole moment of the polyatomic molecule depends on individual dipole moments of bonds and also on the spatial arrangement of various bonds in the molecule.

15.Dipole moment of Be F₂ is zero. Give reason.

Ans. In BeF₂ the dipole moment is zero because the two equal bond dipoles point in opposite directions and cancel the effect of each other.

$\begin{array}{c} F \longrightarrow Be \longrightarrow F \\ \longleftrightarrow & + \longrightarrow \end{array}$

16.Bond dipoles in Be F₂

Give the main features of VSEPR Theory.

Ans. The main postulates of VSEPR theory are as follows :

(i)The shape of a molecule depends upon the number of valence shell electron pairs around the central atom.

(ii)Pairs of electrons in the valence shell repel one another since their electron clouds are negatively charged.

(iii)There pairs of electrons tends to occupy such position in space that minimize repulsion and thus maximize distance between them.

(iv)The valence shell is taken as a sphere with the electron pairs localizing on the sphere at maximum distance from one another.

(v)A multiple bond is treated as it is a single electron pair and two or three electron pairs of a multiple bond is treated as super pair.

(vi)When two or more resonance structures can represent a molecule, the VSEPR nodal is applicable to any such structure.

17.What's difference between lone pair and bonded pair of electrons?

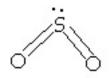
Ans. Lone pair electrons do not take part in bond formation whereas bond pair electrons take part in bond formation.

$18.CO_2$ is linear whereas SO_2 is bend – shaped. Give reason.

Ans. In CO₂, the bond electron are furtherest away from each other forming 180^0 angle. Thus,

CO₂ is linear.

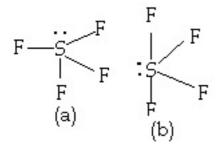
In SO₂, the number of bonding pairs is 4 where it has an lone pair of electron which does not participate in bond formation thereby repulsive strain is experienced.



19.Why does H₂O have bent structure?

Ans. In water molecule, there are two bonding pairs and two lone pairs of electrons. The shape should have been tetrahedral if there were all bp but two lp are present. Thus the shaped is distorted to an angular shape. Because lp – lp repulsion is more than lp – bp repulsion.

20.For the molecule,



Why is structure (b) more stable than structure (a)?

Ans. In (a) the lp is present at axial position so there are three lp - bp repulsions at 90⁰. Whereas in (b) the lp is in an equatorial position are there are two lp - bp repulsions. Hence, arrangement (b) is more stable than (a).

21.How would you attribute the structure of PH₃ molecule using VSEPR model?

Ans. Phosphorus atom has 5 electrons in its outermost orbit. H – atoms contribute one electron each to make in all 8 electron around P – atom. Thus 4 pairs of electrons would be distributed in a tetrahedral manner around the central atom. Three pairs from three P – H

bonds while the fourth pair remains unused. Due to repulsion between the bp and lp, the shape is not of tetrahedral but trigonal pyramidal molecule.

22.In SF_4 molecule, the lp electrons occupies an equatorial position in the trigonal bipyramidal arrangement to an axial position. Give reason.

Ans. In SF₄ molecule, the lp electrons occupies an equatorial position because, lp - bp repulsion is minimum.

23.How is VBT different from Lewis concept?

Ans.In Lewis concept, bond formation is explained in terms of sharing of electron pairs and the Octet rule whereas in VBT bond formation is described in terms of hybridization and overlap of the orbitals.

24.S – orbital does not show any preference for direction. Why?

Ans. S – Orbital does not show any preference for direction because it is spherically symmetrical.

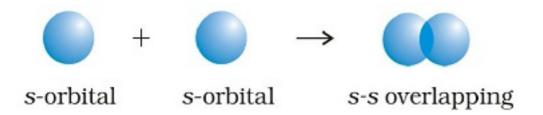
25.Why is s– bond stronger than π - bond?

Ans. Orbitals can overlap to a greater extent in a s - bond due to axial orientation, so s - bond is strong. Whereas, in a pi – bond sideways overlapping is not to an appreciable extent due to the presence of s - bond which restricts the distance between the involved atoms.

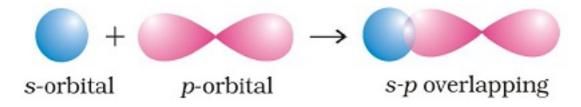
26.What are the different types of s - bond formation?

Ans. s - bond can be formed by any of the following types of combinations of atoms orbitals.

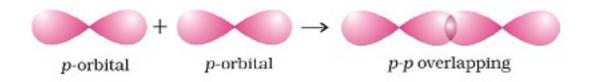
(a) S – S – overlapping : In this case, there is a overlap of two half – filled S – orbitals along the inter nuclear axis.



(b) S- P overlapping : This type of over lapping occurs between half – filled s-orbitals of one atom and half-filled p-orbitals of another atom.

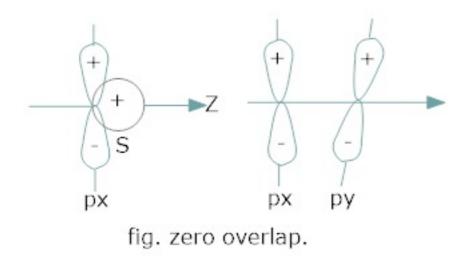


(c) P – P overlapping : This type of overlap takes place between half-filled p-orbitals of the two approaching atoms.



27.What is zero over lap?

Ans. The unsymmetrical overlap of orbitals results in zero overlap i-e; between px-s and px-py orbital



28. the features of hybridisation.

Ans. The main features of hybridization are

(i) The number of hybrid orbitals is equal to number of the atomic orbitals that get hybridized.

(ii) The hybridized orbitals are always equivalent in energy and shape.

(iii) The hybrid orbitals are more effective in forming stable bonds than the pure atomic orbitals.

(iv) The hybrid orbitals orient in a manner to minimize repulsion resulting in a particular geometrical shape.

29.What are the important consolations for hybridisation?

Ans. (i) The orbitals present in the valence shell of the atom are hybridised.

(ii) The orbitals undergoing hybridization should have almost the same energy.

(iii)It is not essential that electrons get promoted prior to hybridization.

It is necessary that only half filled orbitals participate in hybridisation even filled orbitlals can take part.

30.Describe the shape of sp, sp^2 and sp^3 hybrid orbital?

Ans. (i) Sp-hybrid orbital is oriented to an angle 180° .

(ii) S_p^2 -hybrid orbital lie in a plane and is directed towards the corners of equilateral triangle making an angle of 120° .

(iii) Sp^3 -hybrid orbitals are directed towards the four corners of tetrahedron making an angle of $109^{\circ} 28'$

31.Ethylene is a planar molecule whereas acetylene is a linear molecule. Give reason.

Ans. In case of ethylene, C_2H_4 , show Sp^2 hybridization where the four hydrogen atoms

are placed in four corners of a plane sharing 120°

Whereas acetylene shows sp hybridization and shares an angle of 180° and thus it is linear.



32.In H_2O , H_2S , H_2Se , H_2Te , the bond angle decreases though all have the same bent shape. Why?

Ans. In all the four cases, the molecules undergo Sp^3 hybridization forming four hybrid orbitals, two of which are occupied by lp of electrons and two by bp electrons. Thus they are expected to have $109^{0}28^{\circ}$ angle but this does not happen. In case of H_2O molecule, as oxygen is small in size and has high electronegativity value, the bp are closer due to which it is subjected to larger repulsion (bp-bp). In case of H_2S as S atom is larger than O, bp-bp repulsion is less as compared to H_2O and it is true for H_2Se and H_2Te as well.

33.Out of p-orbital and sp-hybrid orbital which has greater directional character and Why?

Ans. Sp-hybrid orbital has greater directional character than p-orbital. Because in case of porbitals, the two lobes are equal in size and equal electron density is distributed whereas in Sp-hybrid orbital, electron density is greater on one side.

34. He_2 does not exist. Explain in terms of LCAO.

Ans. The electronic configuration of helium atom is $1s^2$. Each helium atom contains 2 electrons, therefore, in He_2 molecule there would be 4 electrons. These electrons will accommodated in σ_{1s} and σ^+_{1s} molecular orbitals leading to electronic configuration :

$$He_2 : (\sigma 1s)^2 (\sigma^+ 1s)^2$$

Bond order of He_2 is $\frac{1}{2}(2-2) = 0$

 He_2 molecule is there unstable and does not exist.

35.Dipole moment is a scalar or a vector quantity?

Ans.Dipole moment is a vector quantity and is depicted by a small arrow with tail on the +ve centre and head pointing towards the negative centre.