

Unit

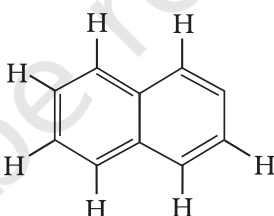
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CHEMICAL BONDING AND MOLECULAR STRUCTURE

I. Multiple Choice Questions (Type-I)

- Isostructural species are those which have the same shape and hybridisation. Among the given species identify the isostructural pairs.
 - $[\text{NF}_3 \text{ and } \text{BF}_3]$
 - $[\text{BF}_4^- \text{ and } \text{NH}_4^+]$
 - $[\text{BCl}_3 \text{ and } \text{BrCl}_3]$
 - $[\text{NH}_3 \text{ and } \text{NO}_3^-]$
- Polarity in a molecule and hence the dipole moment depends primarily on electronegativity of the constituent atoms and shape of a molecule. Which of the following has the highest dipole moment?
 - CO_2
 - HI
 - H_2O
 - SO_2
- The types of hybrid orbitals of nitrogen in NO_2^+ , NO_3^- and NH_4^+ respectively are expected to be
 - sp , sp^3 and sp^2
 - sp , sp^2 and sp^3
 - sp^2 , sp and sp^3
 - sp^2 , sp^3 and sp
- Hydrogen bonds are formed in many compounds e.g., H_2O , HF , NH_3 . The boiling point of such compounds depends to a large extent on the strength of hydrogen bond and the number of hydrogen bonds. The correct decreasing order of the boiling points of above compounds is :
 - $\text{HF} > \text{H}_2\text{O} > \text{NH}_3$

- (ii) $\text{H}_2\text{O} > \text{HF} > \text{NH}_3$
 (iii) $\text{NH}_3 > \text{HF} > \text{H}_2\text{O}$
 (iv) $\text{NH}_3 > \text{H}_2\text{O} > \text{HF}$
5. In PO_4^{3-} ion the formal charge on the oxygen atom of P–O bond is
 (i) + 1
 (ii) – 1
 (iii) – 0.75
 (iv) + 0.75
6. In NO_3^- ion, the number of bond pairs and lone pairs of electrons on nitrogen atom are
 (i) 2, 2
 (ii) 3, 1
 (iii) 1, 3
 (iv) 4, 0
7. Which of the following species has tetrahedral geometry?
 (i) BH_4^-
 (ii) NH_2^-
 (iii) CO_3^{2-}
 (iv) H_3O^+
8. Number of π bonds and σ bonds in the following structure is–



- (i) 6, 19
 (ii) 4, 20
 (iii) 5, 19
 (iv) 5, 20
9. Which molecule/ion out of the following does **not** contain unpaired electrons?
 (i) N_2^+
 (ii) O_2
 (iii) O_2^{2-}
 (iv) B_2

- 10.** In which of the following molecule/ion all the bonds are **not** equal?
- XeF_4
 - BF_4^-
 - C_2H_4
 - SiF_4
- 11.** In which of the following substances will hydrogen bond be strongest?
- HCl
 - H_2O
 - HI
 - H_2S
- 12.** If the electronic configuration of an element is $1s^2 2s^2 2p^6 3s^2 3p^6 3d^2 4s^2$, the four electrons involved in chemical bond formation will be ____.
- $3p^6$
 - $3p^6, 4s^2$
 - $3p^6, 3d^2$
 - $3d^2, 4s^2$
- 13.** Which of the following angle corresponds to sp^2 hybridisation?
- 90°
 - 120°
 - 180°
 - 109°

The electronic configurations of three elements, A, B and C are given below. Answer the questions 14 to 17 on the basis of these configurations.

| | | | | | |
|---|--------|--------|--------|--------|--------|
| A | $1s^2$ | $2s^2$ | $2p^6$ | | |
| B | $1s^2$ | $2s^2$ | $2p^6$ | $3s^2$ | $3p^3$ |
| C | $1s^2$ | $2s^2$ | $2p^6$ | $3s^2$ | $3p^5$ |

- 14.** Stable form of **A** may be represented by the formula :
- A
 - A_2
 - A_3
 - A_4
- 15.** Stable form of **C** may be represented by the formula :
- C
 - C_2

- (iii) C_3
 (iv) C_4
16. The molecular formula of the compound formed from B and C will be
 (i) BC
 (ii) B_2C
 (iii) BC_2
 (iv) BC_3
17. The bond between B and C will be
 (i) Ionic
 (ii) Covalent
 (iii) Hydrogen
 (iv) Coordinate
18. Which of the following order of energies of molecular orbitals of N_2 is correct?
 (i) $(\pi 2p_y) < (\sigma 2p_z) < (\pi^* 2p_x) \approx (\pi^* 2p_y)$
 (ii) $(\pi 2p_y) > (\sigma 2p_z) > (\pi^* 2p_x) \approx (\pi^* 2p_y)$
 (iii) $(\pi 2p_y) < (\sigma 2p_z) > (\pi^* 2p_x) \approx (\pi^* 2p_y)$
 (iv) $(\pi 2p_y) > (\sigma 2p_z) < (\pi^* 2p_x) \approx (\pi^* 2p_y)$
19. Which of the following statement is **not** correct from the view point of molecular orbital theory?
 (i) Be_2 is not a stable molecule.
 (ii) He_2 is not stable but He_2^+ is expected to exist.
 (iii) Bond strength of N_2 is maximum amongst the homonuclear diatomic molecules belonging to the second period.
 (iv) The order of energies of molecular orbitals in N_2 molecule is
 $\sigma 2s < \sigma^* 2s < \sigma 2p_z < (\pi 2p_x = \pi 2p_y) < (\pi^* 2p_x = \pi^* 2p_y) < \sigma^* 2p_z$
20. Which of the following options represents the correct bond order :
 (i) $O_2^- > O_2 > O_2^+$
 (ii) $O_2^- < O_2 < O_2^+$
 (iii) $O_2^- > O_2 < O_2^+$
 (iv) $O_2^- < O_2 > O_2^+$
21. The electronic configuration of the outer most shell of the most electronegative element is
 (i) $2s^2 2p^5$
 (ii) $3s^2 3p^5$
 (iii) $4s^2 4p^5$
 (iv) $5s^2 5p^5$

22. Amongst the following elements whose electronic configurations are given below, the one having the highest ionisation enthalpy is
- (i) $[\text{Ne}]3s^23p^1$
 - (ii) $[\text{Ne}]3s^23p^3$
 - (iii) $[\text{Ne}]3s^23p^2$
 - (iv) $[\text{Ar}]3d^{10}4s^24p^3$

II. Multiple Choice Questions (Type-II)

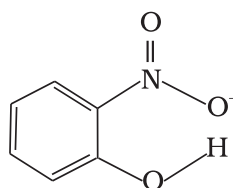
In the following questions two or more options may be correct.

23. Which of the following have identical bond order?
- (i) CN^-
 - (ii) NO^+
 - (iii) O_2^-
 - (iv) O_2^{2-}
24. Which of the following attain the linear structure:
- (i) BeCl_2
 - (ii) NCO^+
 - (iii) NO_2
 - (iv) CS_2
25. CO is isoelectronic with
- (i) NO^+
 - (ii) N_2
 - (iii) SnCl_2
 - (iv) NO_2^-
26. Which of the following species have the same shape?
- (i) CO_2
 - (ii) CCl_4
 - (iii) O_3
 - (iv) NO_2^-
27. Which of the following statements are correct about CO_3^{2-} ?
- (i) The hybridisation of central atom is sp^3 .
 - (ii) Its resonance structure has one C–O single bond and two C=O double bonds.
 - (iii) The average formal charge on each oxygen atom is 0.67 units.
 - (iv) All C–O bond lengths are equal.

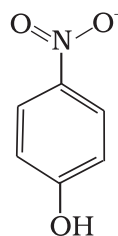
- 28.** Dimagnetic species are those which contain no unpaired electrons. Which among the following are dimagnetic?
- N_2
 - N_2^{2-}
 - O_2
 - O_2^{2-}
- 29.** Species having same bond order are :
- N_2
 - N_2^-
 - F_2^+
 - O_2^-
- 30.** Which of the following statements are **not** correct?
- NaCl being an ionic compound is a good conductor of electricity in the solid state.
 - In canonical structures there is a difference in the arrangement of atoms.
 - Hybrid orbitals form stronger bonds than pure orbitals.
 - VSEPR Theory can explain the square planar geometry of XeF_4 .

III. Short Answer Type

- 31.** Explain the non linear shape of H_2S and non planar shape of PCl_3 using valence shell electron pair repulsion theory.
- 32.** Using molecular orbital theory, compare the bond energy and magnetic character of O_2^+ and O_2^- species.
- 33.** Explain the shape of BrF_5 .
- 34.** Structures of molecules of two compounds are given below :



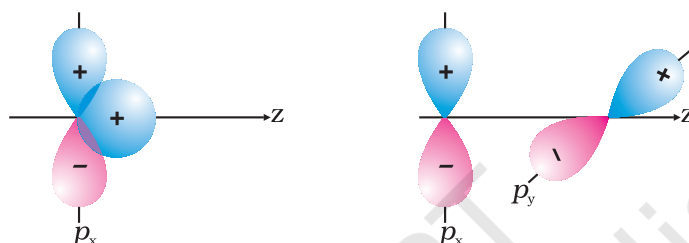
(I)



(II)

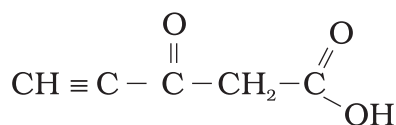
- (a) Which of the two compounds will have intermolecular hydrogen bonding and which compound is expected to show intramolecular hydrogen bonding.
- (b) The melting point of a compound depends on, among other things, the extent of hydrogen bonding. On this basis explain which of the above two compounds 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.

35. Why does type of overlap given in the following figure not result in bond formation?



- 36.** Explain why PCl_5 is trigonal bipyramidal whereas IF_5 is square pyramidal.
- 37.** In both water and dimethyl ether ($\text{CH}_3 - \ddot{\text{O}} - \text{CH}_3$), oxygen atom is central atom, and has the same hybridisation, yet they have different bond angles. Which one has greater bond angle? Give reason.
- 38.** Write Lewis structure of the following compounds and show formal charge on each atom.
 HNO_3 , NO_2 , H_2SO_4
- 39.** 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^+ , N_2^- , N_2^{2+}
- 40.** What is the effect of the following processes on the bond order in N_2 and O_2 ?
 (i) $\text{N}_2 \rightarrow \text{N}_2^+ + e^-$ (ii) $\text{O}_2 \rightarrow \text{O}_2^+ + e^-$
- 41.** Give reasons for the following :
 (i) Covalent bonds are directional bonds while ionic bonds are non-directional.
 (ii) Water molecule has bent structure whereas carbon dioxide molecule is linear.
 (iii) Ethyne molecule is linear.

42. What is an ionic bond? With two suitable examples explain the difference between an ionic and a covalent bond?
43. Arrange the following bonds in order of increasing ionic character giving reason.
N—H, F—H, C—H and O—H
44. Explain why CO_3^{2-} ion cannot be represented by a single Lewis structure. How can it be best represented?
45. 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.



46. Group the following as linear and non-linear molecules :
 H_2O , HOCl , BeCl_2 , Cl_2O
47. 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?
48. Draw the resonating structure of
(i) Ozone molecule
(ii) Nitrate ion
49. Predict the shapes of the following molecules on the basis of hybridisation.
 BCl_3 , CH_4 , CO_2 , NH_3
50. All the C—O bonds in carbonate ion (CO_3^{2-}) are equal in length. Explain.
51. What is meant by the term average bond enthalpy? Why is there difference in bond enthalpy of O—H bond in ethanol ($\text{C}_2\text{H}_5\text{OH}$) and water?

IV. Matching Type

52. 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^3d |
| (iv) NH_4^+ | (d) sp^3 |
| | (e) sp |

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

| Column I | Column II |
|-----------------------------------|--------------------------|
| (i) H_3O^+ | (a) Linear |
| (ii) $\text{HC} \equiv \text{CH}$ | (b) Angular |
| (iii) ClO_2^- | (c) Tetrahedral |
| (iv) NH_4^+ | (d) Trigonal bipyramidal |
| | (e) Pyramidal |

54. 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 |

55. 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 |
| (iii) Ionic solid | (c) H_2 |
| (iv) Covalent solid | (d) HF |
| | (e) O_3 |

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

| Column I | Column II |
|-----------------|------------|
| (i) Tetrahedral | (a) sp^2 |
| (ii) Trigonal | (b) sp |
| (iii) Linear | (c) sp^3 |

V. Assertion and Reason Type

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.

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

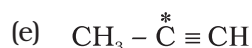
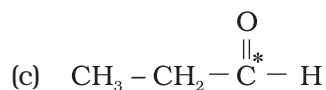
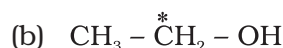
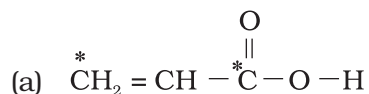
Reason (R) : This is because sodium and chloride ions acquire octet in sodium chloride formation.

- (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.
- 58. Assertion (A) :** Though the central atom of both NH_3 and H_2O molecules are sp^3 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.
- 59. Assertion (A):** Among the two O–H bonds in H_2O 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.

VI. Long Answer Type

- 60.** (i) Discuss the significance/ applications of dipole moment.
 (ii) Represent diagrammatically the bond moments and the resultant dipole moment in CO_2 , NF_3 and CHCl_3 .
- 61.** Use the molecular orbital energy level diagram to show that N_2 would be expected to have a triple bond, F_2 , a single bond and Ne_2 , no bond.
- 62.** 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?
- 63.** Describe hybridisation in the case of PCl_5 and SF_6 . The axial bonds are longer as compared to equatorial bonds in PCl_5 whereas in SF_6 both axial bonds and equatorial bonds have the same bond length. Explain.
- 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.



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

$\sigma 1s < \sigma^* 1s < \sigma 2s < \sigma^* 2s < \sigma 2p_z < (\pi 2p_x \approx \pi 2p_y) < (\pi^* 2p_x \approx \pi^* 2p_y) < \sigma^* 2p_z$

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.

65. Which of the following statements is correct?

- In the formation of dioxygen from oxygen atoms 10 molecular orbitals will be formed.
- All the molecular orbitals in the dioxygen will be completely filled.
- Total number of bonding molecular orbitals will not be same as total number of anti bonding orbitals in dioxygen.
- Number of filled bonding orbitals will be same as number of filled anti bonding orbitals.

66. Which of the following molecular orbitals has maximum number of nodal planes?

- (i) $\sigma^* 1s$
- (ii) $\sigma^* 2p_z$
- (iii) $\pi 2p_x$
- (iv) $\pi^* 2p_y$

67. Which of the following pair is expected to have the same bond order?

- (i) O_2, N_2
- (ii) O_2^+, N_2^-
- (iii) O_2^-, N_2^+
- (iv) O_2^-, N_2^-

68. In which of the following molecules, $\sigma 2p_z$ molecular orbital is filled after $\pi 2p_x$ and $\pi 2p_y$ molecular orbitals?

- (i) O_2
- (ii) Ne_2
- (iii) N_2
- (iv) F_2

ANSWERS

I. Multiple Choice Questions (Type-I)

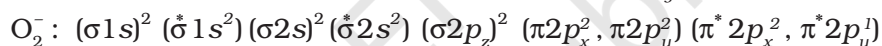
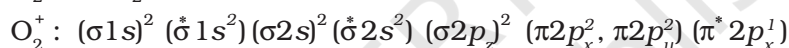
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|----------|----------|----------|-----------|----------|----------|
| 1. (ii) | 2. (iii) | 3. (ii) | 4. (ii) | 5. (ii) | 6. (iv) |
| 7. (i) | 8. (iii) | 9. (iii) | 10. (iii) | 11. (ii) | 12. (iv) |
| 13. (ii) | 14. (i) | 15. (ii) | 16. (iv) | 17. (ii) | 18. (i) |
| 19. (iv) | 20. (ii) | 21. (i) | 22. (ii) | | |

II. Multiple Choice Questions (Type-II)

- | | | |
|-----------------|-----------------|---------------|
| 23. (i), (ii) | 24. (i), (iv) | 25. (i), (ii) |
| 26. (iii), (iv) | 27. (iii), (iv) | 28. (i), (iv) |
| 29. (iii), (iv) | 30. (i), (ii) | |

III. Short Answer Type

32. (i) According to molecular orbital theory electronic configurations of O_2^+ and O_2^- species are as follows :

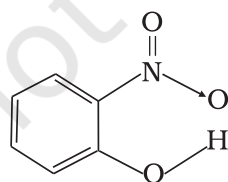


$$\text{Bond order of } O_2^+ = \frac{10 - 5}{2} = \frac{5}{2} = 2.5$$

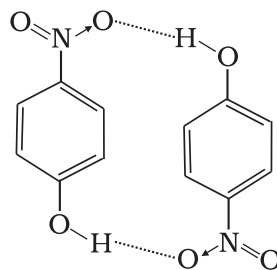
$$\text{Bond order of } O_2^- = \frac{10 - 7}{2} = \frac{3}{2} = 1.5$$

Higher bond order of O_2^+ shows that it is more stable than O_2^- . Both the species have unpaired electrons. So both are paramagnetic in nature.

34. (a) Compound (I) will form intramolecular hydrogen bond because NO_2 and OH group are close together in comparison to that in compound (II).



(I)



(II)

- (b) Compound (II) will have higher melting point because it forms intermolecular hydrogen bonds. Thus, more and more molecules are joined together through hydrogen bond formation.
- (c) Due to intramolecular hydrogen bonding compound (I) will not be able to form hydrogen bonds with water thus will be less soluble in it while compound (II) can form hydrogen bond with water more easily and will be soluble in water.
37. **[Hint :** Dimethyl ether will have larger bond angle. There will be more repulsion between bond pairs of CH_3 groups attached in ether than between bond pairs of hydrogen atoms attached to oxygen in water. The carbon of CH_3 in ether is attached to three hydrogen atoms through σ bonds and electron pairs of these bonds add to the electronic charge density on carbon atom. Hence, repulsion between two $-\text{CH}_3$ groups will be more than that between two hydrogen atoms.]

IV. Matching Type

52. (i) \rightarrow (c) (ii) \rightarrow (a) (iii) \rightarrow (e) (iv) \rightarrow (d)
53. (i) \rightarrow (e) (ii) \rightarrow (a) (iii) \rightarrow (b) (iv) \rightarrow (c)
54. (i) \rightarrow (c) (ii) \rightarrow (d) (iii) \rightarrow (a) (iv) \rightarrow (b)
55. (i) \rightarrow (d) (ii) \rightarrow (e) (iii) \rightarrow (b) (iv) \rightarrow (a)
56. (i) \rightarrow (c) (ii) \rightarrow (a) (iii) \rightarrow (b)

V. Assertion and Reason Type

57. (i) 58. (i) 59. (iv)
65. (i) 66. (ii) 67. (ii) 68. (iii)