

## CHEMICAL BONDING - IV

# Exercise # 1

## PART - I : SUBJECTIVE QUESTIONS

### Section (A) : Dipole moment

- A-1. Inorganic benzene is more reactive than organic benzene. Why?
- A-2. trans-1, 2 dichloro ethene have zero dipole moment while its cis- form has some dipole moment. Explain.
- A-3. Why  $\text{CCl}_4$  have zero dipole moment but  $\text{CHCl}_3$  have some dipole moment ?
- A-4. Arrange in increasing order of dipole moment ;  $\text{H}_2\text{O}$ ,  $\text{H}_2\text{S}$ ,  $\text{BF}_3$ .
- A-5. Dipole moment of LiF was experimentally determined and was found to be 6.32 D. Calculate percentage ionic character in LiF molecule Li—F bond length is 156 pm.

### Section (B) : Intermolecular forces of attraction and H-bonding

- B-1. In which of the following the hydrogen bonding is strongest. Explain briefly ?  
 (A)  $\text{O}-\text{H} \cdots \text{S} (\ell)$  (B)  $\text{S}-\text{H} \cdots \text{O} (\ell)$  (C)  $\text{F}-\text{H} \cdots \text{F} (\text{s})$  (D)  $\text{F}-\text{H} \cdots \text{O} (\ell)$
- B-2. Why  $\text{D}_2\text{O}$  has higher viscosity than  $\text{H}_2\text{O}$  ?

### Section (C) : Applications of H-bond

- C-1. Why glucose, fructose, sucrose etc. are soluble in water though they are covalent compounds ?
- C-2. Ethanol has higher boiling point than diethyl ether. Why ?

## PART - II : OBJECTIVE QUESTIONS

\* Marked Questions may have more than one correct options.

### Section (A) : Dipole moment

- A-1. Which has maximum dipole moment ?



- A-2. Of the following molecules, the one, which has permanent dipole moment, is :  
 (A)  $\text{SiF}_4$  (B)  $\text{BF}_3$  (C)  $\text{PF}_3$  (D)  $\text{PF}_5$
- A-3. Which of the following has the least dipole moment ?  
 (A)  $\text{NF}_3$  (B)  $\text{CO}_2$  (C)  $\text{SO}_2$  (D)  $\text{NH}_3$
- A-4. Which of the following compounds possesses zero dipole moment?  
 (A) Benzene ( $\text{C}_6\text{H}_6$ ) (B) Carbon tetrachloride  
 (C) Boron trifluoride (D) All of these
- A-5. Which of the following statements is false for  $\text{XeO}_3\text{F}_2$  ?  
 (A) Hybridisation of central atom xenon is  $\text{sp}^3\text{d}$ . (B) The compound is non-polar.  
 (C) The compound has  $\text{p}\pi\text{-d}\pi$  bonding. (D) None.
- A-6. The geometry of  $\text{H}_2\text{S}$  and its dipole moment are : [JEE-1999, 2/80]  
 (A) angular and non zero (B) angular and zero  
 (C) linear and non zero (D) linear and zero

- A-7.** Which of the following has been arranged in order of decreasing dipole moment ?  
 (A)  $\text{CH}_3\text{Cl} > \text{CH}_3\text{F} > \text{CH}_3\text{Br} > \text{CH}_3\text{I}$  (B)  $\text{CH}_3\text{F} > \text{CH}_3\text{Cl} > \text{CH}_3\text{Br} > \text{CH}_3\text{I}$   
 (C)  $\text{CH}_3\text{Cl} > \text{CH}_3\text{Br} > \text{CH}_3\text{I} > \text{CH}_3\text{F}$  (D)  $\text{CH}_3\text{F} > \text{CH}_3\text{Cl} > \text{CH}_3\text{I} > \text{CH}_3\text{Br}$
- A-8.** The gaseous HX molecule has a measured dipole moment of 4.0 D, which indicates that it is a very polar molecule. The separation between the nuclei in this molecule is  $2.67 \times 10^{-8}$  cm then the percentage ionic character in HX molecule is :  
 (A) 78% (B) 31.25% (C) 50.25% (D) None of these
- A-9.** The dipole moment of chlorobenzene is 1.73 D. The dipole moment of p-dichlorobenzene is expected to be :  
 (A) 3.46 D (B) 0.00 D (C) 1.73 D (D) 1.00 D

### Section (B) : Intermolecular forces of attraction and H-bonding

- B-1.** Which of the following is not correctly matched with respect to the intermolecular forces existing amongst the molecules (Hydrogen bonding is not taken as dipole-dipole attraction) ?  
 (A) Benzene – London dispersion forces  
 (B) Orthophosphoric acid – London dispersion force, hydrogen bonding.  
 (C) Hydrochloric acid – London dispersion force, dipole-dipole attraction  
 (D) Iodine monochloride – London dispersion force
- B-2.** Which of the following models best describes the bonding within a layer of the graphite structure ?  
 (A) metallic bonding (B) ionic bonding  
 (C) non-metallic covalent bonding (D) van der Waals forces
- B-3.** Which of the following factor is responsible for van der Waals forces ?  
 (A) Instantaneous dipole-induced dipole interaction.  
 (B) Dipole-induced dipole interaction and ion-induced dipole interaction.  
 (C) Dipole-dipole interaction and ion-induced dipole interaction.  
 (D) All of these.
- B-4.** Which of the following bonds/forces is weakest ?  
 (A) Covalent bond (B) Ionic bond (C) Hydrogen bond (D) London force
- B-5.** In which of the following compound, intra-molecular H-bonding is not observed :  
 (A) O-hydroxy benzaldehyde (B) O-nitrophenol  
 (C) Chloral hydrate (D) Boric acid
- B-6.** Consider the following sets of H-bonds
- P :  $\begin{array}{c} \text{---O---H---} \text{---N---} \\ | \\ \text{---N---H---} \text{---N---} \\ | \\ \text{---} \end{array}$

R :  $\begin{array}{c} | \\ \text{---N---} \text{---H---} \text{---N---} \\ | \\ \text{---} \end{array}$

Q :  $\begin{array}{c} \text{---O---H---} \text{---O---} \\ | \\ \text{---} \end{array}$

S :  $\begin{array}{c} | \\ \text{---N---} \text{---H---} \text{---O---} \\ | \\ \text{---} \end{array}$
- The correct order of H-bond strengths is :  
 (A)  $Q > P > S > R$  (B)  $R > Q > S > P$   
 (C)  $R > S > P > Q$  (D)  $P > Q > R > S$
- B-7.** Which of the following compounds would have significant intermolecular hydrogen bonding ?  
 $\text{HF}, \text{CH}_3\text{OH}, \text{N}_2\text{O}_4, \text{CH}_4$   
 (A)  $\text{HF}, \text{N}_2\text{O}_4$  (B)  $\text{HF}, \text{CH}_4, \text{CH}_3\text{OH}$   
 (C)  $\text{HF}, \text{CH}_3\text{OH}$  (D)  $\text{CH}_3\text{OH}, \text{CH}_4$

### Section (C) : Applications of H-bond

- C-1.** Amongst  $\text{NH}_3$ ,  $\text{PH}_3$ ,  $\text{AsH}_3$  and  $\text{SbH}_3$  the one with highest boiling point is :  
 (A)  $\text{NH}_3$  because of lower molecular weight (B)  $\text{SbH}_3$  because of higher molecular weight  
 (C)  $\text{PH}_3$  because of H-bonding (D)  $\text{AsH}_3$  because of lower molecular weight
- C-2.** Which of the following statement is not true ?  
 (A)  $\text{CCl}_4$  has higher boiling point than  $\text{CHCl}_3$ .  
 (B) The  $\text{HF}_2^-$  ion exists in the solid state and in liquid HF solution, but not in dilute aqueous solutions.  
 (C) Hydrogen bonding maintains the planar  $\text{H}_3\text{BO}_3$  units in layers in solid state.  
 (D) None of these.
- C-3.**  $\text{S}_1$  : In the solid  $\text{B}(\text{OH})_3$  units are hydrogen bonded together into two-dimensional sheets with almost hexagonal symmetry.  
 $\text{S}_2$  :  $\text{Na}_2\text{CO}_3$  can be isomorphous with  $\text{Na}_2\text{SO}_3$  as both have similar formula type.  
 $\text{S}_3$  :  $\text{XeO}_3\text{F}_2$  has one lone pair of electron on central xenon atom.  
 $\text{S}_4$  :  $\text{D}_2\text{O}$  has higher boiling point than  $\text{H}_2\text{O}$   
 (A) T F T F (B) T F F T (C) T T F F (D) T T T T
- C-4.** The correct order of boiling point is :  
 (A)  $\text{H}_2\text{O} < \text{H}_2\text{S} < \text{H}_2\text{Se} < \text{H}_2\text{Te}$  (B)  $\text{H}_2\text{O} > \text{H}_2\text{Se} > \text{H}_2\text{Te} > \text{H}_2\text{S}$   
 (C)  $\text{H}_2\text{O} > \text{H}_2\text{S} > \text{H}_2\text{Se} > \text{H}_2\text{Te}$  (D)  $\text{H}_2\text{O} > \text{H}_2\text{Te} > \text{H}_2\text{Se} > \text{H}_2\text{S}$
- C-5.** Which of the following compounds has the highest boiling point  
 (A)  $\text{HCl}$  (B)  $\text{HBr}$  (C)  $\text{H}_2\text{SO}_4$  (D)  $\text{HNO}_3$
- C-6.** Which is correct about  $\text{D}_2\text{O}$   
 (A) Its boiling point is higher than that of  $\text{H}_2\text{O}$  (l)  
 (B)  $\text{O}-\text{D}-\cdots-\text{O}$  bond is stronger than  $\text{O}-\text{H}-\cdots-\text{O}$  bond.  
 (C)  $\text{D}_2\text{O}(\text{s})$  sinks in  $\text{H}_2\text{O}$  (l).  
 (D) all the above are correct.

### PART - III : ASSERTION / REASONING

Each question has 5 choices (A), (B), (C), (D) and (E) out of which ONLY ONE is correct.

- (A) STATEMENT-1 is true, STATEMENT-2 is true and STATEMENT-2 is correct explanation for STATEMENT-1  
 (B) STATEMENT-1 is true, STATEMENT-2 is true and STATEMENT-2 is not correct explanation for STATEMENT-1  
 (C) STATEMENT-1 is true, STATEMENT-2 is false  
 (D) STATEMENT-1 is false, STATEMENT-2 is true  
 (E) Both STATEMENTS are false

- Statement-1** :  $\text{NF}_3$  has little tendency to act as a donor molecule.  
**Statement-2** : The highly electronegative F atoms attract electrons and these moments partly cancel the moment from the lone pair.
- Statement-1** : Fluoromethane ( $\text{CH}_3\text{F}$  ;  $\mu = 1.85 \text{ D}$ ) has a smaller dipole moment than chloromethane ( $\text{CH}_3\text{Cl}$  ;  $\mu = 1.87 \text{ D}$ )  
**Statement-2** : Fluorine has less electron affinity than that of chlorine.
- Statement-1** : Acetylene is not soluble in  $\text{H}_2\text{O}$  but is highly soluble in acetone.  
**Statement-2** : Acetylene forms inter molecular H-bond with acetone easily but not with  $\text{H}_2\text{O}$  as water molecular themselves are highly associated through inter molecular H-bonds.
- Statement-1** : Crystals of hydrated calcium sulphate (gypsum :  $(\text{CaSO}_4 \cdot 2 \text{H}_2\text{O})$ ) are soft and easily cleaved.  
**Statement-2** : Crystals of anhydrous calcium sulphate (anhydrite :  $\text{CaSO}_4$ ) are very hard and very difficult to cleave.
- Statement-1** : In case of persulphuric acid, the  $K_1 \gg K_2$ .  
**Statement-2** : The anion of persulphuric acid is intermolecular hydrogen bonded.

6. **Statement-1** : Ortho boric acid crystal are hard and cannot be broken easily into the powder form.  
**Statement-2** : In the solid state  $B(OH)_3$  units are hydrogen bonded together into two dimensional sheets.
7. **Statement-1** : The crystal structures of  $NaHCO_3$  and  $KHCO_3$ , both show intermolecular hydrogen bonding but are different.  
**Statement-2** : In  $NaHCO_3$  the  $HCO_3^-$  ions are linked together through intermolecular hydrogen bonds into an infinite chain, while in  $KHCO_3$ ,  $HCO_3^-$  ions form dimeric anions through intermolecular hydrogen bonds.
8. **Statement-1** : Fluorine ( $F_2$ ) is gas while iodine ( $I_2$ ) is solid at room temperature.  
**Statement-2** : A larger molecule or heavy atom is more polarizable and has larger dispersion forces because it has many electrons some of which are less tightly held and are farther from the nucleus.
9. **Statement-1** : Noble gases are liquefied at very low temperature. Hence they have low boiling points.  
**Statement-2** : Noble gases being monoatomic have no other interatomic forces except weak dispersion forces.

## Exercise # 2

### PART - I : SUBJECTIVE QUESTIONS

1. Why  $CH_3 - O - H$  is having more dipole moment than  $CH_3 - S - H$ ?
2. Out of trimethylamine and trimethylphosphine, which one has higher dipole moment?
3. Benzene has zero dipole moment and therefore, experiences no dipole-dipole forces but it is a liquid rather than a gas at room temperature. Explain.
4.  $Br_2$  and  $ICl$  have the same number of electrons but  $ICl$  has higher boiling point than  $Br_2$ . Why, explain?
5. Explain why o-hydroxybenzaldehyde is a liquid at room temperature while p-hydroxybenzaldehyde is a high melting solid.
6. If bond length of  $HCl$  is  $1.2 \text{ \AA}$  and dipole moment is  $(\mu) = 1.44 \text{ D}$ , then find the % covalent character in  $HCl$  ( $q_e = 4.8 \times 10^{-10} \text{ esu}$ )
7. Why  $CH_3Cl$  is having higher dipole moment than  $CH_3F$ ?

### PART - II : OBJECTIVE QUESTIONS

#### Single choice type

1. **S<sub>1</sub>** : In  $CrO_5$ , the oxidation number of Cr is +6.  
**S<sub>2</sub>** : Out of  $CH_3Cl$  and  $CHCl_3$ ,  $CH_3Cl$  has higher dipole moment  
**S<sub>3</sub>** : Hybridisation of sulphur in  $SO_3$  and in its trimer is the same,  $sp^2$ .  
(A) T F T (B) T T F (C) T F F (D) T T T
2. Among the  $XeF_2$ ,  $SF_2Cl_2$ ,  $XeOF_2$ ,  $ICl_2^-$ ,  $IOCl_4^-$  and  $F_2ClO^+$   
**S<sub>1</sub>** :  $XeF_2$ ,  $ICl_2^-$ ,  $XeOF_2$  have zero dipole moment  
**S<sub>2</sub>** :  $IOCl_4^-$  and  $F_2ClO^+$  have different electronic arrangement (geometry) at central atom  
**S<sub>3</sub>** :  $SF_2Cl_2$ ,  $IOCl_4^-$  and  $F_2ClO^+$  have equal number of lone pairs of electrons at the central atom.  
**S<sub>4</sub>** : All bond angle in each of species,  $XeOF_4$ ,  $IOCl_4^-$ ,  $SF_2Cl_2$  and  $F_2ClO^+$  are identical  
The correct order for the above statements is :  
(A) F T T F (B) F F F F (C) T T F F (D) T F T F
3. Which of the following would be expected to have a dipole moment of zero on the basis of symmetry?  
(A)  $SOCl_2$  (B)  $OF_2$  (C)  $SeF_6$  (D)  $ClF_5$
4. If molecule  $MX_3$  has Zero dipole moment, the hybrid orbitals used by M (Atomic No. < 21) are  
(A) Pure p (B) sp hybrid (C)  $sp^2$  hybrid (D)  $sp^3$  hybrid

5. Given the species  $N_2$ ,  $CO$ ,  $CN^-$  and  $NO^+$ . Which of the following statement is incorrect.
 

(A) All the species are diamagnetic (B) All the species are isoelectronic  
(C) All the species have dipole moment (D) All the species are linear
6. Which of the following are incorrect for dipole moment ?
 

(A) Lone pair of elements present on central atom can give rise to dipole moment  
(B) Dipole moment is vector quantity  
(C)  $PF_5$  (g) molecule has non zero dipole moment  
(D) Difference in electronegativities of combining atom can lead to dipole moment
7. Which of the following orders are correct regarding mentioned properties
 

(A)  $SO_3 < CCl_4 < XeF_2$  (Bond angle).  
(B)  $SOF_2 > SOCl_2 > SOBr_2$  (Bond angle)  
(C)  $CH_3COO^- > CO_3^{2-} > CH_3COCH_3$  (C — O bond length)  
(D)  $CH_3Cl > CH_3F < CD_3F$  (dipole moment).
8.  $CH_3Cl$  has more dipole moment than  $CH_3F$  because :
 

(A) electron affinity of chlorine is greater than that of fluorine.  
(B) the charge separation is larger in  $CH_3Cl$  compared to  $CH_3F$ .  
(C) the repulsion between the bond pairs and non-bonded pairs of electrons is greater in  $CH_3Cl$  than  $CH_3F$ .  
(D) chlorine has higher electronegativity than fluorine.
9.  $S_1$  : In ozone molecule, O — O bond lengths are equal, this can be explained on the basis of resonance.  
 $S_2$  : Ion-dipole attraction is responsible for hydration of ions.  
 $S_3$  : Intermolecular H-bonding decreases the boiling point.  
 $S_4$  : A symmetrical molecule with identical bonds have non zero dipole moment.  
 (A) T F T F (B) T T F T (C) T T F F (D) T T T F
10. H - bonding is maximum in
 

(A)  $C_6H_5OH$  (B)  $C_6H_5COOH$  (C)  $CH_3CH_2OH$  (D)  $CH_3COCH_3$
11. Select the correct statement.
 

(A) The order of Xe—F bond length in various fluorides of Xenon is  $XeF_2 < XeF_4 < XeF_6$   
(B)  $PH_5$  can undergo  $sp^3d$  hybridisation to have octahedral geometry.  
(C) Dipole moment of  $CH_3F$  is greater than that of  $CH_3Cl$   
(D) Increasing strength of hydrogen bonding is  $Cl-H \cdots Cl < N-H \cdots N < O-H \cdots O < F-H \cdots F$
12. Which of the following compounds would have significant intermolecular hydrogen bonding ?  
 $HF, CH_3OH, N_2O_4, CH_4, NH_3(l)$ 

(A)  $HF, N_2O_4, NH_3(l)$  (B)  $HF, CH_4, CH_3OH$  (C)  $HF, CH_3OH, NH_3(l)$  (D)  $CH_3OH, CH_4, NH_3(l)$
13. Which one of the following does not have intermolecular H-bonding ?
 

(A)  $H_2O$  (B) o-nitro phenol (C)  $HF$  (D)  $CH_3COOH$
14. The increasing order of the strength of hydrogen bond in the following mentioned linkages is :  
 (i)  $O-H \cdots S$  (ii)  $S-H \cdots O$  (iii)  $F-H \cdots F$  (iv)  $F-H \cdots O$   
 (A) (i) < (ii) < (iv) < (iii) (B) (ii) < (i) < (iv) < (iii) (C) (i) < (ii) < (iii) < (iv) (D) (ii) < (i) < (iii) < (iv)
15. Give the correct order of initials T or F for following statements. Use T if statement is true and F if it is false.  
 $S_1$  :  $HF$  boils at a higher temperature than  $HCl$   
 $S_2$  :  $HBr$  boils at lower temperature than  $HI$   
 $S_3$  : Bond length of  $N_2$  is less than  $N_2^+$   
 $S_4$  :  $F_2$  has higher boiling point than  $Cl_2$   
 (A) T F T T (B) T T F F (C) T T T F (D) T T T T

16.  $S_1$  : The polarising power of a cation and polarisability of an anion, both are directly proportional to their sizes.  
 $S_2$  :  $H_2^+$  and  $He_2^+$  have same bond order but  $H_2^+$  is more stable than  $He_2^+$ .  
 $S_3$  : The strength of hydrogen bond does not depend at all on the availability of the lone pair of electrons on the atom forming H-bond.  
 $S_4$  :  $OF_2$  and  $Cl_2O$  both are  $sp^3$  hybridised and bond angle in  $Cl_2O$  is greater than  $109^\circ 28'$ .  
 (A) T F F T (B) F F F T (C) F T F T (D) T T T T
17. Which of the following has minimum melting point  
 (A) CsF (B) HCl (C) HF (D) LiF
18. **Consider the following statements.**  
 $S_1$  : The percentage of s-character in the orbital forming S – S bonds and P – P bonds in  $S_8$  and  $P_4$  molecules respectively are same.  
 $S_2$  : In  $SF_4$  the bond angles, instead of being  $90^\circ$  and  $180^\circ$  are  $89^\circ$  and  $177^\circ$  respectively due to the repulsions between lone pair and bond pairs of electrons.  
 $S_3$  : Aqueous  $H_3PO_4$  is syrupy (i.e more viscous than water)  
 $S_4$  :  $SiO_2$  crystal may be considered as giant molecule in which eight-membered rings are formed with alternate silicon and oxygen atoms.  
 Of these :  
 (A)  $S_1$  &  $S_4$  are correct only. (B)  $S_2$ ,  $S_3$  &  $S_4$  are correct only.  
 (C)  $S_1$ ,  $S_2$ ,  $S_3$  &  $S_4$  are correct. (D)  $S_1$ ,  $S_2$  &  $S_3$  are correct only.
19. Select the correct statement for the sulphuric acid.  
 (I) It has high boiling point and viscosity.  
 (II) There are two types of bond lengths in its bivalent anion.  
 (III)  $p\pi-d\pi$  bonding between sulphur and oxygen is observed.  
 (IV) Sulphur has the same hybridisation that is of boron in diborane.  
 (A) II and III only (B) II, III and IV only  
 (C) I, III and IV only (D) III and IV only
20. Intermolecular hydrogen bonding increases the enthalpy of vaporization of a liquid due to the :  
 (A) decrease in the attraction between molecules.  
 (B) increase in the attraction between molecules.  
 (C) decrease in the molar mass of unassociated liquid molecules.  
 (D) increase in the effective molar mass of hydrogen - bonded molecules.

### More than one choice type

21. The halogen form compounds among themselves with formula  $XX'$ ,  $XX'_3$ ,  $XX'_5$  and  $XX'_7$  where X is the heavier halogen. Which of the following pairs representing their structures and being polar and non-polar are correct?  
 (A)  $XX'$  – Linear – polar (B)  $XX'_3$  – T-shaped – polar  
 (C)  $XX'_5$  – square pyramidal – polar (D)  $XX'_7$  – Pentagonal bipyramidal – non-polar
22. Which of the following is/are correct statement(s) for dipole moment ?  
 (A) Lone pair of electrons present on central atom can give rise to dipole moment.  
 (B) Dipole moment is vector quantity.  
 (C)  $CO_2$  molecule has dipole moment.  
 (D) Difference in electronegativities of combining atoms can lead to dipole moment.
23. Which of the following molecules have intermolecular hydrogen bonds ?  
 (A)  $KH_2PO_4$  (B)  $H_3BO_3$   
 (C)  $C_6H_2CO_2H$  (D)  $CH_3OH$

24. Which is **correct** statement :
- (A) Borazine has higher intermolecular force of attraction as compared to benzene.  
 (B)  $K_{a2}$  fumaric acid is more than  $K_{a2}$  of maleic acid due to intra molecular hydrogen bonding in maleic acid.  
 (C) The O – O bond length in  $O_2[AsF_4]$  is shorter than  $KO_2$ .  
 (D) The bond angle order in halogen – S – halogen is  $OSF_2 < OSeCl_2 < OSBr_2$
25. Select the correct statement(s).
- (A)  $Br_2$  and  $ICl$  have the same number of electrons and thus both have nearly the same boiling points.  
 (B)  $N_2H_4$  is pyramidal about each N-atom.  
 (C) In  $P_4S_3$  molecule, there are six P-S bonds, three P-P bonds and ten lone pairs of electrons (on all atoms).  
 (D) In  $ClO_4^-$ , all Cl – O bonds are identical and there is strong  $p\pi - d\pi$  bonding between chlorine and oxygen atoms.
26. Which of following is correct
- |                                   |   |
|-----------------------------------|---|
| (A) $PH_3 < AsH_3 < SbH_3 < NH_3$ | order of boiling point                                  |
| (B) $D_2O(s) > H_2O(l)$           | order of density  |
| (C) $Mn > Ca > Sr > Rb$           | order of metallic bond strength                         |
| (D) $H_2 < CO_2 < H_2O$           | increasing order of intermolecular forces of attraction |

### PART - III : MATCH THE COLUMN

- |   |   |
|---|---|
| 1. <b>Column – I</b><br>(A) $SO_3$ (gas)<br>(B) $OSF_4$<br>(C) $SO_3F^-$<br><br>(D) $ClOF_3$  | <b>Column – II</b><br>(p) Polar with $p\pi-d\pi$ bonds and identical S–O bond, lengths.<br>(q) One lone pair and $p\pi - d\pi$ bond.<br>(r) Non-polar with $p\pi - p\pi$ and $p\pi - d\pi$ bonds. Identical S–O bond lengths.<br>(s) Polar with $p\pi - d\pi$ bond. |
| 2. <b>Column – I</b><br>(A) $IF_2^-$<br>(B) $ClF_3$<br>(C) $XeO_3F_2$<br>(D) $SF_4$   | <b>Column – II</b><br>(p) $sp^3d$<br>(q) polar<br>(r) one of the bond angles is $180^\circ$ .<br>(s) one lone pair  |
| 3. <b>Column-I</b><br>(A) $HCl < HF$<br>(B) $PH_3 < NH_3$<br>(C) $H_2O < D_2O$<br>(D) $F_2 < Cl_2$  | <b>Column-II</b><br>(p) Strength of hydrogen bonding<br>(q) Dipole moment<br>(r) Boiling point<br>(s) Bond energy   |
| 4. <b>Column-I</b><br>(A) Blue vitriol<br>(B) Gypsum<br>(C) Pure orthophosphoric acid<br>(D) Chloral hydrate  | <b>Column-II</b><br>(p) Ionic bond<br>(q) Covalent bond<br>(r) Hydrogen bond<br>(s) Resonance stabilisation   |
| 5. <b>Column-I</b><br>(A) Liquid bromine<br>(B) Solid hydrogen fluoride<br>(C) Solution of sodium fluoride in water<br>(D) Liquid methylamine<br>(E) Noble gas clathrate. | <b>Column-II</b><br>(p) Hydrogen bond<br>(q) Ion-dipole force<br>(r) Dispersion force.<br>(s) Dipole induced dipole interaction.  |

## PART - IV : COMPREHENSION

### COMPREHENSIONS :

Read the following passage carefully and answer the questions.

#### Comprehension : 1

The degree of polarity of a covalent compound is measured by the dipole moment ( $\mu_{\text{bond}}$ ) of the bond defined as:

$$\mu_{\text{bond}} = \text{Charge on one of the poles} \times \text{bond length}$$

$\mu_{\text{bond}}$  is a vector quantity. The dipole moment of a molecule is the vector addition of all the bond dipole moments present in it. For a triatomic molecule, containing two bond's like  $\text{H}_2\text{O}$ ,  $\mu_{\text{molecule}}$  is given by

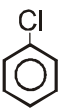
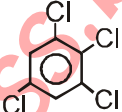
$$\mu_{\text{molecule}}^2 = \mu_{\text{bond}}^2 + \mu_{\text{bond}}^2 + 2\mu_{\text{bond}} \cdot \mu_{\text{bond}} \cos \theta$$

$\theta$  = bond angle

The % ionic character of a bond is calculated using the equations

$$\% \text{ ionic character} = \frac{\mu_{\text{obs}}}{\mu_{\text{ionic}}} \times 100$$

$\mu_{\text{ionic}}$  = dipole moment when the molecule is assumed to be completely ionic.

- Which of the following molecule has non-zero dipole moment :  
(A)  $\text{XeF}_2$  (B)  $\text{ClF}_3$  (C)  $\text{XeO}_2\text{F}_4$  (D)  $\text{XeF}_4$
- The dipole moment of  is 1.5 D. The dipole moment of  will be -  
(A) 0 D (B) 1.5 D (C) 2.86 D (D) 2.25 D
- Which of the following compound has Zero dipole moment -  
(A)  $\text{PCl}_3$  (B)  $\text{PCl}_2\text{F}_3$  (C)  $\text{PCl}_3\text{F}_2$  (D)  $\text{PClF}_4$

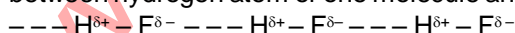
#### Comprehension : 2

Molecular geometry is the general shape of a molecule as determined by the relative positions of the atomic nuclei. VSEPR model predicts the shape of the molecules & ions in which valence shell electron pairs are arranged about the atom as far away from one another as possible, thus minimizing pair repulsion information about the geometry of a molecule can sometimes be obtained from an experimental quantity called dipole moment.

- The dipole moment of a triatomic molecule  $\text{AX}_2$  was found to be equal to the bond moment of  $\text{A}-\text{X}$  bond. Which of the following information regarding geometry of the molecule can be drawn from the above observation.  
(A) Molecule is linear  
(B) Molecule is V shaped with  $\angle \text{X}-\text{A}-\text{X} = 90^\circ$   
(C) Molecule is V shaped with  $\angle \text{X}-\text{A}-\text{X} = 120^\circ$   
(D) Molecular geometry can not be predicted with the given information
- Which of the following inter-halogen compounds is non-polar in nature:  
(A)  $\text{ClF}_3$  (B)  $\text{BrF}_5$  (C)  $\text{IF}_7$  (D)  $\text{BrCl}$

#### Comprehension : 3

Nitrogen, oxygen and fluorine are the highly electronegative elements. When they are tied to a hydrogen atom to form covalent bond, the electrons of the covalent bond are shifted towards the more electronegative atom. This partially positively charged hydrogen atom forms a bond with the other electronegative atom. This bond is called as hydrogen bond and is weaker than covalent bond. For example, in  $\text{HF}$  molecule, the hydrogen bond exists between hydrogen atom of one molecule and fluorine atom of another molecule as depicted :



Here, hydrogen bond acts as a bridge between atoms which holds one atom by covalent bond and the other by hydrogen bond. Hydrogen bond is represented by a dotted line (---) while a solid line represents the covalent bond. Thus, 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.

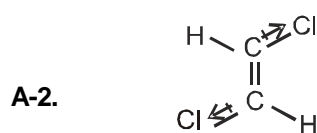


# Answers

## EXERCISE - 1

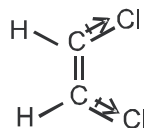
### PART-I

- A-1.** Inorganic benzene ( $N_3B_3H_6$ ) contains polar covalent B – N bonds while benzene ( $C_6H_6$ ) contains non-polar covalent C–C bonds.



trans-form

Two chlorine atom lie opposite direction, so net dipole moment will be zero



cis-form

Two chlorine atom lie on the same side of C = C. So there will be some net dipole moment

- A-3.**  $CCl_4$  is a symmetrical and non polar molecule while  $CHCl_3$  is an unsymmetrical and polar molecule.
- A-4.**  $BF_3 < H_2S < H_2O$ .
- A-5.** 84.5%
- B-1.** Very strong hydrogen bonding occurs in the alkali metal hydrogen fluorides of formula  $M[HF_2]$ ; there is a linear symmetrical anion having an over all, F–H–F distance of 2.26 Å.  
 $[F - H \cdots F]^- \longleftrightarrow [F \cdots H - F]^-$   
 $F^- + HF \longrightarrow [FHF]^-$ ;  $\Delta H = -161 \pm 8 \text{ kJ mol}^{-1}$
- B-2.** Deuterium is more electropositive than hydrogen. Therefore, stronger H-bonding is found in  $D_2O$  than in  $H_2O$ .  $D_2O$  is also denser than  $H_2O$ .
- C-1.** These compounds contain polar–OH groups which can form H-bonds with water.
- C-2.** In ethanol, there is H-bonding but in diethyl ether, there is no H-bonding (because O-atom is attached to C-atom) but there exists weak dipole-dipole attraction in diethyl ether.

### PART-II

- |                 |                 |                 |                 |                 |
|-----------------|-----------------|-----------------|-----------------|-----------------|
| <b>A-1.</b> (A) | <b>A-2.</b> (C) | <b>A-3.</b> (B) | <b>A-4.</b> (D) | <b>A-5.</b> (D) |
| <b>A-6.</b> (A) | <b>A-7.</b> (A) | <b>A-8.</b> (B) | <b>A-9.</b> (B) | <b>B-1.</b> (D) |
| <b>B-2.</b> (C) | <b>B-3.</b> (D) | <b>B-4.</b> (D) | <b>B-5.</b> (D) | <b>B-6.</b> (D) |
| <b>B-7.</b> (C) | <b>C-1.</b> (B) | <b>C-2.</b> (D) | <b>C-3.</b> (B) | <b>C-4.</b> (D) |
| <b>C-5.</b> (C) | <b>C-6.</b> (D) |                 |                 |                 |

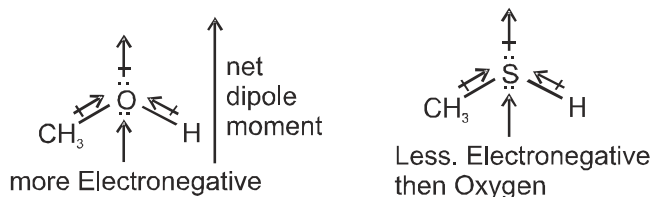
### PART-III

- |               |               |               |               |               |
|---------------|---------------|---------------|---------------|---------------|
| <b>1.</b> (A) | <b>2.</b> (B) | <b>3.</b> (A) | <b>4.</b> (B) | <b>5.</b> (C) |
| <b>6.</b> (D) | <b>7.</b> (A) | <b>8.</b> (A) | <b>9.</b> (A) |               |

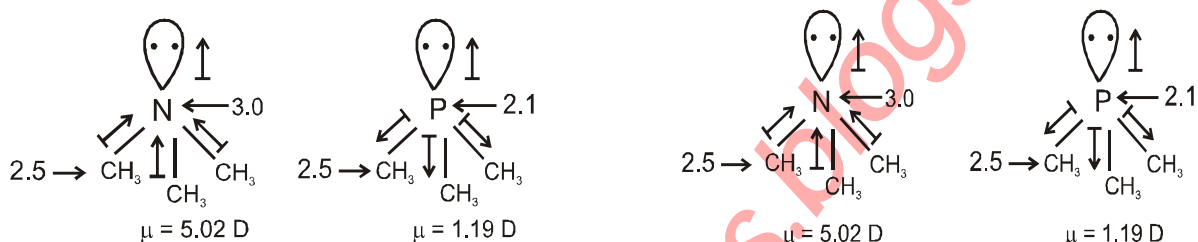
## EXERCISE - 2

### PART-I

1.  $\text{CH}_3 - \text{O} - \text{H}$  is having more dipole moment than  $\text{CH}_3\text{SH}$  because in  $\text{CH}_3\ddot{\text{O}}\text{H}$  the central oxygen atom is more electronegative than sulphur atom. So net dipole moment is more in  $\text{CH}_3\ddot{\text{O}}\text{H}$ .



2. Nitrogen is more electronegative than phosphorus.



So, dipole moment of trimethylamine is greater than trimethyl phosphine.

3. Benzene molecules are held together by dispersion London forces (a type of van der Waal's forces)
4.  $\text{ICl}$  has dipole - dipole attraction due to polar nature but  $\text{Br}_2$  being non-polar experiences very weak dispersion London forces.

5. has intramolecular H-bonding, hence, it does not associate with other molecules to significant extent. Therefore, it exists in liquid state. However, in , due to intermolecular H-bonding, there is association of molecules leading to solid state.
- o-hydroxy benzaldehyde

6. 80% covalent
7. Dipole moment is product of charge and the bond length. In  $\text{CH}_3\text{F}$  the charge is higher but due to greater bond length in  $\text{CH}_3\text{Cl}$  the dipole moment comes out to be higher (remember as a fact)

### PART-II

- |               |             |               |               |             |
|---------------|-------------|---------------|---------------|-------------|
| 1. (B)        | 2. (A)      | 3. (C)        | 4. (C)        | 5. (C)      |
| 6. (C)        | 7. (D)      | 8. (B)        | 9. (C)        | 10. (B)     |
| 11. (D)       | 12. (C)     | 13. (B)       | 14. (A)       | 15. (B)     |
| 16. (B)       | 17. (B)     | 18. (C)       | 19. (C)       | 20. (B)     |
| 21. (A,B,C,D) | 22. (A,B,D) | 23. (A,B,C,D) | 24. (A,B,C,D) | 25. (B,C,D) |
| 26. (B,C,D)   |             |               |               |             |

**PART-III**

1.  $(A - r) ; (B - s) ; (C - p) ; (D - q)$
2.  $(A - p, r) ; (B - p, q) ; (C - p, r) ; (D - p, q, s)$
3.  $(A - p, q, r, s) ; (B - p, q, r, s) ; (C - p, q, r, s) ; (D - r, s)$
4.  $(A - p, q, r, s) ; (B - p, q, r, s) ; (C - q, r) ; (D - q, r)$
5.  $(A - r) ; (B - p, q, r) ; (C - p, q, r) ; (D - p, r) ; (E - p, r, s).$

**PART-IV**

- |        |        |        |        |        |
|--------|--------|--------|--------|--------|
| 1. (B) | 2. (B) | 3. (C) | 4. (C) | 5. (C) |
| 6. (A) | 7. (A) | 8. (C) |        |        |

**EXERCISE - 3**

**PART-I**

- |        |        |      |
|--------|--------|------|
| 1. (A) | 2. (B) | 3. 4 |
|--------|--------|------|

**PART-II**

- |        |        |        |        |        |
|--------|--------|--------|--------|--------|
| 1. (D) | 2. (C) | 3. (A) | 4. (D) | 5. (B) |
|--------|--------|--------|--------|--------|

## Advanced Level Problems

### PART - I : OBJECTIVE QUESTIONS

#### Single choice type

- Among the following compounds the one that is polar and has central atom with  $sp^3$  hybridisation is :  
(A)  $H_2CO_3$  (B)  $SiF_4$  (C)  $BF_3$  (D)  $HClO_2$
- Which of the following are polar  
(A)  $XeF_4$  (B)  $SO_3$  (C)  $XeOF_4$  (D)  $ICl_4^-$
- Which of the following statements is true?  
(A) The dipole moment of  $NF_3$  is zero (B) The dipole moment of  $NF_3$  is less than  $NH_3$   
(C) The dipole moment of  $NF_3$  is more than  $NH_3$  (D) The dipole moment of  $NH_3$  is zero
- Among the following compounds, the correct order of the polarity of the bonds is :  
 $SbH_3, AsH_3, PH_3, NH_3$ .  
(A)  $SbH_3 < AsH_3 < PH_3 < NH_3$  (B)  $AsH_3 < SbH_3 = PH_3 < NH_3$   
(C)  $PH_3 < AsH_3 < SbH_3 < NH_3$  (D)  $AsH_3 < PH_3 < SbH_3 < NH_3$
- Among the following, van der Waals forces are maximum in :  
(A)  $HBr$  (B)  $LiBr$  (C)  $LiCl$  (D)  $AgBr$
- Intermolecular hydrogen bond is present in which of the following pair of molecules ?  
(A)  $SiH_4$  and  $SiF_4$  (B)  $CH_3 - \overset{\overset{O}{\parallel}}{C} - CH_3$  and  $CHCl_3$   
(C)  $H - \overset{\overset{O}{\parallel}}{C} - OH$  and  $CH_3 - \overset{\overset{O}{\parallel}}{C} - OH$  (D)  $CH_3OCH_3$  and  $H_2O_2$
- Which of the following exhibits H-bonding ?  
(A)  $CH_4$  (B)  $H_2Se$  (C)  $N_2H_4$  (D)  $H_2S$
- Bicarbonate ( $HCO_3^-$ ) exists in  $KHCO_3$  and  $NaHCO_3$  respectively as :  
(A) Dimeric and polymeric chain like structure. (B) Polymeric chain and dimeric structure.  
(C) Dimeric and trimeric structure. (D) Trimeric and dimeric structure.
- Covalent compounds have low melting point because  
(A) Covalent bond is less exothermic  
(B) Covalent molecules have definite shape  
(C) Covalent bond is weaker than ionic bond  
(D) Covalent molecules are held by weak Vander Waal's force of attraction

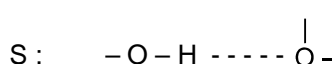
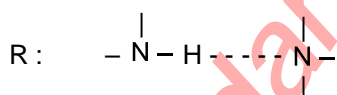
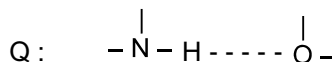
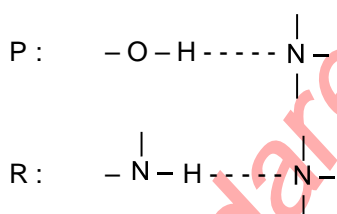
10.  $\text{SnCl}_2 \cdot 2\text{H}_2\text{O}$  readily loses one molecule of water at  $80^\circ\text{C}$  because :  
 (A) One water molecule is coordinated to lone pair of electrons on  $\text{SnCl}_2$  and the other is hydrogen bonded to coordinated water molecule.  
 (B) One water molecule is bonded to  $\text{SnCl}_2$  by coordinate covalent bond and the other is held by ion-dipole attractive forces to central metal ion.  
 (C) Both the water molecules are coordinated to lone pairs of electrons on pyramidal  $\text{SnCl}_2$ .  
 (D) Both the water molecules are bonded to pyramidal  $\text{SnCl}_2$  by hydrogen bonds.
11. The pairs of bases in DNA are held together by  
 (A) Hydrogen bonds (B) Ionic bonds (C) Phosphate groups (D) Deoxyribose groups
12. The bond that determines the secondary structure of a protein is  
 (A) Coordinate bond (B) Covalent bond (C) Hydrogen bond (D) Ionic bond
13. Pure phosphoric acid is very viscous, because :  
 (A) It is a strong acid (B) It is tribasic acid (C) It is hygroscopic  
 (D) It has  $\text{PO}_4^{3-}$  groups which are bonded by many hydrogen bonds
14. Which of the following is least volatile ?  
 (A) HF (B) HCl (C) HBr (D) HI
15. Which of the following is false ?  
 (A) Van der Waals forces are responsible for the formation of molecular crystals.  
 (B) Branching lowers the boiling points of isomeric organic compounds due to reduction in the van der Waals force of attraction.  
 (C) In graphite, van der Waals forces act between the carbon layers.  
 (D) Boiling point of  $\text{NH}_3$  is greater than  $\text{SbH}_3$ .

### More than one choice type

16. Which of the following statements is/are correct :  
 (A) Individual oxidation number of two sulphur atoms in thiosulphate ( $\text{S}_2\text{O}_3^{2-}$ ) ion are +4 and 0, however the average oxidation number of sulphur is +2.  
 (B) The reason for  $K_{a2} \ll K_{a1}$  for peroxymonosulphuric acid is, intramolecular H-bonding in the anion of acid after first ionisation.  
 (C)  $\text{NH}_3$  has a higher boiling point than  $\text{SbH}_3$ , because of H-bonding between  $\text{NH}_3$  molecules.  
 (D) Among HCl, HBr and HI, HI is the strongest acid while HCl is the weakest acid while among HOCl, HOBr and HOI, HOI is the strongest acid while HOI is the weakest acid.
17. Which of the following are polar ?  
 (A)  $\text{XeF}_4$  (B)  $\text{XeF}_6$  (C)  $\text{XeOF}_4$  (D)  $\text{XeF}_5^-$
18. Which of the following statement(s) is/are correct?  
 (A) Ethyne gas is more soluble in acetone than in water.  
 (B)  $\text{CH}_3\text{F}$  is more polar than  $\text{CD}_3\text{F}$  due to deuterium (D) being less electronegative than hydrogen (H).  
 (C) Silyl isocyanate ( $\text{SiH}_3\text{NCO}$ ) is linear in shape while methyl isocyanate ( $\text{CH}_3\text{NCO}$ ) is bent in shape.  
 (D) All of these
19. Which of the following statements is **correct** regarding phosphoric acid ?  
 (A)  $p\pi-d\pi$  back bonding exist between O and P  
 (B) The anion is resonance stablized  
 (C) It is a dibasic acid  
 (D) Inter molecular H bonding between molecules make it a syrupy (viscous) liquid.
20. The critical temperature of water is higher than that of  $\text{O}_2$  because the  $\text{H}_2\text{O}$  molecule has :  
 (A) fewer electrons than  $\text{O}_2$  (B) two covalent bonds  
 (C) V - shape (D) dipole moment

## PART - II : SUBJECTIVE QUESTIONS

- Which among the following will have zero dipole moment.  
 $\text{H}_2\text{O}$ ,  $\text{CO}_2$ ,  $\text{BF}_3$ ,  $\text{NH}_3$
- Out of the given bonds, which one is most polar?  
(a) C–O                      (b) C–F                      (c) O–F                      (d) N–F
- HF forms H-bonding with acetylene while it is a non polar molecule. Explain.
- It is said that two –OH groups present on a carbon are unstable. But chloral hydrate  $\text{CCl}_3\text{CH}(\text{OH})_2$  is fairly stable. Explain.
- Thio - ethers have higher boiling point than ethers. Explain.
- Two atoms A and B have electronegativities as 2.1 and 3.5. These two atoms are covalently single bonded and compound AB has experimental dipole moment equal to 1.4 D. Calculate AB bond length in Å (Use Hanny smith formula for calculation of percentage ionic character).
- Dipole moment of LiF was experimentally determined and was found to be 6.0 D. Calculate percentage of ionic character in LiF molecule. LiF bond length is 2.5 Å.
- The gaseous metallic chloride ( $\text{MCl}$  type) molecule has a measured dipole moment of 9.0 D, which indicates that it is a very polar molecule. The separation between the nuclei in this molecule is  $2.25 \times 10^{-8}$  cm. Calculate the percentage ionic character in KCl molecule. [ $1.6 \times 10^{-19}$ ,  $e = 4.8 \times 10^{-10}$  esu]
- The bond angle in  $\text{H}_2\text{S}$  is  $97^\circ$  and its dipole moment is 1.5 D. The S – H bond distance is 0.15 nm. Therefore, approximate percentage ionic character of S – H bond is (neglect the effect of dipole moment of lone pair on sulphur atom in  $\text{H}_2\text{S}$ ). (Given [ $\cos 97^\circ = -0.12$ ] and  $\sqrt{0.88} = 0.94$ )
- Arrange the following in the increasing order as stated below :  
(i) Polarisability (for the same cation) :  $\text{O}^{2-}$ ,  $\text{F}^-$ ,  $\text{N}^{3-}$   
(ii) Covalent character : LiI, LiBr, LiF  
(iii) H-bond strength :



(iv) Boiling point :  $\text{NH}_3$ ,  $\text{AsH}_3$ ,  $\text{PH}_3$

- Answer the following questions :  
(P) The number of  $p\pi-d\pi$  bonds in  $\text{S}_3\text{O}_9$ .  
(Q) The bond order of the underlined species ;  $\text{NOH}\underline{\text{SO}}_4$ .  
(R) The number of vacant hybrid orbitals which participate in the formation of 3-centre 2 electron bonds i.e., banana bonds in diborane structure.  
(S) The number of intramolecular H-bonds formed in a molecule of chloral hydrate.

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(P)    (Q)    (R)    (S)

12. The dipole moment of HBr is  $2.60 \times 10^{-30}$  Cm and the inter-atomic distance is 1.41 Å. What is the per cent ionic character of HBr ?
13. A diatomic molecule has a dipole moment of 1.2 D. If bond length is 1.0 Å, what percentage of an electronic charge exists on each atom.
14. Arrange the following  
 (i)  $N_2$ ,  $O_2$ ,  $F_2$ ,  $O_2^{+}$  in increasing order of bond dissociation energy.  
 (ii) O, S, F, Cl, N; in increasing strength of hydrogen bonding ( $X-H \cdots X$ ).  
 (iii)  $N_3^-$ ,  $BF_3$ ,  $NH_3$ ,  $XeF_4$  in increasing bond angle.
15. Explain the H-bonding in crystalline  $NaHCO_3$  and  $KHCO_3$  ?
16. The dipole moment of a molecule is really the vectorial sum of the individual bond moment present in it. To compute the dipole moment it is necessary to find out the values of various bond moment. in the following table dipole moment of different bonds are as given.

Bond	$\overrightarrow{H-C}$	$\overrightarrow{C-Cl}$	$\overrightarrow{C=O}$
Bond moments	0.4 D	1.5 D	2.5 D

The group moments of few group as given

Group	$NO_2$	$OH$	$CN$	$CH_3$
direction of dipole	toward N	towards O	toward N	away from $CH_3$
Dipole moment	4D	1.6 D	3.8 D	0.4 D

(i) In  $CH_3CCl_3$  (I),  $CHCl_3$  (II) and  $CH_3Cl$  (III) the normal tetrahedral bond angle is maintained. Also given

$\cos 70.5^\circ = \frac{1}{3}$ . Find dipole moments of the given compounds. (given due to -I effect of Cl, the Bond moment

of  $H-C$  bond directed toward the H in  $CHCl_3$ .)

(ii) In the acetone molecule considering the normal planer structure, find the observed dipole moment of acetone molecule.

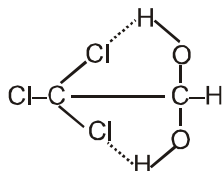
# Answers

## PART-I

- |            |            |            |               |         |
|------------|------------|------------|---------------|---------|
| 1. (D)     | 2. (C)     | 3. (B)     | 4. (C)        | 5. (D)  |
| 6. (C)     | 7. (C)     | 8. (A)     | 9. (D)        | 10. (A) |
| 11. (A)    | 12. (C)    | 13. (D)    | 14. (A)       | 15. (D) |
| 16. (B, D) | 17. (B, C) | 18. (A, C) | 19. (A, B, D) | 20. (D) |

## PART-II

1.  $\text{CO}_2$   $\text{:}\ddot{\text{O}}\equiv\text{C}\equiv\ddot{\text{O}}\text{:}$  and  $\text{BF}_3$
2. C-F bond is most polar due to maximum difference in electronegativity.
3.  $\text{H}-\text{C}\equiv\text{C}-\text{H}$   $\text{sp}$  Hybridisation.  
Due to  $\text{sp}$  hybridisation of C atom, the electronegativity of C is more and hence can form H-bond with HF.
4. The structure of chloral hydrate is as follows



Its stability is due to intermolecular H-bonding.

5. Thio-ethers have higher boiling points due to their higher molecular weights than that of ethers.
6.  $1 \text{ \AA}$
7. 50%.
8. 83.33%
9. 16 %
10. (i)  $\text{N}^{3-} > \text{O}^{2-} > \text{F}^-$  (ii)  $\text{LiI} > \text{LiBr} > \text{LiCl}$  (iii)  $\text{P} > \text{S} > \text{R} > \text{Q}$  (iv)  $\text{NH}_3 > \text{AsH}_3 > \text{PH}_3$
11. 

6	3	2	2
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(P) (Q) (R) (S)
12. 11.5%
13. 25%
14. (i)  $\text{F}_2 < \text{O}_2 < \text{O}_2^+ < \text{N}_2$  (ii)  $\text{S} < \text{Cl} < \text{N} < \text{O} < \text{F}$  (iii)  $\text{XeF}_4 < \text{NH}_3 < \text{BF}_3 > \text{N}_3^-$
15. Inter molecular H-bonding in both but  $\text{KHCO}_3$  forms dimers through H-bonding and  $\text{NaHCO}_3$  form infinite long chains through H-bonding.
16. (i) I = 1.9 D, II = 1.1 D, III = 1.9 D (ii) 2.9 D