

#### **CHEMISTRY**

#### **BOOKS - VK JAISWAL ENGLISH**

### **CHEMICAL BONDING (ADVANCED)**

#### Level 1

- **1.** On decreasing intermolecular distance below the optimum distance (where potential energy is minimum), there is steep increase in potential energy due to:
  - A. Increase in force of attraction between electrons and nucleus
  - B. Increase in stability of bonded atoms
  - C. Equal probability of finding bonding electrons near to either of nuclei

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#### **Answer: D**



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**2.** Identify the correct sequence of increasing number of  $\pi$ -bonds in the structure of the following molecules:

(I) 
$$H_2S_2O_6$$
 (II)  $H_2S_2O_3$  (III)  $H_2S_2S_5$ 

A. I, II and III

B. II, I and III

C. II, III and I

D. I, III and II

#### **Answer: C**



<b>3.</b> $C_2H_2$ is isostructural with					
A. $H_2O_2$					
B. $NO_2$					
C. $SnCl_2$					
D. $CO_2$					
Answer: D					
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<b>4.</b> The shapes of nitrite and nitrile respectively are:					
A. Linear and angular					
B. Angular and linear					
C. Both angular					
D. Both linear					

#### **Answer: B**



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- 5. Linear structure is assumed by:
- (I) NCO (II) $CS_2$  (III) $\overset{+}{N}O_2$  (IV)Solid  $BeH_2$ 
  - A. all four
  - B. (II), (III) and (IV)
  - C. (I), (II) and (III)
  - D. (II) and (III)

#### **Answer: C**



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**6.** Among the oxides of nitrogen  $N_2O$ , NO and  $NO_2$ , molecules with unpaired electrons are:

A.  $N_2O$  and NO $B.NO \text{ and } NO_2$ 

 $C. N_2O$  and  $NO_2$ 

D.  $NO_2$  and Its dimer

### **Answer: B**



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7. Which of the following pair consists of only network solid?

A.  $SiO_2, P_4O_{10}$ 

B.  $P_4O_{10}$ ,  $SO_3$ 

 $C. P_3O_{10}, P_4O_6$ 

D. Diamond,  $SiO_2$ 

Answer: D

**8.** Which of the following ions does not have S-S linkage?

A.  $S_2O_8^{2\,-}$ 

B.  $S_2O_6^{2\,-}$ 

C.  $S_2O_5^{2\,-}$ 

D.  $S_2O_3^{2\,-}$ 

#### Answer: A



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**9.** Among  $KO_2, KAlO_2, CaO_2$  and  $NO_2^+$ , unpaired electrons is present in :

A.  $NO_2^+$  and  $CaO_2$ 

 $B. KO_2$  and  $KAlO_2$ 

C.  $KO_2$  only

#### **Answer: C**



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- **10.** Structure of  $S_2Cl_2$  is analogous to :
  - A.  $SOCl_2$
  - B.  $CO_2$
  - $\mathsf{C}.\,H_2S$
  - D.  $H_2O_2$

#### **Answer: D**



**11.** Number of P-H, P-O-P, P-O-H and P=O bonds in sodium dihydrogen pyrophosphate respectively are:

- A. 1, 1, 1, 2
- B. 0, 1, 2, 2
- $\mathsf{C.}\ 0,\,1,\,2,\,4$
- D. 2, 0, 0, 2

#### **Answer: C**



**12.** Two hybrid orbitals have a bond angle of  $120^{\circ}$ . The percentage of scharacter in the hybrid orbitals is nealy:

- A. 0.25
- B. 0.33
- C. 0.5

D. 0.66

**Answer: B** 



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**13.** The state of hybridisation of central atom in dimer of  $BH_3$  and  $BeH_2$ 

IS:

A.  $sp^2,\,sp^2$ 

B.  $sp^3,\,sp^2$ 

 $\mathsf{C.}\,sp^3,\,sp^3$ 

D.  $sp^2,\,sp^3$ 

#### **Answer: B**



**14.** In  $NO_2$  molecule N atom undergoes in :

A.  $\mathit{sp}^3$  hybridization

B.  $sp^2$  hybridization

C. sp hybridization

D.  $sp^2d$  hybridization

#### **Answer: B**



#### 15. The strongest P-O bond is found in the molecules

A.  $F_3PO$ 

 $\mathsf{B.}\,Cl_3PO$ 

 $\mathsf{C}.\,Br_3PO$ 

D.  $(CH_3)_3PO$ 

#### **Answer: A**



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**16.**  $O_2F_2$  is an unstable yellow orange solid and  $H_2O_2$  is a colourless liquid, both have O-O bond and O-O bond length in  $H_2O_2$  and  $O_2F_2$  respectively is :

- A. 1.22Å, 1.48Å
- B. 1.48Å, 1.22Å
- C. 1.22Å, 1.22Å
- D. 1.48Å, 1.48Å

#### **Answer: B**



17. The bond length of the S-O bond is maximum in which of the following compounds?

A.  $SOCl_2$ 

B.  $SOBr_2$ 

 $\mathsf{C}.\,SOF_2$ 

D. All have same length

#### **Answer: B**



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**18.**  $\angle FAsF$  bond angle in  $AsF_3Cl_2$  molecule is :

A.  $90^{\circ}$  and  $180^{\circ}$ 

B.  $120^{\circ}$ 

 $\mathsf{C.\,90}^\circ$ 

D.  $180^{\circ}$ 

#### Answer: A



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19. Which of the following has largest bond angle?

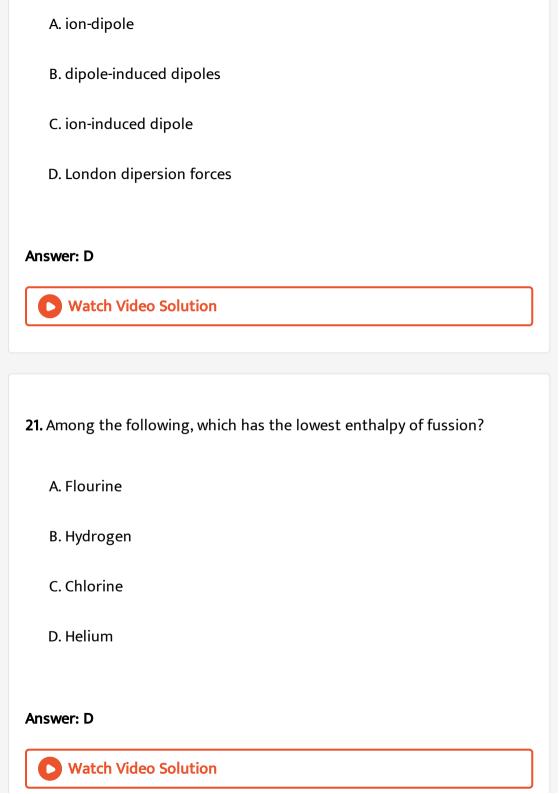
- A.  $H_2O$
- B.  $F_2O$
- $\mathsf{C}.\,Cl_2O$
- D.  $H_2S$

#### **Answer: C**



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**20.** The boiling points of noble gases are illustrative of the operation of forces of the type :



**22.** Out of the two compounds shown below, the vapour pressure of B at a particular temperature is expected to be:

- A. higher than that of A
- B. lower than that of A
- C. same as that of A
- D. can be higher or lower depending upon the size of the vessel

#### Answer: A



- A. ionic forces
- B. covalent bonds
- C. intramolecular H-bonds
- D. covalent as well as H-bonds

#### **Answer: D**



- **24.** The boiling points of methanol, water and dimethyl ether are respectively  $65^{\circ}C$ ,  $100^{\circ}C$  and  $34.5^{\circ}C$ . Which of the following best explains these wide variations in b.p. ?
  - A. The molecular mass increases from water (18) to methanol(32) to dimethyl ether(74)
  - B. The extent of H-bonding decreases from water to methanol while it is absent in ether

C. The extent of intramolecular H-bonding decreases from ether to methanol to water

D. The density of water is  $1.00g \cdot mL^{-1}$ , methanol  $0.7914g \cdot mL^{-1}$ and that of dimethyl ether is  $0.7137q \cdot mL^{-1}$ 

#### Answer: B



## 25. In ice, the length of H-bonds:

A, is less than that of covalent bonds

B. is greater than that of covalent bonds

C. is same as that of covalent bonds

D. can be less greater or same as that of covalent bonds

#### Answer: B



**26.** The correct order of the strength of H-bonds is :

A. 
$$H...F > H...O > H...N$$

$$\operatorname{B.}H.\dots N>H\dots O>H\dots F$$

$$\mathsf{C}.\,H.\ldots O>H\ldots N>H\ldots F$$

$$\mathsf{D}.\,H.\ldots F>H\ldots \,N>H\ldots \,O$$

#### **Answer: A**



- **27.** o-nitrophenol can be easily stream distilled whereas p-nitrophenol cannot be. This is because of :
  - A. Strong intermolecular hydrogen bonding in o-nitrophenol
  - B. Strong intramolecular hydrogen bonding in o-nitrophenol
  - C. Strong intramolecular hydrogen bonding in p-nitrophenol

D. dipole moment of p-nitrophenol is larger than that of o-nitrophenol

#### **Answer: B**



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- 28. What is not true about ice?
  - A. It has open cage like structure
  - B. It has less density than water
  - C. Each O atom is surrounded by 4 H atoms
  - D. Each O atom has four H-bonds around it

#### Answer: D



**29.** When ice melts to form liquid water at  $0\,^\circ$  C, there is a contraction in volume . This is due to

A. the molecules contracting in size

B. a partial disruption of the hydrogen bonded network of ice on melting

C. the adsorption of heat during the melting point process

D. the dissolving in ait into the water during the melting process

#### **Answer: B**



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**30.** In an ionic compound  $A^+X^-$  the degree of covalent bonding is greatest when

A.  $A^+$  and  $X^-$  ion are small

B.  $A^{\,+}$  is small and  $X^{\,-}$  is large

C. $A^+ \; { m and} \; X^-$ ions are approximately of the same size				
D. $X^{-}$ is small and $A^{+}$ is large				

#### **Answer: B**



#### 31. In which of the following species the bonds are non-directonal?

- A.  $NCl_3$
- B. RbCl
- $\mathsf{C}.\,BeCl_2$
- D.  $BCl_3$

#### Answer: B



**32.** Which of the following when dissolved in water forms a solution which is non-conducting?

A. Green vitriol

B. Chile or Indian salt petre

C. Alcohol

D. Potash alum

#### **Answer: C**



33. Which of the following statements about LiC and NaCl is wrong?

A. liCl has lower melting points than NaCl

B. LiCl dissolves more in organic solvents whereas NaCl does not

C. LiCl would ionise in water less than NaCl

D. Fused LiCl would be less conducting than fused NaCl

# Answer: C Watch Video Solution

**34.** Which of the following substance has the highest melting point? .

A. NaCl

B. KCl

 $\mathsf{C}.\,MgO$ 

D. BaO

#### **Answer: C**



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**35.** The stability of ionic crystal depends principally on

A. high electron affinity of anion forming species

B. the lattice energy of crystal

C. low I.E of cation forming solid

D. low heat of sublimation of cation forming solid

#### **Answer: B**



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#### **36.** In which of the following solvents, KI has highest solubility?

A. 
$$C_6H_6(~\in~=0)$$

$$\mathsf{B.}\left(CH_{3}\right)_{2}\!CO(~\in~=2)$$

C. 
$$CH_3OH(~\in~=32)$$

D. 
$$\mathbb{C}Cl_4(\ \in\ =0)$$

#### Answer: C



**37.** Among  $LiCl, RbCl, BeCl_2, MgCl_2$ , the compounds with greatest and least ionic character respectively are

- A. LiCl and RbCl
- $B.\,RbCl$  and  $BeCl_2$
- C. RbCl and  $MgCl_2$
- D.  $MgCl_2$  and  $BeCl_2$

#### **Answer: B**



- **38.** The compouds with the highest degree of covalency is :
  - $\mathsf{A.}\ NaCl$
  - B.  $MgCl_2$
  - $\mathsf{C}.\,AgCl$
  - D. CsCl

#### Answer: C



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- **39.** The salt having the least solubility in water is:
  - A.  $BaCl_2$
  - $\mathsf{B.}\,Ba(NO_3)_2$
  - $\mathsf{C}.\,MgSO_4$
  - $\mathsf{D.}\,BaSO_4$

#### **Answer: D**



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**40.** The solubility of  $Na_2, SO_4, BeSO_4, MgSO_4$  and  $BaSO_4$  will follow the order

A. 
$$BaSO_4 > BeSO_4 > MgSO_4 > Na_2SO_4$$

$$\mathrm{B.}\,Na_2SO_4>BeSO_4>MgSO_4>BaSO_4$$

$$\mathsf{C.}\,BeSO_4>MgSO_4>BaSO_4>Na_2SO_4$$

D. 
$$MgSO_4 > BeSO_4 > Na_2SO_4 > BaSO_4$$

#### **Answer: B**



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**41.** "Solubility of alkali metal fluorides increases down the group " Select correct explanation for given statement:

A. Hydration energy increases and lattice energy decreases down the group

B. Both energy decrease down the group but decrease in hydration energy is rapid

C. Both energy decrease down the group but decrease in lattice energy is rapid

D. Both energy increase down the group but increase in hydration energy is rapid

Answer: C

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# **42.** Covalency favoured in the following case :

A. smaller cation

B. larger anion

C. large charge on cation anions

D. all of these

Answer: D



**43.** The melting point of RbBr is  $682^{\circ}C$ , while that of NaF is  $988^{\circ}C$ . The principle reason that melting point of NaF is much higher than that of RbBr is that :

- A. The molar mass of NaF is smaller than that of RbBr
- B. The bond of RbBr has more covalent character than the bond in

NaF

- C. The difference in electronegativity between Rb and Br is smaller than the difference between Na and F
- D. The inter-nuclear distance,  $r_c + r_a$  is greater for RbBr than for NaF

#### Answer: D



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44. In which of the following compounds B-F bond length is shortest?

A.  $BF_4^{\,-}$ B.  $BF_3ar{r}NH_3$  $\mathsf{C}.\,BF_3$ D.  $BF_3\bar{r}N(CH_3)_3$ **Answer: C Watch Video Solution** 45. Which of the following pair of molecules will have permanent dipole moment? A.  $NO_2$  and  $CO_2$  $B. NO_2$  and  $O_3$  $\mathsf{C}.\,SiF_4$  and  $CO_2$ D.  $SiF_4$  and  $NO_2$ **Answer: B** 

**46.** The observed dipole moment of HCl is 1.03D. If the bond length of HCL is  $1.275 \rm{\AA}$ , then the percent ionic character of H-Cl bond is

A. 0.6

B. 0.39

C. 0.29

D. 0.17

#### Answer: D



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**47.** The dipole moment of o, p and m-dichlorobenzene will be in the order

A. o > p > m

 $\mathtt{B.}\,p>o>m$ 

 $\mathsf{C}.\,m>o>p$ 

 $\mathrm{D.}\,o>m>p$ 

#### **Answer: D**



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48. Which of the following molecule has highest dipole moment?

A.  $BF_3$ 

B.  $NH_3$ 

C.  $NF_3$ 

 $\operatorname{D.}B_2H_6$ 

#### **Answer: B**



**49.** In terms of polar character the correct, the correct order is .

A. 
$$NH_3 < H_2O > HF < H_2S$$

$${\rm B.} \, H_2 S < N H_3 < H_2 O < H F$$

$$\mathsf{C.}\,H_2O < NH_3 < H_2S < HF$$

D. 
$$HF < H_2O < NH_3 < H_2O$$

#### **Answer: B**



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**50.** The correct order of dipole moment is :

A. 
$$CH_4>NF_3>NH_3>H_2O$$

B. 
$$NF_3>CH_4>NH_3>H_2O$$

C. 
$$NH_3>NF_3>CH_4>H_2O$$

D. 
$$H_2O>NH_3>NF_3>CH_4$$

#### **Answer: A**



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#### 51. The correct order of S-O bond length is:

A. 
$$SO_3^{2-} > SO_4^{2-} > SO_3 > SO_2$$

$${\rm B.}\,SO_3^{2\,-}\,>SO_4^{2\,-}\,>SO_2>SO_3$$

$$\mathsf{C.}\,SO_4^{2-} > SO_3^{2-} > SO_2 > SO_3$$

$${\rm D.}\,SO_4^{2-} > SO_3^{2-} > SO_3 > SO_2$$

#### **Answer: B**



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**52.** What is not true about rasonance?

A. The resonating structures are hypothetical

- B. The unpaired electrons in various resonating structures are same
- C. Hybrid structure is most energetic
- D. Hybrid structure is least energetic

#### Answer: C



- **53.** Which of the following conditions is not correct for resonating structures?
  - A. The contributing structures must have the same number of unpaired electrons
  - B. The contributing structures should have almost similar energies
  - C. The contributing structures should be so written that unlike charges reside on atoms that are far apart

D. The positive charge should be present on the electropositive element and the negative charge on the electronegative element

#### Answer: C



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**54.** A molecule is described by three Lewis structures having energies  $E_1$ ,  $E_2$ , and  $E_3$ , respectively. The energies of these structures follow the order

 $E_1 > E_2 > E_3$ , respectively. If the experimental energy of the molecules

is  $E_0$ , the resonance energy is

A. 
$$(E_1 + E_2 + E_3) - E_0$$

B.  $E_0-E_3$ 

C.  $E_0-E_1$ 

D.  $E_0-E_2$ 

#### Answer: B



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**55.** For compounds ,

A: Tetracynoethene

B: Carbon dioxide

C: Benzene

D:1,3-Butaidene.

Ratio of  $\sigma$  and  $\pi$  bonds is in order :

A. 
$$A = B < C < D$$

$$\mathsf{B.}\,A = B < D < C$$

$$\mathsf{C.}\, A = B = C = D$$

$$\mathsf{D}.\, C < C < A < B$$

#### **Answer: A**



## **56.** In a compound

The number of sigma and pi bonds respectively are :  $C = C \\ | C_{2H_5}$ 

- A. 19, 11
- B. 19, 5
- C. 13, 11
- D. 7, 3

### **Answer: A**



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57. Which of the following does not contain a coordinate bond?

- A.  $H_3O^+$
- B.  $BF_{\scriptscriptstyle 4}^{\,-}$

D.  $NH_4^{\ +}$ 

**Answer: C** 



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58. Which of the following does not contain coordinate bond?

A.  $PH_4^{\,+}$ 

 $\mathsf{B.}\,NO_2$ 

 $\mathsf{C}.\,O_3$ 

D.  $CO_3^{2-}$ 

Answer: D



59. Which of the following halides is inert towards hydrolysis at room temperature?

A.  $SiCl_4$ 

B.  $PCl_3$ 

C.  $NCl_3$ 

D.  $NF_3$ 

## **Answer: D**



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60. Which of the following trihalides off nitrogen behaves as weakest base?

A.  $NF_3$ 

B.  $NCl_3$ 

 $\mathsf{C}.\,NBr_3$ 

D.  $NI_3$ 

Answer: A



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61. Increasing order of stability of the +2 oxidation of the ions?

A. 
$$Ca^{2+} < Ba^{2+} < Sr^{2+}$$

$${\rm B.}\, Pb^{2+}\, < Ge^{2+}\, < Sn^{2+}$$

C. 
$$Ge^{2+} < Sn^{2+} < Pb^{2+}$$

D. 
$$Cu^{2+,\,tAu^{2+}\,<\,Ag^{2+}}$$

#### **Answer: C**



**62.** The number of three centre two electron bonds in a molecule of diborane is\_\_\_\_\_.

**A.** 0

B. 2

**C**. 4

D. 6

# Answer: B



**63.** In which of the following compounds octet is complete and incomplete for all atoms :

A.	$Al_2Cl_6$	$Al_2(CH_3)_6$	$AlF_3$	Dimer of	$BeCl_2$	Dimer of	$BeH_2$
	IC	$Al_2(CH_3)_6$ $IC$	IC	C		C	
В.	$Al_2Cl_6$	$Al_2(CH_3)_6$ $IC$	$AlF_3$	Dimer of	$BeCl_2$	Dimer of	$BeH_2$
	C	IC	IC	C		IC	
_	$Al_2Cl_6$	$Al_2(CH_3)_6 \ IC$	$AlF_3$	Dimer of	$BeCl_2$	Dimer of	$BeH_2$
C.	C	IC	C	IC		IC	

D.  $Al_2Cl_6 \ Al_2(CH_3)_6 \ AlF_3 \ Dimer of \ BeCl_2 \ Dimer of \ BeH_2$   $IC \ IC \ IC \ IC \ IC \ Answer: C$ Watch Video Solution

**64.** In which of the following metal to metal bond is present?

B. Stannous chloride

A. Cupric chloride

C. Mercurous chloride

D. Mercutic chloride

**Answer: C** 



**65.** On decreasing intermolecular distance below the optimum distance (where potential energy is minimum), there is steep increase in potential energy due to:

A. Increase in force of attraction between electrons and nucleus

B. Increase in stability of bonded atoms

C. Equal probability of fining bonding electrons near to either of nuclei

D. Increase in interelectronic and internuclear repulsion

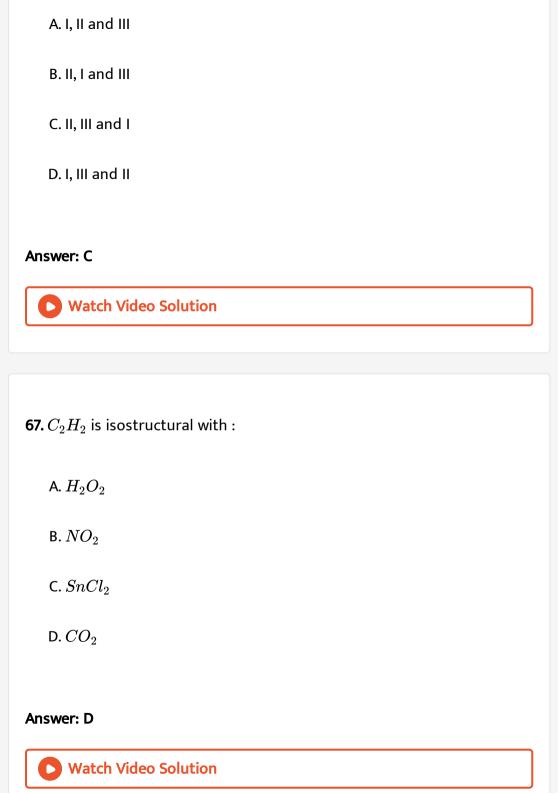
#### **Answer: D**



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**66.** Identify the correct sequence of increasing number of  $\pi$ -bonds in the structure of the following molecules:

(I) $H_2S_2O_6$  (II) $H_2S_2O_3$  (III) $H_2S_2S_5$ 



68. The shapes of nitrite and nitrile respectively are: A. Lineat and angular B. Angular and linear C. Both angular D. Both linear **Answer: B Watch Video Solution** 69. Linear structure is assumed by: (I) NCO (II) $CS_2$  (III) $\overset{+}{N}O_2$  (IV)Solid  $BeH_2$ A. all four B. (II), (III) and (IV) C. (I), (II) and (III)

D. (II) and (III)

#### **Answer: C**



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**70.** Among the oxides of nitrogen  $N_2O,\,NO\,$  and  $NO_2$ , molecules with unpaired electrons are:

A.  $N_2O$  and NO

 $B. NO \text{ and } NO_2$ 

 $\mathsf{C}.\,N_2O$  and  $NO_2$ 

 ${\sf D.}\ NO_2$  and Its dimer

#### **Answer: B**



71. Which of the following pair consists of only network solid?

A.  $SiO_2, P_4O_{10}$ 

 $\operatorname{B.}P_4O_{10},SO_3$ 

C.  $P_3O_{10}, P_4O_6$ 

D. Diamond,  $SiO_2$ 

#### **Answer: D**



# 72. Which of the following ions does not have S-S linkage?

- A.  $S_2O_8^{2\,-}$ 
  - B.  $S_2O_6^{2\,-}$
  - C.  $S_2O_5^{2\,-}$
  - D.  $S_2O_3^{2\,-}$

#### **Answer: A**



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**73.** Among  $KO_2, KAlO_2, CaO_2$  and  $NO_2^+$ , unpaired electrons is present in :

- A.  $NO_2^+$  and  $CaO_2$
- $B. KO_2$  and  $KAlO_2$
- C.  $KO_2$  only
- D.  $CaO_2$  only

## **Answer: C**



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**74.** Structure of  $S_2Cl_2$  is analogous to :

A.  $SOCl_2$ B.  $CO_2$  $\mathsf{C}.\,H_2S$ D.  $H_2O_2$ **Answer: D** Watch Video Solution 75. Number of P-H, P-O-P, P-O-H and P-O bonds in sodium dihydrogen pyrophosphate respectively are: A. 1, 1, 1, 2B.0, 1, 2, 2C. 0, 1, 2, 4D. 2, 0, 0, 2**Answer: C** 

**76.** Two hybrid orbitals have a bond angle of  $120^{\circ}$ . The percentage of scharacter in the hybrid orbitals is nealy:

- A. 0.25
- B. 0.33
- C. 0.5
- D. 0.66

## **Answer: B**



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**77.** The state of hybridisation of central atom in dimer of  $BH_3$  and  $BeH_2$ 

IS:

A.  $sp^2,\,sp^2$ 

- B.  $sp^3,\,sp^2$
- $\mathsf{C.}\,sp^3,\,sp^3$
- D.  $sp^2,\,sp^3$

#### **Answer: B**



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# **78.** In $NO_2$ molecule N atom undergoes in :

- A.  $\mathit{sp}^3$  hybridization
- ${\sf B.}\,sp^2$  hybridization
- C. sp hybridization
- D.  $sp^2d$  hybridization

## **Answer: B**



**79.** The strongest P-O bond is found in the molecules

A.  $F_3PO$ 

 $\mathsf{B.}\,\mathit{Cl}_{3}\mathit{PO}$ 

C.  $Br_3PO$ 

D.  $(CH_3)_3PO$ 

## **Answer: A**



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**80.**  $O_2F_2$  is an unstable yellow change solid and  $H_2O_2$  is a colourless liquid, both have O-O bond and O-O bond length in  $H_2O_2$  and  $O_2F_2$  respectively is :

A. 1.22Å, 1.48Å

B. 1.48Å, 1.22Å

C. 1.22Å, 1.22Å

D. 1.48Å, 1.48Å

**Answer: B** 



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**81.** The bond length of the S-O bond is maximum in which of the following compounds?

A.  $SOCl_2$ 

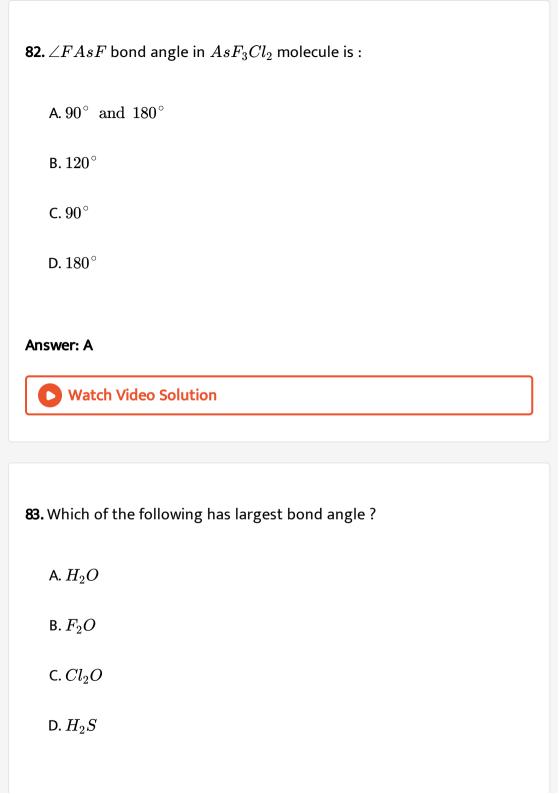
 $\mathsf{B.}\,SOBr_2$ 

 $\mathsf{C.}\,SOF_2$ 

D. All have same length

# Answer: B





# Answer: C



**84.** The boiling points of noble gases are illustrative of the operation of forces of the type :

- A. ion-dipole
- B. dipole-induced dipoles
- C. ion-induced dipole
- D. London dipersion forces

#### Answer: D



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85. Among the following, which has the lowest enthalpy of fussion?

- A. Flourine
- B. Hydrogen
- C. Chlorine
- D. Helium

### **Answer: D**



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**86.** Out of the two compounds shown below, the vapour pressure of B at a particular temperature is expected to be:

- A. higher than that of A
- B. lower than that of A

- C. same as that of A

  D. can be higher or lower depending upon the size of the vessel

  Answer: A
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- 87. The crystal lattice of ice is mostly formed by
  - A. ionic forces
  - B. covalent bonds
  - C. intramolecular H-bonds
  - D. covalent as well as H-bonds

#### **Answer: D**



**88.** The boiling points of methanol, water and dimethyl ether are respectively  $65^{\circ}C$ ,  $100^{\circ}C$  and  $34.5^{\circ}C$ . Which of the following best explains these wide variations in b.p. ?

A. The molecular mass increase from water (18) to methanol(32) to diethyl ether(74)

B. The extent of H-bonding decrease from water too methanol while it is absent in ether

C. The extent of intramolecular H-bondin decrease from ether to methanol to water

D. The density of water is  $1.00g\cdot mL^{-1}$ , methanol  $0.7914g\cdot mL^{-1}$  and that of diethyl ether is  $0.7137g\cdot mL^{-1}$ 

#### Answer: B



- 89. In ice, the length of H-bonds:
  - A. is less than that of covalent bonds
  - B. is greater than that of covalent bonds
  - C. is same as that of covalent bonds
  - D. can be less greater or same as that of covalent bonds

#### **Answer: B**



- **90.** The correct order of the strength of H-bonds is :
  - A.  $H.\ldots F > H.\ldots O > H.\ldots N$
  - $\operatorname{B.}H.\dots N>H\dots O>H\dots F$
  - $\mathsf{C}.\,H.\dots\,O>H\dots\,N>H\dots\,F$
  - $\mathsf{D}.\,H.\dots F>H\dots N>H\dots O$

#### **Answer: A**



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- **91.** o-nitrophenol can be easily stream distilled whereas p-nitrophenol cannot be. This is because of :
  - A. Strong intermolecular hydrogen bonding in o-nitrophenol
  - B. Strong intramolecular hydrogen bonding in o-nitrophenol
  - C. Strong intramolecular hydrogen bonding in p-nitrophenol
  - D. dipole moment of p-nitrophenol is larger than that of o-nitrophenol

#### Answer: B



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92. What is not true about ice?

- A. It has open cage like structure
- B. It has less density than water
- C. Each O atom is surrounded by 4 H atoms
- D. Each O atom has four H-bonds around it

#### **Answer: D**



- **93.** When ice melts to form liquid water at  $0^{\circ}$  C, there is a contraction in volume . This is due to
  - A. the molecules contracting in size
  - B. a partial disruption of the hydrogen bonded network of ice on
    - melting
  - C. the adsorption of heat during the melting point process
  - D. the dissolving in ait into the water during the melting process

#### **Answer: B**



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**94.** In an ionic compound  $A^+X^-$  the degree of covalent bonding is greatest when

- A.  $A^+$  and  $X^-$  ion are small
- B.  $A^{\,+}$  is small and  $X^{\,-}$  is large
- C.  $A^+$  and  $X^-$  ions are approximately of the same size
- D.  $X^-$  is small and  $A^+$  is large

#### **Answer: B**



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95. In which of the following species the bonds are non-directional?

A.  $NCl_3$ B. RbClC.  $BeCl_2$ D.  $BCl_3$ **Answer: B** Watch Video Solution 96. Which of the following when dissolving in water forms a solution which is non-conducting? A. Green vitriol B. Chlile or Indian salt petre C. Alcohol D. Potash alum **Answer: C** 

- 97. Which of the following statements about LiCl and NaCl is wrong?
  - A. liCl has lower melting points than NaCl
  - B. LiCl dissolves more in organic solvents whereas NaCl does not
  - C. LiCl would ionise in water less than NaCl
  - D. Fused LiCl would be less conducting than fused NaCl

#### **Answer: C**



- **98.** Which of the following substance has the highest melting point? .
  - A. NaCl
  - $\mathsf{B.}\ KCl$
  - $\mathsf{C}.\,MgO$

D.	BaO
<b>–</b> .	$\boldsymbol{\mathcal{L}}$

#### Answer: C



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- 99. The stability of ionic crystal principally depends on :
  - A. high electron affinity of anion forming species
  - B. the lattice energy of crystal
  - C. low I.E of cation forming solid
  - D. low heat of sublimation of cation forming solid

#### **Answer: B**



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100. In which of the following solvents, KI has highest solubility?

A. 
$$C_6H_6(~\in~=0)$$

B.  $(CH_3)_2CO(\in = 2)$ 

 $C.CH_3OH(\in =32)$ 

D.  $CCl_4(\in = 0)$ 

## Answer: C



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**101.** Amongst LiCl, RbCl,  $BeCl_2$  and  $MgCl_2$ , the compounds whith the greatrest and the least ionic character respecitely are:

A. LiCl and RbCl

 $B.\,RbCl$  and  $BeCl_2$ 

 $\mathsf{C}.\,RbCl$  and  $MgCl_2$ 

D.  $MgCl_2$  and  $BeCl_2$ 

# **Answer: B**



102. The compouds with the highest degree of covalency is:

A. NaCl

B.  $MgCl_2$ 

 $\mathsf{C}.\,AgCl$ 

D. CsCl

# Answer: C



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103. The salt having the least solubility in water is:

A.  $BaCl_2$ 

B.  $Ba(NO_3)_2$ 

C.  $MgSO_4$ 

D.  $BaSO_4$ 

#### **Answer: D**



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**104.** The solubility of  $Na_2SO_4$ ,  $BeSO_4$ ,  $MgSO_4$  and  $BaSO_4$  in water follow the order :

A. 
$$BaSO_4 > BeSO_4 > MgSO_4 > Na_2SO_4$$

$${\tt B.}\ Na_2SO_4>BeSO_4>MgSO_4>BaSO_4$$

C. 
$$BeSO_4 > MgSO_4 > BaSO_4 > Na_2SO_4$$

D. 
$$MgSO_4 > BeSO_4 > Na_2SO_4 > BaSO_4$$

## Answer: B



**105.** "Solubility of alkali metal fluorices increases down the group " Select correct explanation for given statement:

A. Hydrogen energy increase and lattice energy decrease down the group

B. Both energy decrease down the group but decrease in hydration energy is rapid

C. Both energy decrease down the group but decrease in lattice energy is rapid

D. Both energy increase down the group but increase in hydration energy is rapid

#### Answer: C



A. smaller cation B. larger anion C. large charge on cation anions D. all of these Answer: D **Watch Video Solution 107.** The melting points RbBr is  $682^{\circ}C$ , while that of NaF is  $988^{\circ}$ . The principal reason of this fact is: A. The molar mass of NaF is smaller than that of RbBr B. the bond of RbBr has more covalent character than the bond in NaF C. the difference in electronegativity between Rb and Br is maller the difference between Na and F

D. the intermoleculear distance,  $r_c + r_a$  is greater for RbBr than for

NaF

### **Answer: D**



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108. In which of the following compounds B-F bond length is shortest?

A.  $BF_{\scriptscriptstyle 4}^{\,-}$ 

B.  $BF_3\bar{r}NH_3$ 

 $\mathsf{C}.\,BF_3$ 

D.  $BF_3\bar{r}N(CH_3)_3$ 

# **Answer: C**



109. Which of the following is wheat fruit? A.  $NO_2$  and  $CO_2$ B.  $NO_2$  and  $O_3$  $C. SiF_4$  and  $CO_2$ D.  $SiF_4$  and  $NO_4$ **Answer: B Watch Video Solution** 

**110.** The observed dipole moment of HCl is 1.03 D. If the H-Cl bond length is 1.275 Å, calculate the per cent ionic character in HCl.

- A. 0.6
- B. 0.39
- C. 0.29
- D. 0.17

# **Answer: D**



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**111.** The dipole moment of o, p and m-dichlorobenzene will be in the order :

A. 
$$o > p > m$$

$$\mathtt{B.}\,p>o>m$$

$$\mathsf{C}.\,m>o>p$$

# **Answer: D**



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112. Which of the following molecules has highest dipole moment?

A.  $BF_3$ 

B.  $NH_3$ 

 $\mathsf{C}.\,NF_3$ 

D.  $B_2H_6$ 

# **Answer: B**



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# A. $NH_3 < H_2O > HF < H_2S$

B. 
$$H_2S < NH_3 < H_2O < HF$$

113. In terms of polar of the following order is correct?

C. 
$$H_2O < NH_3 < H_2S < HF$$

D. 
$$HF < H_2O < NH_3 < H_2O$$

# **Answer: B**



114. The correct order of the dipole moment is:

A. 
$$CH_4>NF_3>NH_3>H_2O$$

$$\operatorname{B.}NF_3 > CH_4 > NH_3 > H_2O$$

C. 
$$NH_3 > NF_3 > CH_4 > H_2O$$

$$\mathsf{D}.\,H_2O>NH_3>NF_3>CH_4$$

### **Answer: A**



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115. The correct order of S-O bond length is:

A. 
$$SO_3^{2-} > SO_4^{2-} > SO_3 > SO_2$$

B. 
$$SO_3^{2-} > SO_4^{2-} > SO_2 > SO_3$$

$$\mathsf{C.}\ SO_4^{2-} > SO_3^{2-} > SO_2 > SO_3$$

D. 
$$SO_4^{2-} > SO_3^{2-} > SO_3 > SO_2$$

## **Answer: B**



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- 116. What is not true about rasonance?
  - A. The resonating structures are hypothetical
  - B. The unpaired electrons in various resonating structures are same
  - C. Hybrid structure is most energetic
  - D. Hybrid structure is least energetic

# **Answer: C**



**117.** Which of the following conditions is not correct for resonating structures?

A. The contributing structures must have the same number of unpaired electrons

- B. The contributing structures should have almost similar energies
- C. The contributing structures should be so written that unlike charges reside on atoms that are far apart
- D. The positive charge should be present on the electropositive element and the negative charge on the electronegative element

# **Answer: C**



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118. A molecule may be represented by three structures having energies

 $E_1,\,E_2\,\,{
m and}\,\,E_3$ , respetively. The energies of these structures follow the

order  $E_2 < E_2 < E_1$ , respectively. If the experimental bond energy of the molecule is  $E_0$ , the resonance energy is :

A. 
$$(E_1+E_2+E_3)-E_0$$

B. 
$$E_0-E_3$$

C. 
$$E_0-E_1$$

D. 
$$E_0-E_2$$

# **Answer: B**



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# 119. A:tetracyanomethane B:carbondioxide C:Benzene C:1,3-But-di-ene

Ratio of  $\sigma$  and  $\pi$  bonds is in order:

A. 
$$A=B < C < D$$

$$\operatorname{B.} A = B < D < C$$

$$\mathsf{C.}\, A = B = C = D$$

$$\mathsf{D}.\, C < C < A < B$$

# Answer: A



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# 120. In a compound

The number of sigma and pi bonds respectively are  $:\stackrel{|}{C}=\stackrel{|}{C}_{NC}$ 

 $NC = M(CO)_3$ 

B. 19, 5

C. 13, 11

D.7, 3

# **Answer: A**



<b>121.</b> Which of the following does not contain a coordinate bond?							
A. $H_3O^+$							
B. $BF_4^-$							
C. $HF_2^{-}$							
D. $NH_4^{+}$							
Answer: C							
Watch Video Solution							
122. Which of the following molecules does not have co-ordinate bond?							
<b>122.</b> Which of the following molecules does not have co-ordinate bond? $ {\rm A.} \ PH_4^{\ +} $							
A. $P{H_4}^+$							
A. $P{H_4}^+$ B. $NO_2$							

# Answer: D



**123.** Which of the following halides is inert towards hydrolysis at room temperature?

- A.  $SiCl_4$
- B.  $PCl_3$
- $\mathsf{C}.\,NCl_3$
- D.  $NF_3$

# **Answer: D**



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124. Amongst the following trihalides, which one is least basic?

A.  $NF_3$ 

B.  $NCl_3$ 

 $\mathsf{C}.\,NBr_3$ 

D.  $NI_3$ 

# **Answer: A**



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A. 
$$Ca^{2+} < Ba^{2+} < Sr^{2+}$$

C. 
$$Ge^{2+} < Sn^{2+} < Pb^{2+}$$

B.  $Pb^{2+} < Ge^{2+} < Sn^{2+}$ 

D. 
$$Cu^{2+,\,tAu^{2+}\,<\,Ag^{2+}}$$



**Answer: C** 

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125. Increasing order of stability of the +2 oxidation of the ions?

**126.** The number of three centre two electron bonds in a molecule of diborane is :

**A.** 0

B. 2

 $\mathsf{C.}\ 4$ 

D. 6

## **Answer: B**



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**127.** In which of the following compounds octet is complete and incomplete for all atoms :

A. 
$$Al_2Cl_6$$
  $Al_2(CH_3)_6$   $AlF_3$  Dimer of  $BeCl_2$  Dimer of  $BeH_2$   $IC$   $IC$   $IC$   $C$   $C$   $C$   $C$   $C$   $B.  $Al_2Cl_6$   $Al_2(CH_3)_6$   $AlF_3$  Dimer of  $BeCl_2$  Dimer of  $BeH_2$   $C$   $IC$   $IC$   $C$   $IC$$ 

 $Al_2Cl_6$   $Al_2(CH_3)_6$   $AlF_3$  Dimer of  $BeCl_2$  Dimer of  $BeH_2$ CICIC $Al_2Cl_6$   $Al_2(CH_3)_6$   $AlF_3$  Dimer of  $BeCl_2$  Dimer of  $BeH_2$ IC CIC - ICIC**Answer: C** 

128. In Which of the following metal to metal bond is present?



A. Cupric chloride

B. Stannous chloride

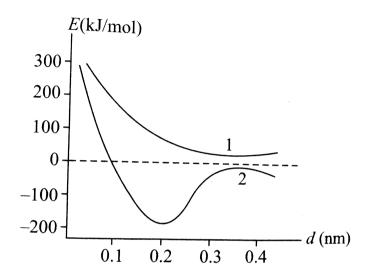
C. Mercurous chloride

D. Mercutic chloride



**Answer: C** 

**1.** Consider the given figure showing the formation of  $H_2^+$  ion depending on intermuclear distance versus potential energy of the system.



- A. Curve-1 represents the most stable state of the system for  $H_2^{\,+}$
- B. Curve-2 represents the most stable state of the system for  $H_2^{\,+}$
- C. Curve-1 indicates that the molecular hydrogen ion is formed
- D. Curve-2 represents the energy level of the antibonding region

# **Answer: B**



2. Molecule having non-pola as well as polar bonds but the molecule as a whole is polar:

A.  $S_2F_2$ 

B.  $N_2O_4$ 

 $\mathsf{C}.\,Si_2H_6$ 

D.  $I_2Cl_6$ 

# Answer: A



- **3.** Choose the correct statement regarding  $SeOCl_2$  molecule :
  - A. It does not contain plane of symmtry
    - B. Cl-S-Cl` bond angle is greater than 'Cl-Se-O' bond angle
    - C. Lone pair has greater than 33.% s-character
    - D. Central atom used one d-ordital in bonding

# Answer: D



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- **4.** In which species X-O bond order is 1.5 and contains  $p\pi-d\pi$  bond (s)
  - A.  $IO_2F_2$
  - B.  $HCOO^-$
  - $\mathsf{C.}\,SO_3^{2\,-}$
  - D.  $XeO_2F_2$

# Answer: A



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5. Which of the following species has polar and non-polar bonds but molecule as a whole is non-plonar?

A. 
$$S_2O_3^{2-}$$

$$\mathrm{B.}\left(SCN\right)_{2}$$

$$\mathsf{C.}\,Be_2Cl_4$$

# D. $Si_2H_6$

#### Answer: D



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# **6.** The incorrect statement (s) regarding $\overline{\phantom{a}}(C)X_3$

A. If electronegativity of surrounding element X is less than 2.5, then central carbon atom used almost 33% s-character in their hybrid bonding orbital

B. If electronegativity of surrounding element X is less than 2.5, then central carbon atom used almost 25% s-character in their hybrid bonding orbital

- C. If X is F, then species should be polar and pyramidal
- D. If X is H, then species should be polar and planar

### **Answer: D**



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- 7. Consider following compounds,
- (i)  $H_3X HCS$  (ii)  $H_3Y NCS$  (iii)  $(H_3Y)_2O$  (iv)  $(H_2X)_2O$

The incorrect statement regarding given compounds is:

- A. If Y is carbon in compounds (ii) and (iii), then both are bent
- B. If X is silicon in compounds (i) and (iv), then both are linear
- C. If X is carbon and Y is silicon, the compound (i) is more basic than

compound (ii)

D. If X is silicon and Y is carbon, then X-O-X bond angle compounds (iii)

is greater than Y-O-Y bond angle in compounds (iv)

## **Answer: B**



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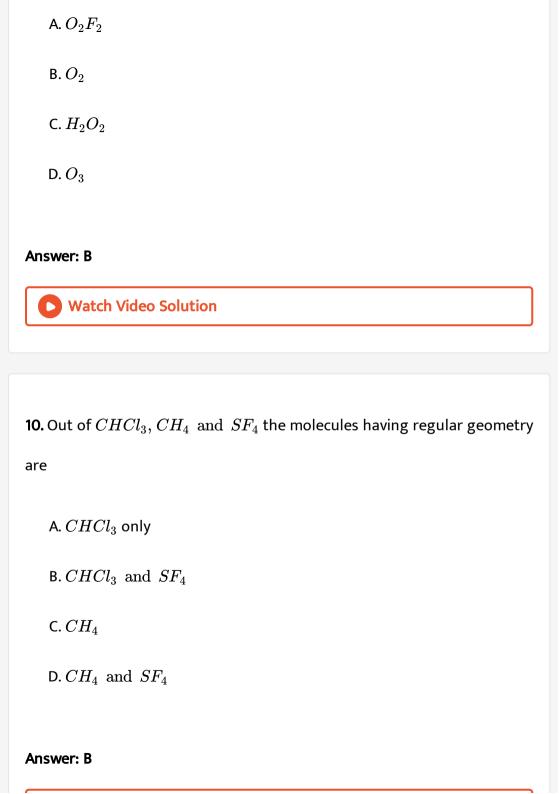
- **8.** The incorrect statement about carbene  $(CH_2)$  is :
  - A. In singlet carbene, carbon is  $sp^2$ -hybridized whereas in triplet carbene, carbon is sp-hybridized
  - B. Triplet carbene is less stable than singlet carbene
  - C. Stability order of singlet halocarbene is :CHF>CHCl>CHBr
  - D. None of the above

## **Answer: B**



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9. The lowest O-O bond length in the following molecule is:



11.	When	iodine	is dissolved	in aqueous	potassium	iodine,	the shape	of
th	e speci	es form	ed is :					

A. linear

B. angular

C. triangular

D. see-saw

# **Answer: A**



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12. Which of the following set of species have planar structure?

A.  $I_3^-, CH_3, Cl_3^-, SiF_6^{2-}$ 

 $\operatorname{B.}I_3^{\,+},ICl_4^{\,-},Al_2Cl_6,TeCl_4$ 

C.  $SCl_2, N_2O_5, SF_4$ 

D.  $I_{2}Cl_{6}, XeF_{2}, BrF_{4}^{-}, XeF_{5}^{-}$ 

# **Answer: D**



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13. Which of the following compounds have the same no. of lone pairs with their central atom?

(I)  $XeF_5^-$  (II)  $BrF_3$  (III)  $XeF_2$  (IV) $H_3S^+$  (V) Triplet methylene

A. (IV) and (V)

B. (I) and (III)

C. (I) and (II)

D. II, IV and V

# **Answer: C**



14. Given the correct order of initials T or F for following statements. Use

T if statements is true and F it it is false:

 $(I)(CH_3)_2P(CF_3)_3$  is non-palar and  $(CH_3)_3P(CF_3)_2$  is polar molecule

(II) $CH_3\widehat{P}CH_3$  bond angle are equal in  $(CH_3)_3P(CF_3)_2$  molecule

(III)  $PF_3$  will be more soluble in polar solvent than  $SiF_4$ 

A. TTF

B. FFT

C. FFF

D. FTT

## **Answer: D**



- **15.** The correct sequence of polarity of the following molecule
- (1) Benzene (2)Inorganic Benzene (3)  $PCl_3F_2$  (4) $PCl_2F_3$

16. Which among the followiing molecules is not perfect flat?

- A.  $B_3N_3H_6$
- $\mathsf{B.}\, C_3 N_3 (NH_2)_3$
- $\mathsf{C}.\,SO_3$
- D.  $C_3N_3(N_3)_3$

### **Answer: D**



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17. Which of the following structure(s) is /are non-planar?

- A.  $Na_3B_3O_6$
- B.  $I_2Cl_6$
- C. Sheet silicate

D. Inorganic graphite layer

**Answer: C** 



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- **18.** Nodal planes of  $\pi$ -bonds in  $CH_2=C=C=CH_2$  are located in,
  - A. All are in molecular plane
  - B. Two in molecular plane and one in a plane perpendicular to molecular plane which contains C-C sigma-bond
  - C. On is molecular plane and two in plane perpendicular to molecular plane which contains C-C sigma-bonds
  - D. Two in molecular plane and one in a perpendicular to molecular plane which bisects C-C sigma-bonds at right angle.

## Answer: B



**19.** Which of the following have X-O-X linkage?

(where X is central atom):

(i) $Cr_2O_7^{2-}$  (ii) $S_2O_3^{2-}$  (iii) pyrosilicate (iv) Hyponitrous acid

- A. (i) and (iii)
- B. (iii) and (iv)
- C. (i), (iii) and (iv)
- D. (i) and (ii)

# **Answer: A**



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**20.** Oxidation state of 'S' in peroxodisulphuric acid and sodium tetrathionate:

A. +6, +5, 0

$$B. +6, +6, +6$$

$$C. +6, +4, +2$$

$$D. +6, +2, 0$$

# Answer: A



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# **21.** Structure of $Na_2igl[B_4O_5(OH)_4igr]\cdot SH_2O$ contains

A. two triangular and two tetrahedral units

B. three triangular and one tetrahedral units

C. all tetrahedral units

D. all triangular units

# Answer: A



A.  $(CN)_2$ 

B.  $OCN^-$ 

 $\mathsf{C}.\,XeF_2$ 

D.  $XeF_4$ 

#### **Answer: D**



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# 23. Incorrect match is:

## **Electron geometry**

- (a) Tetrahedron
- (b) Trigonal bipyramidal
- (c) Octahedron
  (d) Pentagonal bipyramidal

# Possible molecular shape from respective electron geometry

Bent

Triangular planar Square pyramidal

Pentagonal planar



**24.** Consider the following reactions:

 $MX_4 + X'_2 \rightarrow MX_4X_2'$ 

If atomic number of M is 52 and X and X' are halogens and X' is more electonegative than X. Then choose the correct statement regarding the given information:

A. Both X' atoms occupy axial positions which are formed by overlapping of p and d-orbitals only

B. All M-X bond lengths are identical in both  $MX_4$  and  $MX_4X{^\prime}_2$  compounds

C. Central atom  ${\cal M}$  does not use any valence non-axial set of d-orbitals in hybridization of final product

D. Hybridization of central atom  ${\cal M}$  remains same in both reactant and final product

# **Answer: C**



# 25. Select the Incorrect statements:

	Statement	Shape	Example
(a)	Bond pair has > ?5% p-character	&	HF
(b)	Reduction in axial bond angle is more than that of in equatorial bond angle	×	SF <sub>4</sub>
(c)	Two axial d-orbitals and one non-axial d-orbital are used in hybridization	*	XeF <sub>5</sub>
(d)	Two p-orbitals are used in hybridization	2	SnCl <sub>2</sub>



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**26.** In which of the following species, d-obitals having xz an yz two nodal planes involved in hybridization of central atoms?

A. 
$$IO_2F_2^{\,-}$$

B. 
$$ClF_4^{\,-}$$

$$\mathsf{C}.\,IF_7$$

D. None of these

## **Answer: C**



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**27.** The correct order of increasing s character ( in percentage ) in the hybrid orbitals in below molecules / ions is ( assume all hybrid orbitals are exactly equivalent) :

$$CO_3^{2\,-}$$
  $XeF_4$   $I_3^ NCl_3$   $BeCl_2(g)$ 

A. 
$$II < III < IV < I < V$$

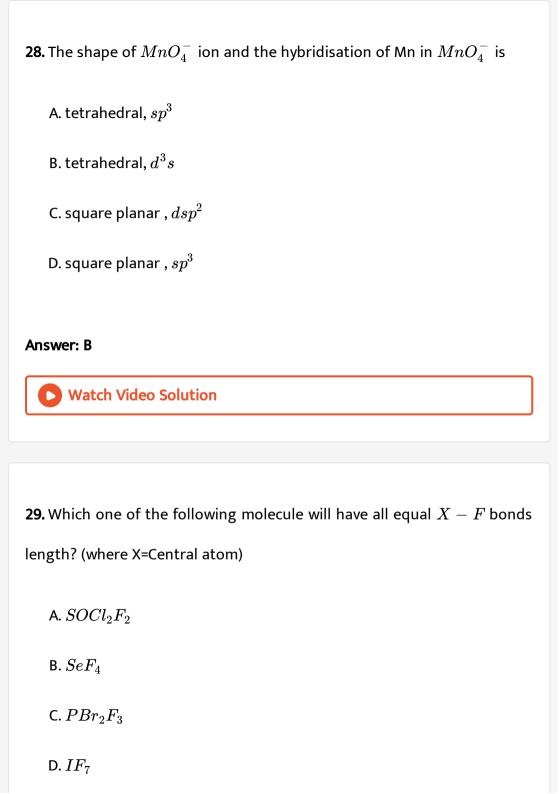
$$\mathsf{B}.\,II < IV < III < V < I$$

$$\mathsf{C}.\,III < II < I < V < IV$$

$$\mathsf{D}.\,II < IV < III < I < V$$

# **Answer: A**







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**30.** Consider the following information (F=F or Cl)

Molecule	P-X(axial) bond length	P-X(Equitorial) bond length
PF <sub>5</sub>	а	ь
PF <sub>4</sub> CH <sub>3</sub>	A settle of County of the or	of someontal and Lorenza a una
PF <sub>3</sub> (CH <sub>3</sub> ) <sub>2</sub>	the standage of the vi	more on the Mark No.
PCl <sub>5</sub>	g	h

According to given information choose the incorrect order of bond length:

A. 
$$g > a > d > b$$

$$\operatorname{B.} g > e > f > b$$

$$\mathsf{C}.\, f > d > a > b$$

D. 
$$c > f > d > b$$

**Answer: C** 



**31.** In which of the following cases C-C bond length will be highest?

A. 
$$CH_3-CF_3$$

$$\mathsf{B.}\,FCH_2-CH_2F$$

$$\mathsf{C.}\,F_2CH-CHF_2$$

D. 
$$CF_3-CF_3$$

#### **Answer: B**



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**32.** In which of the following cases C-C bond length will be highest?

(I) In  $N_2F_4$ , d-orbitals are contracted by electronegative flourine atoms,

but d-orbitals contraction is not possible by H-atoms in  $N_2H_4$ 

(II) The N-N bond energy in  $N_2F_4$  is more than N-N bond energy in

 $N_2H_4$ 

(III) The N-N bond length in  $N_2F_4$  is more than that of in  $N_2F_4$ 

(IV) The N-N bond length in  $N_2F_4$  is less than that of in  $N_2H_4$ choose the correct codes: A. I, II and III B. I and III C. II and IV D. II and III **Answer: B** View Text Solution 33. The correct order of equatorial FSF bond angle in the following compound. (I) $SF_4$  (II)  $OSF_4$  (III)  $H_2CSF_4$ A. III > II > I $\mathrm{B.}\,I > III > II$ C.I > II > III

D.II > III > I

Answer: D



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**34.** Incorrect orders of bond angle is:

A. 
$$Ocl_2 > SF_2 > AsH_2 > H_2Se$$

B. 
$$NH_3>PF_3>PH_3>H_2S$$

C. 
$$XeO_4 > ClO_4^- > SO_4^- > CF_4$$

D. 
$$N(SiH_3)_3 > O(SiH_3)_2 > OMe_2$$

Answer: C::D



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**35.** Minimum  ${\cal F}-{\cal S}-{\cal F}$  bond angle present in :

A.  $OF_2 < ClO_2 < H_2O < Cl_2O$ B.  $OF_2 < H_2O < Cl_2O < ClO_2$ C.  $OF_2 < H_2O < ClO_2 < Cl_2O$ D.  $ClO_2 < OF_2 < H_2O < Cl_2O$ Answer: B

A.  $SSF_2$ 

B.  $SF_6$ 

 $\mathsf{C}.\,SF_2$ 

**Answer: D** 

D.  $F_3SSF$ 

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36. The correct order of increasing bond angle is

**37.** The correct order for bond angles is :

A. 
$$NO_2^->NO_2^+>NO_2$$

$${\rm B.}\,NO_2^{\,+}\,>NO_2^{\,+}\,>NO_2$$

$${\sf C.}\ NO_2 > NO_2^+ > NO_2^-$$

D. 
$$NO_2^+ > NO_2 > NO_2^-$$

#### **Answer: D**



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38. Which one is correct for bond angle?

A. 
$$PF_3 > PCl_3$$

$$\operatorname{B.}\mathit{OCl}_2 = \mathit{ClO}_2$$

$$\mathsf{C.}\,NF_3>NH_3$$

D. 
$$PCl_3 > PF_3$$

#### **Answer: D**



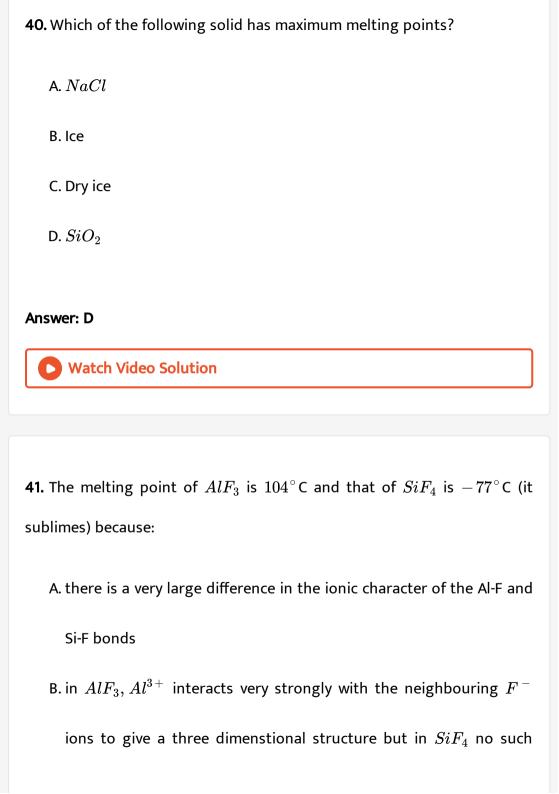
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**39.** In molecules of the type  $AX_2I_n$  (where I represents lone pair and n is its number) there exists a bond between element A and X. The  $\angle XAX$  bond angle

- A. Always decrease if n increases
- B. Always increase if n increases
- C. Will be maximum for n=3
- D. Generally decrease if n decreases

### **Answer: C**





interaction is possible

C. the silicon ion in the tetrahedral  $SiF_4$  is not shielded effectively from the flouride ions whereas in  $AlF_3$ , the  $Al^{3\,+}$  ion is shielded on all sides

D. the attractive forces between the  $SiF_4$  molecules are strong whereas those between the  $AlF_3$  molecules are weak

# **Answer: B**



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**42.** The correct order of boiling point is:

A.  $T_2 < D_2 > H_2$ 

B. n-pentane < neo-pentane

 $\mathsf{C}.\,Xe < Ar < He$ 

D. m-nitrophenol > o-nitrophenol

### **Answer: D**



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**43.** Cis butene dioic acid  $\stackrel{K_{a_1}(\ -H^+\ )}{\longleftrightarrow} X_1^- \stackrel{K_{a_2}(\ -H^+\ )}{\longleftrightarrow} X_2^-$ 

Trans-butene dioic acid 
$$\stackrel{K_{a_1}(\;-H^+\;)}{\longleftarrow} Y_1^- \stackrel{K_{a_2}(\;-H^+\;)}{\longleftarrow} Y_2^-$$

The incorrect statement regarding above information is:

- A.  $X_2^{2\,-}$  species is more basic than  $Y_2^{2\,-}$  species
- B.  $X_1^{\,-}$  species is more basic than  $Y_1^{\,-}$  species
- C.  $K_{a_1}$  is greater than  $K^{\prime}{}_{a_1}$
- D.  $K_{a_2}$  is greater than  $K^{\prime}_{a_2}$

### **Answer: B**



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**44.** Which of the following is not a best representation of the H-bond?

В.

D. None

#### **Answer: C**



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# 45. The H-Bonds in solid HF can be best represented as:

A. 
$$H-F\ldots H-F\ldots H-F$$

C.

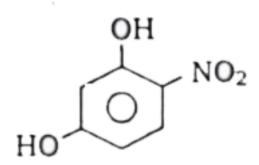
D.

**Answer: C** 



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**46.** The type of molecular forces of attraction present in the following compounds is :



- A. Intermolecular H-bonding
- B. Intramolecular H-bonding
- C. van der waal's force
- D. All of these

### **Answer: D**



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- 47. Which of the following interaction lies in the range of 8-42kH/mol?
  - A.  $H_2$ . .  $H_2O$
  - B. HCl. . HCl
  - C.  $F^- \dots HF$
  - D.  $HCN...NH_3$

#### **Answer: D**



- **48.** The incorrect order is:
  - A. Covalent character :  $PbCl_4 > CaCl_2 > SrCl_2 > BaCl_2$

B. Thermal stability :  $PbF_4 > PbCl_4 > PbBr_4 > PbI_4$ 

C. Melting point : KF > KCl > KBr > KI

D. Boiling point :  $CHCl_3 > CH_3Cl > CCl_4$ 

#### Answer: D



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**49.** If  $CdI_2$  is pink in colour, the  $CdCl_2$  will be ' ' coloured.

A. Yellow

B. Red

C. Blue

D. connot be predicted

## Answer: D



50. The correct solubility order is/are

(I)  $CaCO_3 > SrCO_3 > BaCO_3$  (II) $Li_2CO_3 < Na_2CO_3 > K_2CO_3$ 

 $\text{(III)} K_2 CO_3 < Rb_2 CO_3 < Cs_2 CO_3 \text{ (IV)} Na_2 CO_3 > K_2 CO_3 > Rb_2 CO_3$ 

A. II, IV

B. I, IV

C. II, III, IV

D. I, II, III

### **Answer: D**



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**51.** On heating to  $400-500^{\circ}C$ , relatively unstable hydrides and carbonates decompose. Which of the following will decompose when heated to  $400-500^{\circ}C$ ?

 $(I)LiH\ (II)NaH\ (III)Li_{2}CO_{3}\ (IV)Na_{2}CO_{3}$ 

A. 
$$II, III$$

B. I, II, III

 $\mathsf{C}.\,I,\,III$ 

D. III, IV

#### Answer: A



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**52.** Both  $N(SiH_3)_3$  and  $NH(SiH_3)_2$  compounds have trigonal planar skeleton. Incorrect statement about both compounds is :

A. SiNSi bond angle in  $NH(SiH_3)_2 > SiNSi$  bond angle in

$$N(SiH_3)_3$$

B. N-Si bond length in  $NH(SiH_3)_2>N-Si$  bond length in

$$N(SiH_3)_3$$

C. N-Si bond length in  $NH(SiH_3)_2 < N-Si$  bond length in

$$N(SiH_3)_3ig)$$

 $N(SiH_3)_2$ 

D. Back bonding strength in  $NH(SiH_3)_2>\,\,$  Back bonding strength in

# Answer: B



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**53.** The incorrect statement regarding  $O(SiH_3)_2$  and  $OCl_2$  molecule is/are

A. The strength of back bonding is more in  $O(SiH_3)_2$  molecule than

 $Ocl_2$  molecule

B. Si-O-Si bond angle in  $O(SiH_3)_2$  is greater than Cl-O-Cl bond angle in  $OCl_2$ 

C. The nature of back in both molecules is  $2p_x-3d_\pi$ 

D. Hybridisation of central O-atom in both molecules is same

**Answer: D** 



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**54.** Among following molecule N-Si bond length is shortest:

A.  $N(SiH_3)_3$ 

B.  $NH(SiH_3)_2$ 

C.  $NH_2(SiH_3)$ 

D. All have equal N-Si bond length

**Answer: C** 



**55.** Which of the following molecule has weakest  $(p\pi-d\pi)$  back bonding?

A.  $ClO_2$ 

B.  $N(SiH_3)_3$ 

C.  $SiF_3$ 

D.  $O(SiH_3)_2$ 

### **Answer: A**



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**56.** "hybridization of central atom does not always change due to back bonding". This statement is valid for which of the following compounds?

(i) $\mathbb{C}l_3^-$  (ii) $\mathbb{C}l_2$  (iii) $O(SiH_3)_2$  (iv) $N(SiH_3)_3$ 

A. (i) and (ii)

B. (i) and (iii)

C. (ii)	and	(iii)
D. All		

### **Answer: A**



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**57.** The geometry with respect of the central atom of the following molecules are

$$N(SiH_3)_3, Me_3N, (SiH_3)_3P$$

- A. planar, pyramidal, planar
- B. planar, pyramidal, pyramidal
- C. pyramidal, pyramidal
- D. pyramidal, planar, pyramidal

### **Answer: B**



**58.** Incorrect statement regarding  $BF_3NH_3$  molecule is :

- A. FBF bond angle  $\,< 120^{\circ}$
- B. HNH bond angle  $\,>109^{\circ}\,28^{\,\prime}$
- C. Exhibits intermolecular H-bond
- D. hybridization of N-atom is  $sp^3$

#### **Answer: D**



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**59.** In which of the following molecules  $\mu$ (oberved) is found to be greater than  $\mu$  (theoretical):

- A.  $CHCl_3$
- В. 📝

$$\begin{array}{c} NO_2 \\ NO_2 \\ NO_2 \end{array}$$

## **Answer: D**

C.



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**60.** Among the following, the molecule with the highest dipole moment is

:

A.  $CH_3Cl$ 

B.  $CH_2Cl_2$ 

 $\mathsf{C.}\,CHCl_3$ 

 $\mathsf{D.}\,CCl_4$ 

# Answer: A



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**61.** Which of the following compounds has dipole moment approximately equal to that of chlorobenzene?

- A. o-dichlorobenzene
- B. m-dichlorobenzene
- C. p-dichlorobenzene
- D. p-chloronitrobenzene

#### **Answer: B**



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**62.** Which of the following bonds have lowest bond energy?

B. N-N $\mathsf{C}.H-H$ D.O-O**Answer: D** Watch Video Solution 63. The bond having the minimum bond energy is: A. C-CB.O-O $\mathsf{C}.\,S-S$  $\mathsf{D}.\,P-P$ **Answer: B** Watch Video Solution

A. C-C

**64.** Arrange in increasing order of extent of hydrolysis  $[CCl_4, MgCl_2, AlCl_3, PCl_5, SiCl_4].$ 

A. 
$$CCl_4 < MgCl_2 < AlCl_3 < SiCl_4 < PCl_5$$

$$\operatorname{B.}CCl_4 < AlCl_3 < MgCl_2 < PCl_5 < SiCl_4$$

C. 
$$CCl_4 < SiCl_4 < PCl_5 < AlCl_3 < MgCl_2$$

$$\hbox{D.} \ CCl_4 < PCl_5 < SiCl_4 < AlCl_3 < MgCl_2$$

#### **Answer: A**



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**65.** Inorganic benzene reacts with HCl to form a compounds  $B_3N_3H_9Cl_3$ .

The protonation occurs at:

A. B-atom

B. N-atom

- C. First at B-atom than rearranges into N-atom
- D. first at N-atom then rearranges into B-atom

#### **Answer: B**



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- **66.** Select the correct statements about hydrolysis of  $BCl_3$  and  $NCl_3$ :
  - A.  $NCl_3$  is hydrolysed and gives HOCl but  $BCl_3$  is not hydrolysed
  - B. Both  $NCl_3 \ {
    m and} \ BCl_3$  on hydrolysis gives HCl
  - C.  $NCl_3$  on hydrolysis give HOCl but  $BCl_3$  gives HCl
  - D. Both  $NCl_3$  and  $BCl_3$  on hydrolysis gives HOCl

### **Answer: C**



67. The incorrect statement regarding molecular orbital (s) is:

A. If there is a nodal plane perpendicular to the intermoleculear axis and lying between the nuclei of bonded atoms then corresponding orbitals is antibonding M.O.

B. If a nodal plane lies in the inter-nuclear axis, then corresponing orbitals is (pi) bonding M.O.

C. The  $\sigma$ - bonding molecular orbital does not contain nodal planes containing the internuclear axis

D. The  $\,\delta$ -bonding molecular orbitals possesses three nodal planes containing the internuclear axis

# Answer: D



**68.** Which of the following species absorb maximum energy in its HOMO-LUMO electronic transition?

- A.  $O_2$
- $\mathrm{B.}\,N_2^{\,-}$
- $\mathsf{C}.\,C_2$
- D.  $N_2$

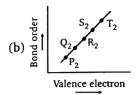
#### **Answer: D**



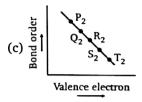
**69.** If P to T are second period p-block elements then which of the following graph show corrrect relation between valence electrons in  $P_2 o T_2$ 

(a) 
$$P_2$$

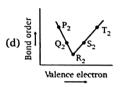
$$Valence electron$$



В.



C.



D.

#### Answer: A



- 70. Which of the following facts given is not correct?
- (I) Bond length order,  $H_2^{\,-}-H_2^{\,+}>H_2$
- (II)  $O_2^+,NO,N_2^-$  have same bond order of  $2rac{1}{2}$
- (II) Bond order can assume any value including zero upto four

(IV)  $NO_3^-$  and  $BO_3^-$  have same order for X-O bond (where X is central atom)

**71.**  $N_2$  and  $O_2$  are converted to monocations  $N_2^+$  and  $O_2^+$  respectively,

A. I, II and III

B. I and IV

C. II and IV

D. I and II

### **Answer: B**



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which is wrong statement:

- A. In  $N_2^+$  , the N-N bond weakens
- B. In  ${\cal O}_2^+$  , the O-O bond order increase
- C. In  $O_2^+$ , the paramagnetism decrease

D.  $N_2^+$  becomes diamagnetic

**Answer: D** 



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72. In which of the following transformations, the bond order has increased and the magnetic behaviour has changed?

A. 
$$C_2^{\,+}\, o C_2$$

B. 
$$NO^- o NO$$

$$\mathsf{C}.\,O_2\to O_2^+$$

D. 
$$N_2 
ightarrow N_2^{\,+}$$

#### **Answer: A**



73. H.O.M.O (Highest Occupied Molecular Orbital ) of CO molecular is:

A. Non-bonding M.O. with slight antiboding character

B. Non-bonding M.O. with slight boding character

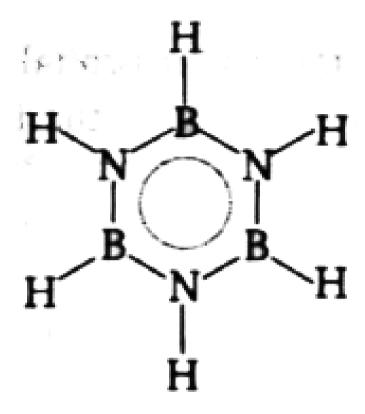
C. Pure non-bonding M.O.

D. None of above

#### **Answer: A**



# **74.** The structure of $B_3N_3H_6$ is as follows:



How may derivations structures of  $B_3N_3H_4X_2$  can be derived from the basic structure, by the replacement of two hydrogen atoms?

- A. 2
- B. 3
- **C**. 4
- D. 5

### **Answer: C**



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# 75. Correctly match is:

- A.  $d_{x^2-y^2}$  atomic orbital -one nodal plane
- B.  $p_y$  atomic orbital -Two nodal planes
- C.  $\sigma_{p_x}$  - $\psi$ (gerade)
- D.  $\Pi_{p_y}$  - $\psi$ (ungerade)

# **Answer: C**



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**76.** Select correct statement (s):

A. Acidic strength of HBr>HCl but reverse is true for their

reducing property

B. Basic strength of  $PH_3>AsH_3$  but reverse is true for their bond angle

C. Dipole moment of  $CH_3Cl>CH_3F$  but reverse is true for their  $H\widehat{C}H$  bond angle

D.  $K_{a_1}$  of fumaric acid is higher than maleic acid but reverse is true for their  $K_{a_2}$ 

#### **Answer: C**



**77.** Molecule having non-polar as well as polar bonds but the molecule as a whole is polar

A.  $S_2F_2$ 

- B.  $N_2O_4$
- $\mathsf{C.}\,Si_2H_6$
- $\operatorname{D.}I_2Cl_6$

#### **Answer: A**



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- **78.** Choose the correct statement regarding  $SeOCl_2$  molecule :
  - A. It does not contain plane of symmtry
  - B. Cl-S-Cl` bond angle is greater than 'Cl-Se-O' bond angle
  - C. Lone pair has greater than 33.% s-character
  - D. Central atom used one d-ordital in bonding

## Answer: D



**79.** In which species, X-O bond order is 1.5 and contains  $p\pi-d\pi$  bond(s)

A.  $IO_2F_2$ 

 $\mathsf{B.}\,HCOO^-$ 

 $\mathsf{C.}\,SO_3^{2\,-}$ 

D.  $XeO_2F_2$ 

## Answer: A



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**80.** Which of the following species has polar and non-polar bonds but molecule as a whole is non-plonar?

A. 
$$S_2O_3^{2\,-}$$

 $\mathrm{B.}\left(SCN\right)_{2}$ 

 $\mathsf{C.}\,Be_2Cl_4$ 

D.  $Si_2H_6$ 

#### **Answer: D**



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**81.** The incorrect statement (s) regarding  $\overline{\phantom{a}}(C)X_3$ 

A. If electronegativity of surrounding element X is less than 2.5, then central carbon atom used almost 33% s-character in their hybrid bonding orbital

- B. If electronegativity of surrounding element X is less than 2.5, then central carbon atom used almost 25% s-character in their hybrid bonding orbital
- C. If X is F, then species should be polar and pyramidal
- D. If X is H, then species should be polar and planar

## **Answer: D**



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82. Consider following compounds,

(i) 
$$H_3X - HCS$$
 (ii)  $H_3Y - NCS$  (iii)  $(H_3Y)_2O$  (iv)  $(H_2X)_2O$ 

The incorrect statement regarding given compounds is:

A. If Y is carbon in compounds (ii) and (iv), then both are bent

B. If X is silicon in compounds (i) and (iii), then both are linear

C. If X is carbon and Y is silicon, the compound (i) is more basic than

compound (ii)

D. If X is silicon and Y is carbon, then X-O-X bond angle compounds (iii)

is greater than Y-O-Y bond angle in compounds (iv)

#### **Answer: B**



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**83.** The incorrect statement about carbene  $(CH_2)$  is :

A. In singler carbene, carbon is  $sp^2$ -hybridized whereas in triplet

carbene, carbon is sp-hybridized

B. Triplet carbene is less stable than singlet carbene

84. The lowest O-O bond length in the following molecule is:

C. Stability order of singlet halocarbenes is

D. None of the above

#### **Answer: B**



- A.  $O_2F_2$
- $B.O_2$
- $\mathsf{C}.\,H_2O_2$
- $D.O_3$

### **Answer: B**



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**85.** Out of  $CHCl_3,\,CH_4\,$  and  $SF_4\,$  the molecules do not having regular geometry are:

A.  $CHCl_3$  only

 $B. CHCl_3$  and  $SF_4$ 

 $\mathsf{C}.\,CH_4$ 

D.  $CH_4$  and  $SF_4$ 

### **Answer: B**



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**86.** When iodine is dissolved in aqueous potassium iodide, the shape of the species formed is :

B. angular C. triangular

A. linear

D. see-saw

# **Answer: A**



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A. 
$$I_3^-, CH_3, Cl_3^-, SiF_6^{2-}$$

 $\mathsf{B}.\ I_3^{+},\ ICl_4^{-},\ Al_2Cl_6,\ TeCl_4$ 

87. Which of the following set of species have planar structure?

C.  $SCl_2, N_2O_5, SF_4$ 

 ${\rm D.}\, I_{2}Cl_{6}, XeF_{2}, BrF_{4}^{\,-}, XeF_{5}^{\,-}$ 

### Answer: D



**88.** Which of the following compounds have the same no. of lone pairs with their central atom?

- (I)  $XeF_5^-$  (II)  $BrF_3$  (III)  $XeF_2$  (IV) $H_3S^+$  (V) Triple methylene
  - A. (IV) and (V)
  - B. (I) and (III)
  - C. (I) and (II)
  - D. II, IV and V

### Answer: C

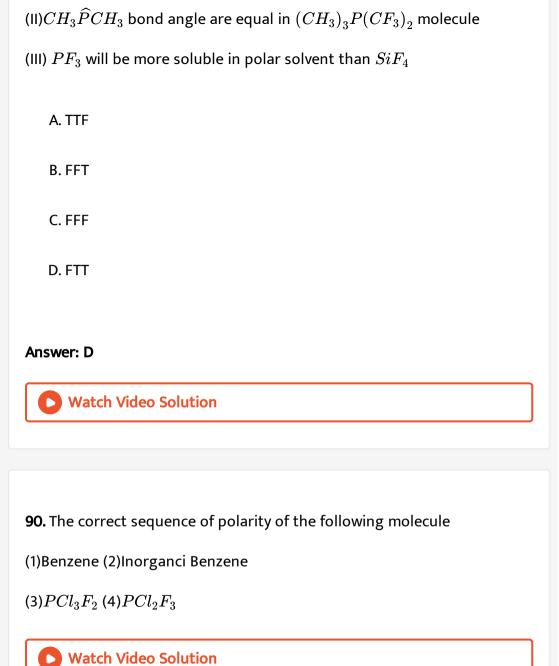


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**89.** Given the correct order of initials T or F for following statements. Use

T if statements is true and F it it is false:

(I) $(CH_3)_2P(CF_3)_3$  is non-palar and  $(CH_3)_3P(CF_3)_2$  is polar molecule



**91.** Which among the followiing molecules is not perfect flat?

A.  $B_3N_3H_6$ 

B.  $C_3N_3(NH_2)_3$ 

 $\mathsf{C}.\,SO_3$ 

D.  $C_3N_3(N_3)_3$ 

### **Answer: B**



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**92.** Which of the following structure(s) is /are non-planar?

A.  $Na_3B_3O_6$ 

 $\operatorname{B.}I_{2}Cl_{6}$ 

C. Sheet silicate

D. Inorganic graphite layer

#### **Answer: C**



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- **93.** Nodal planes of  $\pi$ -bonds in  $CH_2=C=C=CH_2$  are located in,
  - A. All are in molecular plane
  - B. Two in molecular plane and one in a plane perpendicular to molecular plane which contains C-C sigma-bond
  - C. On is molecular plane and two in plane perpendicular to molecular plane which contains C-C sigma-bonds
  - D. Two in molecular plane and one in a perpendicular to molecular plane which bisects C-C sigma-bonds at right angle.

### **Answer: B**



94. Which of the following have X-O-X linkage?

(where X is central atom):

(i) $Cr_2O_7^{2-}$  (ii) $S_2O_3^{2-}$  (iii) pyrosilicate (iv) Hyponitrous acid

- A. (i) and (iii)
- B. (iii) and (iv)
- C. (i), (iii) and (iv)
- D. (i) and (ii)

#### Answer: A



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**95.** Select the correct statements:

A.  $HSO_5^-$  ion has one S-O-H linkage

B. Number of B-O-B linkage in Borax is equal to number of P-O-P

linkage in  $P_4O_{10}$ 

C. Hybridization of both sulphur in  $H_2S_2O_8$  (pyrosulphurous acid ) is

same but oxidation state of both sulphur are different

D. Tetra-polyphosphoric acid has four P-O-P and no P-P linkage

### **Answer: C**



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96. Oxidation state of 'S' in peroxodisulphuric acid and sodium tetrathionate:

A. +6, +5, 0

B. + 6, + 6, + 6

C. +6, +4, +2

D. +6, +2, 0

### Answer: A



**97.** Structure of  $Na_2igl[B_4O_5(OH)_4igr]\cdot 8H_2O$  contains

A. two triangular and two tetrahedral units

B. three triangular and one tetrahedral units

C. all tetrahedral units

D. all triangular units

### **Answer: A**



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98. Which of the following molecular species is not linear?

A.  $(CN)_2$ 

 $\mathsf{B.}\,OCN^{\,-}$ 

 $\mathsf{C.}\,XeF_2$ 

D.  $XeF_2$ 

### Answer: D



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### 99. Incorrect match is:

#### **Electron geometry**

Possible molecular shape from respective electron geometry

(a) Tetrahedron

- (b) Trigonal bipyramidal
- Bent Triangular planar
- (c) Octahedron (d) Pentagonal bipyramidal

Square pyramidal Pentagonal planar



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# **100.** Consider the following reactions:

$$MX_4 + X'_2 \rightarrow MX_4X_2'$$

If atomic number of M is 52 and X and X' are halogens and X' is more electonegative than X. Then choose the correct statement regarding the given information:

A. Both X' atoms occupy axial positions which are formed by overlaping of p and d-orbitals only

- B. All M-X bond lengths are identical in both  $MX_4$  and  $MX_4X{^\prime}_2$  compounds
- C. Central atom 'M' does not use anyone valence non-axial set of dorbitals in hybridization of final product
- D. Hybridization of central atom 'M' remains same in both reactant and final product

### **Answer: C**



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### 101. Select the correct statements:

	Statement	Shape	Example
(a)	Bond pair has > 75% p-character	&	HF
(b)	Reduction in axial bond angle is more than that of in equatorial bond angle	×	SF <sub>4</sub>
(c)	Two axial d-orbitals and one non-axial d-orbital are used in hybridization	*	XeF <sub>5</sub>
(d)	Two p-orbitals are used in hybridization	2	SnCl <sub>2</sub>

**102.** In which of the following species, d-obitals having xz an yz two nodal planes involved in hybridization of central atoms?

- A.  $IO_2F_2^{\,-}$
- B.  $ClF_4^{\;-}$
- $\mathsf{C}.\,IF_7$
- D. None of these

### **Answer: C**



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103. The correct order of increasing s-character (in percentage) in the hybrid orbitals of following molecules/ions is : (I) $CO_3^{2-}$  (II) $XeF_4$  (III)  $I_3^-$  (IV)  $NCl_3$  (V)  $BeCl_2$ 

A. II < III < IV < I < V

 $\mathsf{B}.\,II < IV < III < V < I$ 

 $\mathsf{C}.\,III < II < I < V < IV$ 

 $\mathsf{D}.\,II < IV < III < I < V$ 

### Answer: A



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# **104.** The shapes of $MnO_4^-$ ion and the hybridization of Mn in $MnO_4^-$ is :

- A. tetrahedral,  $sp^3$
- B. tetrahedral,  $d^3s$
- C. square planar ,  $dsp^2$
- D. square planar ,  $sp^3$

## **Answer: B**



**105.** Which one of the following molecule will have all equal X-F bonds

length? (where X=Central atom)

- A.  $SOCl_2F_2$
- B.  $SeF_4$
- C.  $PBr_2F_3$
- D.  $IF_7$

### **Answer: A**



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106. Consider the following information (F=F or Cl)

Molecule	P-X(axial) bond length	P-X(Equitorial) bond length
PF <sub>5</sub>	а	ь
PF <sub>4</sub> CH <sub>3</sub>	A settle of County of the or	of someontal and Lorenza and the
PF <sub>3</sub> (CH <sub>3</sub> ) <sub>2</sub>	ti smalmdddien an fine y	men en la fill "Villa"
PCl <sub>5</sub>	g	h

According to given information choose the incorrect order of bond length:

A. g > a > d > b

B. g > e > f > b

**Answer: C** 

C. f > d > a > b

D. c > f > d > b

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A.  $CH_3 - CF_3$ 

B.  $FCH_2 - CH_2F$ 

C.  $F_2CH - CHF_2$ 

D.  $CF_3 - CF_3$ 

**107.** In which of the following cases C-C bond length will be highest?





Answer: B

**108.** In which of the following cases C-C bond length will be highest?

- (I) In  $N_2F_4$ , d-orbitals are contracted by electronegative flourine atoms, but d-orbitals contraction is not possible by H-atoms in  $N_2H_4$
- (II) The N-N bond energy in  $N_2F_4$  is more than N-N bond energy in  $N_2H_4$
- (III) The N-N bond length in  $N_2F_4$  is more than that of in  $N_2F_4$
- (IV) The N-N bond length in  $N_2F_4$  is less than that of in  $N_2H_4$  choose the correct codes:

A. I, II and III

B. I and III

C. II and IV

D. II and III

### Answer: B



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**109.** The correct order of equatorial FSF bond angle in the following compound.

(I) $SF_4$  (II)  $OSF_4$  (III)  $H_2CSF_4$ 

A. 
$$III > II > I$$

### **Answer: D**



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# 110. Incorrect orders of bond angle is:

A. 
$$Ocl_2 > SF_2 > AsH_2 > H_2Se$$

B. 
$$NH_3>PF_3>PH_3>H_2S$$

C. 
$$XeO_4 > ClO_4^- > SO_4^- > CF_4$$

D. 
$$N(SiH_3)_3 > O(SiH_3)_2 > OMe_2$$

# Answer: C::D



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# 111. Minimum ${\cal F}-{\cal S}-{\cal F}$ bond angle present in :

A.  $SSF_2$ 

B.  $SF_6$ 

 $\mathsf{C}.\,SF_2$ 

D.  $F_3SSF$ 

# **Answer: D**



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112. The correct order of increasing bond angle is

A. 
$$OF_2 < ClO_2 < H_2O < Cl_2O$$

$$\operatorname{B.}OF_2 < H_2O < Cl_2O < ClO_2$$

$$\mathsf{C.}\,OF_2 < H_2O < ClO_2 < Cl_2O$$

$$\mathrm{D.}\,ClO_2 < OF_2 < H_2O < Cl_2O$$

#### **Answer: B**



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113. Bond energies can be obtained by using the following relation:

$$\Delta H(reaction) = \sum$$
 Bond energy of bonds, broken in the reactants

 $-\sum$  Bond energy fo bonds, formed in the products

Bond enegry depends on three factors:

- a. Greater is the bond length, lesser is the bond enegry.
- b. Bond energy increases with the bond multiplicity.
- c. Bond enegry increases with electronegativity difference between the bonding atoms.

Arrange  $N-H,\,O-H,\,$  and F-H bonds in teh decreasing order of bond enegry.

A. 
$$NO_2^- > NO_2^+ > NO_2^-$$

$$\mathsf{B}.\mathit{NO}_2^+ > \mathit{NO}_2^+ > \mathit{NO}_2$$

$$\mathsf{C.}\,NO_2>NO_2^+>NO_2^-$$

D. 
$$NO_2^+>NO_2>NO_2^-$$

### **Answer: D**



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# 114. Which one is correct for bond angle?

A. 
$$PF_3 > PCl_3$$

$$\operatorname{B.}\mathit{OCl}_2 = \mathit{ClO}_2$$

C. 
$$NF_3>NH_3$$

D. 
$$PCl_3 > PF_3$$

### **Answer: D**



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115. In molecules of the type  $AX_2I_n$  (where I represents lone pair and n is its number) there exists a bond between element A and X. The  $\angle XAX$  bond angle

- A. Always decrease if n increase
- B. Always increase if n increase
- C. Will be maximum for n=3
- D. generally decrease if n decrease

# **Answer: C**



A.	NaCl	

B. Ice

C. Dry ice

D.  $SiO_2$ 

### **Answer: D**



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**117.** The melting points of  $AlF_3$  is  $104^\circ$  and that of  $SiF_4$  is  $-77^\circ$  (it sublimes) because:

A. there is a very large difference in the ionic character of the Al-F and

Si-F bonds

B. in  $AlF_3,\,Al^{3\,+}$  interacts very strongly with the neighbouring  $F^{\,-}$  ions to given a three dimenstional structure but in  $SiF_4$  no such interaction is possible

C. the silicon ion in the tetrahedral  $SiF_4$  molecules is not shielded effectively from the flouride ions whereas in  $AlF_3$ , the  $Al^{3\,+}$  ion is

shielded on all sides

D. the attractive forces between the  $SiF_4$  molecules are strong whereas those between the  $AlF_3$  molecules are weak

### Answer: B



# 118. The correct order of boiling point is:

A. 
$$T_2 < D_2 > H_2$$

 ${\bf B.\,n\text{-}pentane}\,<\,{\bf neo\text{-}pentane}$ 

 $\mathsf{C}.\,Xe < Ar < He$ 

D. m-nitrophenol > o-nitrophenol

# **Answer: D**

**119.** Cis butene dioic acid  $\stackrel{K_{a_1}(\;-H^{\,+\;})}{\longleftarrow} X_1^- \stackrel{K_{a_2}(\;-H^{\,+\;})}{\longleftarrow} X_2^-$ 

Trans-butene dioic acid 
$$\stackrel{K_{a_1}(\;-H^{\,+\;})}{\longleftarrow}Y_1^{\,-\;}\stackrel{K_{a_2}(\;-H^{\,+\;})}{\longleftarrow}Y_2^{\,-\;}$$

The incorrect statement regarding above information is:

- A.  $X_2^{2\,-}$  species is more basic than  $Y_2^{2\,-}$  species
- B.  $X_1^{\,-}$  species is more basic than  $Y_1^{\,-}$  species
- C.  $K_{a_1}$  is greater than  $K^{\prime}{}_{a_1}$
- D.  $K_{a_2}$  is greater than  $K^{\prime}{}_{a_2}$

### **Answer: B**



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120. Which of the following is not a best representation of the H-bond?

В.

D. None

### **Answer: C**



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# 121. The H-bonds in solid HF can be best represented as:

A. 
$$H-F\ldots H-F\ldots H-F$$

C.

D.

# **Answer: C**



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# 122. Major product of the following reactions is

- A. Intermolecular H-bonding
- B. Intramolecular H-bonding
- C. van der waal's force
- D. All of these

# Answer: D



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123. Which of the following interaction lies in the range of 8-42kJ/mol?

A.  $H_2.$  .  $H_2O$ 

B. HCl. . HCl

C.  $F^- \dots HF$ 

D.  $HCN...NH_3$ 

**Answer: D** 



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**124.** Assertion (A): The presence of  $CO_2$  in the air accelerates corrosion.

Reason (R):  $CO_2$  is a poisonous gas.

A. Yellow

B. Red

C. Blue

D. connot be predicted

### Answer: D



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# 125. The correct solubility order is/are

(I)  $CaCO_3 > SrCO_3 > BaCO_3$  (II) $Li_2CO_3 < Na_2CO_3 > K_2CO_3$ 

 $(III)K_{2}CO_{3} < Rb_{2}CO_{3} < Cs_{2}CO_{3} \ (IV)Na_{2}CO_{3} > K_{2}CO_{3} > Rb_{2}CO_{3}$ 

A. II, IV

B. I, IV

C. II, III, IV

D. I, II, III

### Answer: D



**126.** On heating to  $400-500^{\circ}C$ , relatively unstable hydrides and carbonates decompose. Which of the following will decompose when heated to  $400-500^{\circ}C$ ?

 $(I)LiH(II)NaH(III)Li_2CO_3(IV)Na_2CO_3$ 

A. II, III

B. I, II, III

 $\mathsf{C}.\,I,\,III$ 

D. III, IV

Answer: A



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**127.** Both  $N(SiH_3)_3$  and  $NH(SiH_3)_2$  compounds have trigonal planar skeleton. Incorrect statement about both compounds is :

A. SiNSi bond angle in  $NH(SiH_3)_2 > SiNSi$  bond angle in  $N(SiH_3)_3$ )

B.N-Si bond length in  $NH(SiH_3)_2>N-Si$  bond length in  $N(SiH_3)_3ig)$ 

C. N-Si bond length in  $NH(SiH_3)_2 < N-Si$  bond length in  $N(SiH_3)_3ig)$ 

D. Back bonding strength in  $NH(SiH_3)_2>\,$  Back bonding strength in  $N(SiH_3)_3ig)$ 

# **Answer: B**



**128.** The incorrect statement regarding  $O(SiH_3)_2$  and  $OCl_2$  molecule is/are

A. The strength of back bonding is more in  $O(SiH_3)_2$  molecule than

 $Ocl_2$  molecule

B. Si-O-Si bond angle in  $O(SiH_3)_2$  is greater than Cl-O-Cl

bond angle in  $OCl_2$ 

C. The nature of back in both molecules is  $2p_x-3d_\pi$ 

D. Hybridisation of central O-atom in both molecules is same

#### **Answer: D**



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**129.** Among following molecule N-Si bond length is shortest:

A.  $N(SiH_3)_3$ 

B.  $NH(SiH_3)_2$ 

 $\mathsf{C.}\ NH_2(SiH_3$ 

D. All have equal N-Si bond length

### Answer: C



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**130.** Which of the following molecule has weakest  $(p\pi-d\pi)$  back bonding?

- A.  $OCl_2$
- B.  $N(SiH_3)_3$
- C.  $SiF_3$
- D.  $O(SiH_3)_2$

### **Answer: A**



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**131.** The geometry with respect to the central atom of the following molecules are :

 $N(SiH_3)_3: Me_3N: (SiH_3)_3P$ 

A. planar, pyramidal, planar

B. planar, pyramidal, pyramidal

C. pyramidal, pyramidal

D. pyramidal, planar, pyramidal

### **Answer: B**



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A. FBF bond angle  $\,< 120^{\circ}$ 

C. Exhibits intermolecular H-bond

B. HNH bond angle  $\,>109^{\circ}\,28^{\,\prime}$ 

**132.** Incorrect statement regarding  $BF_3NH_3$  molecule is :

D. hybridization of N-atom is  $sp^3$ 

# Answer: D

**133.** In which of the following molecules  $\mu$ (oberved) is found to be greater than  $\mu$  (theoretical):

A. 
$$CHCl_3$$

В.

$$(c) \bigcup^{NO_2} NO_2$$

# Answer: D

D.



134. Among the following, the molecule with the highest dipole moment is:

A.  $CH_3Cl$ 

B.  $CH_2Cl_2$ 

 $C. CHCl_3$ 

D.  $CCl_4$ 

# Answer: A



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135. Which of the following is wheat fruit?

A. 0-dichlorobenzene

B. m-dichlorobenzene

C. p-dichlorobenzene

D. p-chloronitrobenzene

### **Answer: B**



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**136.** Which of the following is wheat fruit?

A. 
$$C-C$$

B. N-N

 $\mathsf{C}.H-H$ 

D.O-O

# **Answer: D**



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 $\textbf{137.} \ \mathsf{The} \ \mathsf{bond} \ \mathsf{having} \ \mathsf{the} \ \mathsf{minimum} \ \mathsf{bond} \ \mathsf{energy} \ \mathsf{is} :$ 

A. C-C

$$B.O-O$$

 $\mathsf{C}.\,S-S$ 

D.P-P

#### **Answer: B**



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# 138. The correct increasing order of extent of hydrolysis is

A. 
$$CCl_4 < MgCl_2 < AlCl_3 < SiCl_4 < PCl_5$$

$$\operatorname{B.}CCl_4 < AlCl_3 < MgCl_2 < PCl_5 < SiCl_4$$

$$\mathsf{C.}\,CCl_4 < SiCl_4 < PCl_5 < AlCl_3 < MgCl_2$$

D. 
$$CCl_4 < PCl_5 < SiCl_4 < AlCl_3 < MgCl_2$$

#### **Answer: A**



**139.** Inorganic benzene reacts with HCl to form a compounds  $B_3N_3H_9Cl_3$ 

. The protonation occurs at:

A. B-atom

B. N-atom

C. First at B-atom than rearranges into N-atom

D. first at N-atom then rearranges into B-atom

#### **Answer: B**



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**140.** Select the correct statements about hydrolysis of  $BCl_3$  and  $NCl_3$ :

A.  $NCl_3$  is hydrolysis and gives HOCl but  $BCl_3$  is not hydrolysed

B. Both  $NCl_3$  and  $BCl_3$  on hydrolysis gives HCl

C.  $NCl_3$  on hydrolysis give HOCl but  $BCl_3$  gives HCl

D. Both  $NCl_3$  and  $BCl_3$  on hydrolysis gives HOCl

#### **Answer: C**



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141. The incorrect statement regarding molecular orbital (s) is:

- A. If there is a nodal plane perpendicular to the intermoleculear axis and lying between the nuclei of bonded atoms then corresponding orbitals is antibonding M.O.
- B. If a nodal plane lies in the inter-nuclear axis, then corresponing orbitals is (pi) bonding M.O.
- C. The  $\sigma$ -bonding molecular orbital does not contains the internuclear axis
- D. The  $\,\delta$ -bonding molecular orbitals possesses three nodal planes containing the internuclear axis

Answer: D

142. Which of the following species absorb maximum energy in its

HOMO-LUMO electronic transition?

A.  $O_2$ 

B.  $N_2^-$ 

 $\mathsf{C}.\,C_2$ 

D.  $N_2$ 

#### Answer: D



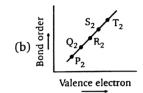
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**143.** If P to T are second period p-block elements then which of the following graph show corrrect relation between valence electrons in  $P_2 o T_2$ 

(a) 
$$\stackrel{\text{by op go}}{\text{go}} = Q_2$$

$$\begin{array}{c} R_2 \\ P_2 \end{array}$$
Valence electron

A.



В.

(c) 
$$\frac{1}{2}$$
  $\frac{1}{2}$   $\frac{1}{2}$   $\frac{1}{2}$   $\frac{1}{2}$   $\frac{1}{2}$  Valence electron

C.

(d) 
$$\stackrel{\text{b}}{\underset{\text{g}}{\text{g}}} \stackrel{\text{b}}{\underset{\text{Q}_2}{\text{Valence electron}}} V_2$$

D.

#### **Answer: A**



- 144. Which of the following facts given is not correct?
- (I) Bond length order,  $H_2^{\,-}\,-\,H_2^{\,+}\,>H_2$

(IV) 
$$NO_3^-$$
 and  $BO_3^-$  have same order for  $X-O$  bond (where X is central atom)

(II) Bond order can assume any value including zero upto four

(II)  $O_2^+,NO,N_2^-$  have same bond order of  $2rac{1}{2}$ 

A. I, II and III

B. I and IV

C. II and IV

D. I and II



**Answer: B** 

145.

# \_\_\_\_\_

A. In  $N_2^{\,+}$  , the N-N bond weakens

respectively, which is wrong statement:

B. In  $O_2^+$ , the O-O bond order increase

 $N_2 \ {
m and} \ O_2$  are converted to monocations  $N_2^+ \ {
m and} \ O_2^+$ 

C. In  $O_2^+$ , the paramagnetism decrease

D.  $N_2^+$  becomes diamagnetic

#### **Answer: D**



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**146.** In which of the following transformations, the bond order has increased and the magnetic behaviour has changed?

A. 
$$C_2^{\,+}\,
ightarrow\, C_2$$

B. 
$$NO^- o NO$$

$$\mathsf{C}.\,O_2\to O_2^+$$

D. 
$$N_2 
ightarrow N_2^+$$

#### **Answer: A**



147. H.O.M.O (Highest Occupied Molecular Orbital ) of CO molecular is:

A. Non-bonding M.O. with slight antiboding character

B. Non-bonding M.O. with slight boding character

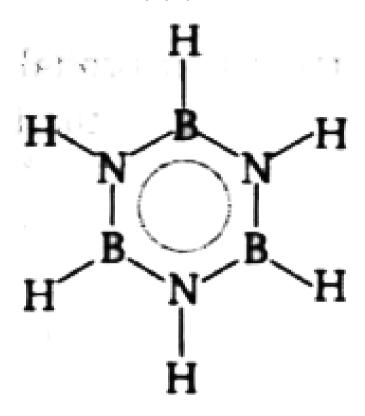
C. Pure non-bonding M.O.

D. None of above

#### **Answer: A**



**148.** The structure of  $B_3N_3H_6$  is as follows:



How may derivations structures of  $B_3N_3H_4X_2$  can be derived from the basic structure, by the replacement of two hydrogen atoms?

- A. 2
- B. 3
- **C**. 4
- D. 5

#### **Answer: C**



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# 149. Correctly match is:

- A.  $d_{x^2-y^2}$  atomic orbital -one nodal plane
- B.  $p_y$  atomic orbital -Two nodal planes
- C.  $\sigma_{p_x}$  - $\psi$ (gerade)
- D.  $P_{p_y}$  - $\psi$ (ungerade)

### **Answer: C**



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**150.** Select correct statement (s):

A. Acidic strength of HBr>HCl but reverse is true for their

reducing property

B. Basic strength of  $PH_3>AsH_3$  but reverse is true for their bond angle

C. Dipole moment of  $CH_3Cl>CH_3F$  but reverse is true for their  $H\widehat{C}H$  bond angle

D.  $K_{a_1}$  of fumaric acid is higher than maleic acid but reverse is true for their  $K_{a_2}$ 

#### **Answer: C**



# Level 3

**1.** Colour of compounds occurs due to phenomenon of polarisation, charge transfer, d-d transition and H.O.M.O.L.U.M.O transition. Most of the

transition metal transition metal complex compound are coloured either due to d-d electrons transition or charge transfer and ionic compounds are coloured due to polarisation of anion. Q. Which of the following is correct about  $KFe^{II}\Big[Fe(CN_6)\Big]$  and  $KFe^{III}\Big[Fe(CN_6)\Big]$  complex compounds?

A. Both are blue coloured compounds because colour arises due to dd electrons transition in Fe cation present outside the complex ion.

B. Both are blue coloured compounds because colour arises due to transfer of electron between  $Fe^{II}$  and  $Fe^{III}$  cation

C. Both are blue coloured compounds because in complexes  $Fe^{II}$ 

D. Complex (I) has blue colour becouse (II) has brown colour

cation shows same d-d transition bond

#### **Answer: B**



**2.** Colour of compounds occurs due to phenomenon of polarisation, charge transfer, d-d transition and H.O.M.O.L.U.M.O transition. Most of the transition metal transition metal complex compound are coloured either due to d-d electrons transition or charge transfer and ionic compounds are coloured due to polarisation of anion. Q. If  $MCl_4$  salt is white, then comment on colour of its iodine salt.

A. Coloured

B. White

C. May be or coloured

D. Black

#### Answer: C



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**3.** Hydrogen bond is the given to the relatively weak secondry interaction between a hydrogen atom bound to an electronegative atom and

another atom which is also generally electronegative and which has one or more lone pairs and can thus acts as a base. We can give the following generalized representation of hydrogen bond.

$$\overset{\delta-}{X}-\overset{\delta+}{H}...Y$$

Bond dissociation energy of H-bond ranges from 8 and 42kJ/ol, and the most commonly encountred hydrogen bonds are  $O-H...O,\,N-H...O$  and F-H...E.

Q. Among molecules of HCl, which of the following forces are present.

A. Interaction between two HCl molecules is found to be greater than 8kJ/mol

- B. Weak dipole-dipole interaction
- C. Weak ion-dipole interactions
- D. All of above

#### Answer: B



**4.** Hydrogen bond is the given to the relatively weak secondry interaction between a hydrogen atom bound to an electronegative atom and another atom which is also generally electronegative and which has one or more lone pairs and can thus acts as a base. We can give the follwoing generalized representation of hydrogen bond.

$$\overset{\delta-}{X}-\overset{\delta+}{H}...Y$$

Bond dissociation energy of H-bond ranges from 8 and 42kJ/ol, and the most commonly encountred hydrogen bonds are  $O-H...O,\,N-H...O$  and F-H...E.

Q.Which of the following interaction has energy between 8-42kJ/mol?

- A.  $Na^+:CCl_4$
- B.  $CHCl_3$  :  $Br^-$
- $C. C_6H_6:CCl_4$
- D.  $H_2O:HCN$

#### **Answer: D**



- **5.** The intermolecular forces of attraction (i.e., H-bonding and van der waal's forces) exist among polar and non-polar species which effect melting point, boiling point, solubility and viscosity of covalent compounds:
- Q. Melting and boling points of halogen increase down the group due to
  - A. Increase in London dispersion forces
  - B. Increase in extent of polarity
  - C. Increase in Molecular mass
  - D. Both (a) and (b)

#### Answer: D

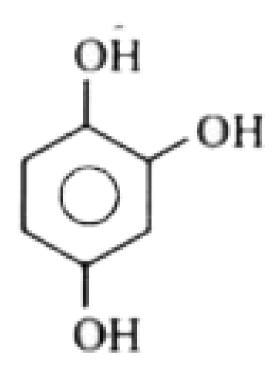


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**6.** The intermolecular forces of attraction (i.e., H-bonding and van der waal's forces) exist among polar and non-polar species which effect melting point, boiling point, solubility and viscosity of covalent

compounds:

Q.The type of molecular force of attraction present in the following compounds is :



A. Intermolecular H-bonding

B. Intramolecular H-bonding

C. van der waal's force

D. All of these

**Answer: D** 

**7.** The intermolecular forces of attraction (i.e., H-bonding and van der waal's forces) exist among polar and non-polar species which effect melting point, boiling point, solubility and viscosity of covalent compounds:

Q. Select the incorrect order of boiling point between the following compounds:

A. 
$$N_3H < CH_3N_3$$

B. 
$$Me_2SO_4 < H_2SO_4$$

$$\mathsf{C.}\, Me_3BO_3 < B(OH)_3$$

D. 
$$BF_3 < BI_3$$

#### Answer: A



**8.** There are five species P, Q, R, S and T. Spectroscopical analysis shows that P,Q and R are homonuclear diatomic species and have their bond order 2.5, 1.5 and 2.5 respectively and rest two species S and T are heteronuclear diatomic species and have bond order 3 and 2 respectively. All homonuclear diatomic species are paramagnetic and all heteronuclear species are diamagnetic in nature. P, R, S and T are monocovalent positive ion and Q is monovalent negative ion:

 $\ensuremath{\mathbf{Q}}.$  According to given information the incorrect match is :

A. 
$$P=N_2^{\,+}$$

$$\mathrm{B.}\,R=O_2^+$$

$$\mathsf{C.}\,S = CO^+$$

D. 
$$T=CN^+$$

#### Answer: C

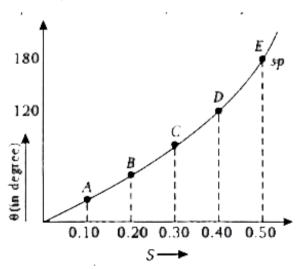


Compound, $MX_n$ type $(n = 2$ or 3 or 4)	Value of cos θ (θ = bond angle between equivalent hybrid orbitals)
P	-0.241
Q	-0.292
R	-0.5
S	-0.325
T	-0.469

9.

In all expected compounds each central atom only uses its s and p-orbitals in hybridization. The relationship between bond angle ' $\theta$ ' and decimal fraction of s and p character present in the equivalent orbitals is given by:

 $\cos\theta=rac{S}{S-1}=rac{P-1}{P},$  S=decimal fraction of s-character in the equivalent hybrid orbital and P=decimal fraction of p-character in the equivalent hybrid orbital.



Q. If the value n is 2 for compound T, then number of lone pair present at central atom of compound T will be :

**A.** 0

B. 1

C. 2

D. 3

#### **Answer: B**



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10. The comcept of redistribution of energy in different orbitals of an atom associated with different energies to give new orbitals of equal (or somethimes it may be non-equal) energy oriented in space in definite directions is called hybridization and formed new orbitals are called hybrid orbitals. The bonds formed by such orbitals are called hybrid bonds. The process of mixing of orbitals itself requires some energy. Thus, some additional energy, is needed for the hybridisation (mixing) of

Q. Select from each set the molecule or ion having the smallest bond angle:

- (i)  $H_2Se$ ,  $H_2Te$  and  $PH_3$
- (ii)  $NO_2^-$  and  $NH_2^-$

atomic orbitals.

- (iii)  $POF_3$  and  $POCl_3$  (X P X angle)
- (iv)  $OSF_2Cl_2$  and  $SF_2(CH_3)_2$  (F-S-F angle)
  - A.  $H_2Se$ ,  $NH_2^-POF_3$  and  $OSF_2Cl_2$
  - B.  $H_2Te, NO_2^-, POF_3$  and  $SF_2(CH_3)_2$

 $\mathsf{C.}\,PH_3,NH_2^{\,-},POCl_3 \,\,\,\mathrm{and}\,\,\,SF_2(CH_3)_2$ 

D.  $H_2Te, NH_2^-, POF_3 \text{ and } SF_2(CH_3)_2$ 

#### Answer: D



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11. The comcept of redistribution of energy in different orbitals of an atom associated with different energies to give new orbitals of equal (or somethimes it may be non-equal) energy oriented in space in definite directions is called hybridization and formed new orbitals are called hybrid orbitals. The bonds formed by such orbitals are called hybrid bonds. The process of mixing of orbitals itself requires some energy. Thus, some additional energy, is needed for the hybridisation (mixing) of atomic orbitals.

Q. Which of the following statement is correct?

A. In  $BrF_3$ , maximum three halogen atoms can lie in same plane

B. In  $CH_2SF_2(CH_3)_2$  molecule all hydrogen atoms which bonded to

 $s-sp^2$ overlapping, lie in equitorial plane

C. In  $OSCl_4,\,Cl-S-Cl$  equitorial bond angle is greater than  $120^\circ$ 

D. Molecules  $IOF_5$  and  $XeO_2F_4$  have similar shape but have different number of lone pairs in whole molecule

#### **Answer: B**



12. The comcept of redistribution of energy in different orbitals of an atom associated with different energies to give new orbitals of equal (or somethimes it may be non-equal) energy oriented in space in definite directions is called hybridization and formed new orbitals are called hybrid orbitals. The bonds formed by such orbitals are called hybrid bonds. The process of mixing of orbitals itself requires some energy. Thus, some additional energy, is needed for the hybridisation (mixing) of atomic orbitals.

Q. In neutral moleule  $XeO_{n_1}F_{n_2}$ , central atom has no lone pair and ratio of  $\frac{n_2}{n_1}$  is two, then which of the following orbitals does not participate in bonding ( $n_1$  and  $n_2$  are natural numbers):

A.  $d_{x^2}$ 

B.  $p_x$ 

 $C.d_{r^2-n^2}$ 

D. None of these

### Answer: D



13. Drago suggested an emprical rule which is compatible with the energetics of hybridization. It will occupy a stereochemically inactive sorbital, and the bonding will be through p-orbitals and bond angles will be nearly  $90^\circ$  if the electronegativity of the surrounding atoms is  $\leq 2.5$ . Q. In which of the following molecule central atom has higher % scharacter in its bond pair:

- A.  $AsH_3$
- B.  $GeH_3$
- $\mathsf{C}.\,P_4$
- D.  $H_2Se$

**Answer: B** 



**14.** Drago suggested an emprical rule which is compatible with the energetics of hybridization. It will occupy a stereochemically inactive sorbital, and the bonding will be through p-orbitals and bond angles will be nearly  $90^\circ$  if the electronegativity of the surrounding atoms is  $\leq 2.5$ . Q. Correct order of bond angle is :

A. 
$$PH_4^{\,+}>OF_2>SF_2>SbH_3>H_2Te$$

B. 
$$OF_2>SF_2>PH_4^{\ +}>SbH_3>H_2Te$$

C. 
$$PH_{\scriptscriptstyle A}^{\,+} > SF_2 > OF_2 > SbH_3 > H_2Te$$

D. 
$$SF_2 > OF_2 > PH_4^{\,+} > SbH_3 > H_2Te$$

#### Answer: A



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**15.** According to hybridisation theory, the % s-character in  $sp, sp^2$  and  $sp^3$  hybrid orbitals is 50, 33.3 and 25 respectively, but this is not true for all the species. When  $\theta$  is the bond angle between equivalent

hybrid orbitals then % s and p-character in hybrid orbitals (when only s and p-orbitals are involved in hybridisation) can be calculated by the following formula:

$$\cos \theta = \frac{S}{S-1} = \frac{P-1}{P}$$

Q. Two elements X and Y combined together to form a covalent compound. If % p-character is found to be 80% in a orbital then the hybridised state of central atom X for the orbital is:

A. 
$$sp^2$$

В.  $sp^3$ 

 $\mathsf{C}.\,sp^4$ 

D.  $sp^5$ 

## **Answer: C**



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According hybridisation theory, the % s-character 16. to in  $sp,\,sp^2 \,\,{
m and}\,\,sp^3$  hybrid orbitals is 50, 33.3 and 25 respectively, but this is not true for all the species. When  $\theta$  is the bond angle between equivalent hybrid orbitals then % s and p-character in hybrid orbitals (when only s and p-orbitals are involved in hybridisation) can be calculated by the following formula:

$$\cos\theta = \frac{S}{S-1} = \frac{P-1}{P}$$

Q. Correct order of P-P bond length in the following compound is :

A. 
$$P_2F_4 < P_2(CH_3)_4 < P_2(CF_3)_4 < P_2H_4$$

$$\mathsf{B}.\, P_2F_4 < P_2(CF_3)_4 < P_2(CH_3)_4 < P_2H_4$$

C. 
$$P_2F_4 < P_2H_4 < P_2(CH_3)_4 < P_2(CF_3)_4$$

D. 
$$P_2F_4 < P_2(CH_3)_4 < P_2H_4 < P_2(CF_3)_4$$

#### **Answer: B**



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17. The correct statement is:

A. in  $CH_2F_2$  the F-C-F bond angle is larger  $109\,^\circ\,28\,'$ 

B. in  $CH_2F_2$  the C-F bond has more than 25% s-character

C. in  $CH_2F_2$  the H-C-H bond angle is larger  $109^{\circ}\,28$  '

D. in  $CH_2F_2$  the C-H bond has more than 25% s-character

# Answer: C,D



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**18.**  $PCl_5$  is an example of a molecule having  $sp^3d$ -hybridisation. Three out of the orbitals in  $sp_xp_y$ -hybridisation while remaining two have  $p_xd_{x^2}$ -hybridisation. If P-atom is attached to substitutuents differ in electronegativity, as in  $PCl_xF_{5-x}$ , then it is has been experimently observed that the more electronegativity substituent occupies the axial position of t.b.p geometry.

Q. The incorrect statement regarding  $PCl_2F_2^-$  molecule will be:

A. given compound is polar

B. both axial position occupied by F-atoms

C. both Cl atoms present in equatorial position

D. one Cl atom present at axial and other Cl atom is present at equatorial position of geometry

#### **Answer: D**



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**19.**  $PCl_5$  is an example of a molecule having  $sp^3d$ -hybridisation. Three out of the orbitals in  $sp_xp_y$ -hybridisation while remaining two have  $p_xd_{x^2}$ -hybridisation. If P-atom is attached to substitutuents differ in electronegativity, as in  $PCl_xF_{5-x}$ , then it is has been experimently observed that the more electronegativity substituent occupies the axial position of t.b.p geometry.

Q. The highest H-C-H bond angle present in :

A.  $CH_2F_2$ 

B.  $CH_4$ 

 $\mathsf{C}.\,CH_3F$ 

D.

#### **Answer: A**



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**20.** The first compound of the noble gasees was made in 1962. Barlett and Lohman had previously used the highly oxidizing compound platium hexaflouride to oxidize dioxygen.

$$O_2 + PtF_6 
ightarrow O_2^+ \left[ PtF_6 
ight]^-$$

The first ionization energy for  $O_2 o O_2$  is  $1165kJ \cdot mol^{-1}$ , which is almost as the value of  $1170kJ \cdot mol^{-1}$  for  $Xe o Xe^+$ . It was predicated that xenon should react with  $PtF_6$ . Experimental showed that when deep of red  $PtF_6$  vapour was mixed with an equal volume of Xe the gases combined immediately at room temperature to produce a solid.

$$Xe + PtF_6 
ightarrow Xe[PtF_6]$$

Q. IUPAC name of first xenon-compounds synthesized by secientist Barlett is:

A. xenonhexaflouroplatinate(IV)

B. xenonhexaflouroplatinate(V)

C. hexafluoroplatinum(V) xenon

D. xenoniumhexafluoroplatinum(V)

#### Answer: B



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**21.** The first compound of the noble gasees was made in 1962. Barlett and Lohman had previously used the highly oxidizing compound platium hexaflouride to oxidize dioxygen.

$$O_2 + PtF_6 
ightarrow O_2^+ \left[ PtF_6 
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 $Xe + PtF_6 
ightarrow Xe[PtF_6]$ 

Q. Noble gases are water insoluble, however decrease their insolubility in water decrease down the group due to increase in :

A. dipole-dipole attraction

B. dipole-induced dipoles

C. instantaneous dipole-induced dipole attraction

D. None of these

#### **Answer: B**



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**22.** The first compound of the noble gasees was made in 1962. Barlett and Lohman had previously used the highly oxidizing compound platium hexaflouride to oxidize dioxygen.

$$O_2 + PtF_6 
ightarrow O_2^+ [PtF_6]^-$$

The first ionization energy for  $O_2 o O_2$  is  $1165kJ\cdot mol^{-1}$ , which is almost as the value of  $1170kJ\cdot mol^{-1}$  for  $Xe o Xe^+$ . It was predicated

that xenon should react with  $PtF_6$ . Experimental showed that when deep of red  $PtF_6$  vapour was mixed with an equal volume of Xe the gases combined immediately at room temperature to produce a solid.

$$Xe + PtF_6 
ightarrow Xe[PtF_6]$$

Q. Which of the following species is not having perfect octahedron structure?

- A.  $XeF_6$
- B.  $SiF_6^{\,2\,-}$
- $\mathsf{C.}\,PCl_6^-$
- D.  $XeO_6^{4\,-}$

#### Answer: A



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formation of molecular orbital formed. Number of atomic orbitals overlapping together is equal to the molecule orbital formed. The two

23. According to MOT, two atomic orbitals overlap resulting in the

atomic orbital thus formed by LCAO (linear combination of atomic orbital) in the phase or in the different phase are known as bonding and antibonding molecular orbitals respectively. The energy of bonding molecular orbital is lower than that of the pure atomic orbitals by an amount  $\Delta$ . This known as the stabilization energy. The enerby of antibonding molecular orbital in increased by  $\Delta'$  (destabilisation energy).

Q . The bond order of  $N_2^-$  is equal to that of

A.  $O_2$ 

B.  $O_2^{2\,-}$ 

 $\mathsf{C.}\,O_2^{\,+}$ 

D. None

#### **Answer: C**



**24.** According to MOT, two atomic orbitals overlap resulting in the formation of molecular orbital formed. Number of atomic orbitals overlapping together is equal to the molecule orbital formed. The two atomic orbital thus formed by LCAO (linear combination of atomic orbital) in the phase or in the different phase are known as bonding and antibonding molecular orbitals respectively. The energy of bonding molecular orbital is lower than that of the pure atomic orbitals by an amount  $\Delta$ . This known as the stabilization energy. The enerby of antibonding molecular orbital in increased by  $\Delta'$  (destabilisation energy).

Q. Which of the following statement(s) is true:

- A. Higher the bond order lesser the bond length.
- B. Higher the bond order greater the bond length.
- C. Higher the bond order lesser the bond energy.
- D. Higher the bond order lesser the number of bonds

**25.** According to MOT, two atomic orbitals overlap resulting in the formation of molecular orbital formed. Number of atomic orbitals overlapping together is equal to the molecule orbital formed. The two atomic orbital thus formed by LCAO (linear combination of atomic orbital) in the phase or in the different phase are known as bonding and antibonding molecular orbitals respectively. The energy of bonding molecular orbital is lower than that of the pure atomic orbitals by an amount  $\Delta$ . This known as the stabilization energy. The enerby of antibonding molecular orbital in increased by  $\Delta'$  (destabilisation energy).

Q. How many nodal plane is present in  $\sigma_{s \text{ and } p}$  bonding molecular orbital ?

A. zero

B. 1

C. 2

#### Answer: A



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26. According to MOT, two atomic orbitals overlap resulting in the formation of molecular orbital formed. Number of atomic orbitals overlapping together is equal to the molecule orbital formed. The two atomic orbital thus formed by LCAO (linear combination of atomic orbital) in the phase or in the different phase are known as bonding and antibonding molecular orbitals respectively. The energy of bonding molecular orbital is lower than that of the pure atomic orbitals by an amount  $\Delta$ . This known as the stabilization energy. The enerby of antibonding molecular orbital in increased by  $\Delta'$  (destabilisation energy).

Q. which of the following combination of orbitals is corrects?

A. (a) (-) +(-) -- (-) (-)

**Answer: C::D** 



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27. According to MOT, two atomic orbitals overlap resulting in the formation of molecular orbital formed. Number of atomic orbitals overlapping together is equal to the molecule orbital formed. The two atomic orbital thus formed by LCAO (linear combination of atomic orbital) in the phase or in the different phase are known as bonding and antibonding molecular orbitals respectively. The energy of bonding molecular orbital is lower than that of the pure atomic orbitals by an amount  $\Delta$ . This known as the stabilization energy. The enerby of antibonding molecular orbital in increased by  $\Delta'$  (destabilisation energy).

Q. Which of the following statements is not correct regarding bonding molecular orbitals?

A. Bonding molecular orbitals possess low energy than the atomic orbitals from which they are formed

B. Bonding molecular orbitals have low electron density the two nuclei

C. Electron in bonding molecular contributes to the attraction between atoms.

D. They are formed when the lobes of the combining atomic orbitals have the same sign.

### **Answer: B**



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**28.** According to MOT, two atomic orbitals overlap resulting in the formation of molecular orbital formed. Number of atomic orbitals overlapping together is equal to the molecule orbital formed. The two

atomic orbital thus formed by LCAO (linear combination of atomic orbital) in the phase or in the different phase are known as bonding and antibonding molecular orbitals respectively. The energy of bonding molecular orbital is lower than that of the pure atomic orbitals by an amount  $\Delta$ . This known as the stabilization energy. The enerby of antibonding molecular orbital in increased by  $\Delta'$  (destabilisation energy).

Q. If x-axis is the molecular axis, then  $\pi$ -moleulcar orbitals are formed by the overlap of :

A.  $S+p_x$ 

B.  $p_x + p_y$ 

 $\mathsf{C.}\,P_z+p_z$ 

D.  $p_x + p_x$ 

### **Answer: C**



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**29.** Polar covalent molecules exhibit dipole moment. Dipole moment is equal to the product of charge separation , q and the bond length d for the bond. Unit of dipole moment is debye. One debye is equal to  $10^{-18}$  esu cm.

Dipole moments is a vector quantity. It has both magnitude and direction. Hence, dipole moment of a molecule depends upon the relative orientation of the bond dipoles, but not on the polarity of bonds alone. A symmetrical structure shows zero dipole moment. Thus, dipole moment helps to predict the geometry of a molecules. Dipole moment values can be distinguish between cis- and trans- isomers, ortho, meta and pareforms of a substance, etc.

Q. Which is a polar molecule?

- A.  $XeF_4$
- $\mathsf{B.}\,BF_3$
- C.  $I_2Cl_6$
- D.  $PCl_2F_3$

### **Answer: D**



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**30.** Polar covalent molecules exhibit dipole moment. Dipole moment is equal to the product of charge separation , q and the bond length d for the bond. Unit of dipole moment is debye. One debye is equal to  $10^{-18}$  esu cm.

Dipole moments is a vector quantity. It has both magnitude and direction. Hence, dipole moment of a molecule depends upon the relative orientation of the bond dipoles, but not on the polarity of bonds alone. A symmetrical structure shows zero dipole moment. Thus, dipole moment helps to predict the geometry of a molecules. Dipole moment values can be distinguish between cis- and trans- isomers, ortho, meta and pareforms of a substance, etc.

Q. A diatomic molecule has a dipole moment of 1.2D. If the bond length is  $1.0 imes 10^{-8}$  cm, what fraction of charge does exist each atom?

- B.0.2
- $\mathsf{C.}\ 0.25$
- $\mathsf{D.}\ 0.3$

### **Answer: C**



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31. Polar covalent molecules exhibit dipole moment. Dipole moment is equal to the product of charge separation , q and the bond length d for the bond. Unit of dipole moment is debye. One debye is equal to  $10^{-18}$  esu cm.

Dipole moments is a vector quantity. It has both magnitude and direction. Hence, dipole moment of a molecule depends upon the relative orientation of the bond dipoles, but not on the polarity of bonds alone. A symmetrical structure shows zero dipole moment. Thus, dipole moment helps to predict the geometry of a molecules. Dipole moment values can be distinguish between cis- and trans- isomers, ortho, meta and pare-

forms of a substance, etc.

Q. Arrange the following compounds in increasing order of dipole moments, toluene (I), o-dichlorobenzene (II), m-dichlorobenzene (III) and p-dichlorobenzene (IV):

A. 
$$IV < I < II < III$$

$$\mathsf{B}.\,I < IV < II < III$$

$$\mathsf{C}.\,IV < I < III < II$$

$$\mathsf{D}.\,IV < II < I < III$$

### Answer: C



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32. Polar covalent molecules exhibit dipole moment. Dipole moment is equal to the product of charge separation , q and the bond length d for the bond. Unit of dipole moment is debye. One debye is equal to  $10^{-18}$  esu cm.

Dipole moments is a vector quantity. It has both magnitude and direction.

Hence, dipole moment of a molecule depends upon the relative orientation of the bond dipoles, but not on the polarity of bonds alone. A symmetrical structure shows zero dipole moment. Thus, dipole moment helps to predict the geometry of a molecules. Dipole moment values can be distinguish between cis- and trans- isomers, ortho, meta and pareforms of a substance, etc.

Q.  $\mu$  of the  $AX_4$  type of molecule is zero. The geometry of it can be :

A. tetrahedral

B. square planar

C. A or B

D. None of these

### Answer: C



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**33.** Polar covalent molecules exhibit dipole moment. Dipole moment is equal to the product of charge separation , q and the bond length d for

the bond. Unit of dipole moment is debye. One debye is equal to  $10^{-18}$  esu cm.

Dipole moments is a vector quantity. It has both magnitude and direction.

Hence, dipole moment of a molecule depends upon the relative orientation of the bond dipoles, but not on the polarity of bonds alone. A symmetrical structure shows zero dipole moment. Thus, dipole moment helps to predict the geometry of a molecules. Dipole moment values can be distinguish between cis- and trans- isomers, ortho, meta and pareforms of a substance, etc.

Q. Which of the following statement is correct regarding  $C_2H_2F_2$  molecule?

- A. One isomer is polar, one is non-polar
- B. Two isomers are polar, one is non-polar
- C. Two isomers are planar, one is non-planar
- D. Two isomers are only possible and planar

### Answer: B



**34.** In general boiling point of covalent compounds is affect due to increasing molecular weight and hydrogen bonding. Thus it is obversed that the boiling point of octanol is greater that water while that of methanol is lesser than water. The other properties which are considerably affected due to hydrogen bond formation are acidic nature, melting point, anamalous behaviour of water below  $4^{\circ}C$ ., It is therefore concluded that hydrogen bonding palys a vutal role in explaning many observable facts.

Q. Which among the following has maximum boiling point?

A. 
$$CH_2-CH_2-\left(CH_2\right)_2-CH_2OH$$

B.  $CH_3-CH_2-CH_2-CH-CH_3$ 

C.  $CH_3-CH_2-CH-CH_2-CH_3$ 
 $CH_3$ 

D.  $CH_3-CH_2-CH_2-CH_3$ 

**35.** In general boiling point of covalent compounds is affect due to increasing molecular weight and hydrogen bonding. Thus it is obversed that the boiling point of octanol is greater that water while that of methanol is lesser than water. The other properties which are considerably affected due to hydrogen bond formation are acidic nature, melting point, anamalous behaviour of water below  $4^{\circ}$  C., It is therefore concluded that hydrogen bonding palys a vutal role in explaning many observable facts.

Q. Which of the following statements is true?

bonding

- A. The lattice structure of ice involves true covalent bond
- B. The lattice structure of ice is the result of dipole-dipole interaction
- C. The lattice structure of ice is the result of intra and inter-molecular hydrogen bond formation
- D. The lattice structure of ice is only due to inter-molecular hydrogen

### **Answer: D**



**36.** In general boiling point of covalent compounds is affect due to increasing molecular weight and hydrogen bonding. Thus it is obversed that the boiling point of octanol is greater that water while that of methanol is lesser than water. The other properties which are considerably affected due to hydrogen bond formation are acidic nature, melting point, anamalous behaviour of water below  $4^{\circ}C$ ., It is therefore concluded that hydrogen bonding palys a vutal role in explaning many observable facts.

Q.  $K_{a_2}$  of maleic acid is lesser than  $K_{a_2}$  of fumaric acid due to:

A. Intramolecular hydrogen bonding in the formed after one proton removal in fumaric acid

B. Intermolecular hydrogen bonding in the ion formed after one proton removal in maleic acid

C. Intramolecular hydrogen bonding in the ion formed after one proton removal in maleic acid

D. Intermolecular hydrogen bonding in the ion formed after one proton removal in fumaric acid

### **Answer: C**



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37. The molecule in which an atom is associated with more than 8 electrons is known as hypervalent molecule and less than 8 electrons is known as hypovalent molecule. All hypervalent molecules must have  $p\pi-d\pi$  bonding but the molecules having back bonding back boding need not to have always  $p\pi-d\pi$  bondin.

Q.Which of the molecule is not hypovalent but complete its octet:

- A.  $AlI_3$
- B.  $AlCl_3$

- C.  $AlBr_3$
- D.  $AlF_3$

### **Answer: D**



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**38.** The molecule in which an atom is associated with more than 8 electrons is known as hypervalent molecule and less than 8 electrons is known as hypovalent molecule. All hypervalent molecules must have  $p\pi-d\pi$  bonding but the molecules having back bonding back boding need not to have always  $p\pi-d\pi$  bondin.

Q. Which of the following is having complete octet:

- A.  $BeCl_2$  (dimer)
- B.  $BeH_2$  (dimer)
- C.  $BeH_2$  (s)
- D.  $BeCl_2$  (s)

### **Answer: D**



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**39.** The molecule in which an atom is associated with more than 8 electrons is known as hypervalent molecule and less than 8 electrons is known as hypovalent molecule. All hypervalent molecules must have  $p\pi-d\pi$  bonding but the molecules having back bonding back boding need not to have always  $p\pi-d\pi$  bondin.

Q.Which of the following molecule is not having  $p\pi-d\pi$  bonding:

- A.  $SO_2$
- B.  $P_4O_{10}$
- $\mathsf{C}.\,PF_3$
- D.  $B_3N_3H_6$

### **Answer: D**



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**40.** Colour of compounds occurs due to phenomenon of polarisation, charge transfer, d-d transition and H.O.M.O.L.U.M.O transition. Most of the transition metal transition metal complex compound are coloured either due to d-d electrons transition or charge transfer and ionic compounds are coloured due to polarisation of anion. Q. If  $MCl_4$  salt is white, then comment on colour of its iodine salt.

A. Coloured

B. White

C. May be or coloured

D. Black

Answer: C



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**41.** Hydrogen bond is the given to the relatively weak secondry interaction between a hydrogen atom bound to an electronegative atom and another atom which is also generally electronegative and which has one or more lone pairs and can thus acts as a base. We can give the following generalized representation of hydrogen bond.

$$\overset{\delta-}{X} - \overset{\delta+}{H}...Y$$

Bond dissociation energy of H-bond ranges from 8 and 42kJ/ol, and the most commonly encountred hydrogen bonds are  $O-H...O,\,N-H...O$  and F-H...E.

Q. Among molecules of HCl, which of the following forces are present.

A. Interaction between two HCl molecules is found to be greater than

8kJ/mol

B. Weak dipole-dipole interaction

C. Weak ion-dipole interactions

D. All of above

## Answer: B

**42.** Hydrogen bond is the given to the relatively weak secondry interaction between a hydrogen atom bound to an electronegative atom and another atom which is also generally electronegative and which has one or more lone pairs and can thus acts as a base. We can give the follwoing generalized representation of hydrogen bond.

$$\overset{\delta-}{X} - \overset{\delta+}{H}...Y$$

Bond dissociation energy of H-bond ranges from 8 and 42kJ/ol, and the most commonly encountred hydrogen bonds are  $O-H...O,\,N-H...O$  and F-H...E.

Q.Which of the following interaction has energy between 8-42kJ/mol?

- A.  $Na^+:CCl_4$
- B.  $CHCl_3$ :  $Br^-$
- $C. C_6H_6:CCl_4$
- D.  $H_2O:HCN$

### **Answer: D**



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- **43.** The intermolecular forces of attraction (i.e., H-bonding and van der waal's forces) exist among polar and non-polar species which effect melting point, boiling point, solubility and viscosity of covalent compounds:
- Q. Melting and boling points of halogen increase down the group due to
  - A. Increase in London dispersion forces
  - B. Increase in extent of polarity
  - C. Increase in Molecular mass
  - D. Both (a) and (b)

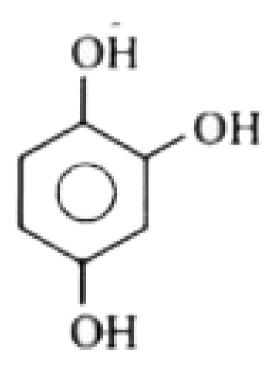
### **Answer: D**



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**44.** The intermolecular forces of attraction (i.e., H-bonding and van der waal's forces) exist among polar and non-polar species which effect melting point, boiling point, solubility and viscosity of covalent compounds:

Q.The type of molecular force of attraction present in the following compounds is :



A. Intermolecular H-bonding

B. Intramolecular H-bonding

- C. van der waal's force
- D. All of these

### **Answer: D**



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- **45.** The intermolecular forces of attraction (i.e., H-bonding and van der waal's forces) exist among polar and non-polar species which effect melting point, boiling point, solubility and viscosity of covalent compounds:
- Q. Select the incorrect order of boiling point between the following compounds:
  - A.  $N_2 H < C H_3 N_3$
  - $\mathrm{B.}\,Me_2SO_4 < H_2SO_4$
  - C.  $Me_3BO_3 < B(OH)_3$
  - D.  $BF_3 < BI_2$

### **Answer: A**

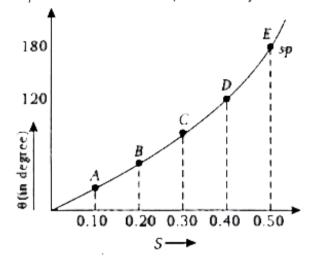


Compound, $MX_n$ type $(n = 2$ or 3 or 4)	Value of cos θ (θ = bond angle between equivalent hybrid orbitals)
P	-0.241
Q	-0.292
R	-0.5
S	-0.325
Т	-0.469

46.

In all expected compounds each central atom only uses its s and p-orbitals in hybridization. The relationship between bond angle  $'\theta'$  and decimal fraction of s and p character present in the equivalent orbitals is given by:

 $\cos\theta=rac{S}{S-1}=rac{P-1}{P},$  S=decimal fraction of s-character in the equivalent hybrid orbital and P=decimal fraction of p-character in the equivalent hybrid orbital.



Q. If the value n is 2 for compound T, then number of lone pair present at central atom of compound T will be :

- **A.** 0
- **B**. 1
- C. 2
- D. 3

**Answer: B** 



**View Text Solution** 

47. The comcept of redistribution of energy in different orbitals of an atom associated with different energies to give new orbitals of equal (or somethimes it may be non-equal) energy oriented in space in definite directions is called hybridization and formed new orbitals are called hybrid orbitals. The bonds formed by such orbitals are called hybrid bonds. The process of mixing of orbitals itself requires some energy. Thus, some additional energy, is needed for the hybridisation (mixing) of atomic orbitals.

Q. Select from each set the molecule or ion having the smallest bond angle:

- (i)  $H_2Se$ ,  $H_2Te$  and  $PH_3$
- (ii)  $NO_2^-$  and  $NH_2^-$
- (iii)  $POF_3$  and  $POCl_3$  (X P X angle)
- (iv)  $OSF_2Cl_2$  and  $SF_2(CH_3)_2$  (F-S-F angle)
  - A.  $H_2Se$ ,  $NH_2^-POF_3$  and  $OSF_2Cl_2$
  - B.  $H_2Te$ ,  $NO_2^-$ ,  $POF_3$  and  $SF_2(CH_3)_2$
  - C.  $PH_3$ ,  $NH_2^-$ ,  $POCl_3$  and  $SF_2(CH_3)_2$

D.  $H_2Te, NH_2^-, POF_3 \text{ and } SF_2(CH_3)_2$ 

### **Answer: D**



**View Text Solution** 

**48.** The comcept of redistribution of energy in different orbitals of an atom associated with different energies to give new orbitals of equal (or somethimes it may be non-equal) energy oriented in space in definite directions is called hybridization and formed new orbitals are called hybrid orbitals. The bonds formed by such orbitals are called hybrid bonds. The process of mixing of orbitals itself requires some energy. Thus, some additional energy, is needed for the hybridisation (mixing) of atomic orbitals.

Q. Which of the following statement is correct?

A. In  $BrF_3$ , maximum three halogen atoms can lie in same plane

B. In  $CH_2SF_2(CH_3)_2$  molecule all hydrogen atoms which bonded to

 $s-sp^2$  overlapping, lie in equitorial plane

C. In  $OSCl_4$ , Cl-S-Cl equitorial bond angle is greater than  $120^\circ$ 

D. Molecules  $IOF_5$  and  $XeO_2F_4$  have similar shape but have different number of lone pairs in whole molecule

### **Answer: B**



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**49.** The comcept of redistribution of energy in different orbitals of an atom associated with different energies to give new orbitals of equal (or somethimes it may be non-equal) energy oriented in space in definite directions is called hybridization and formed new orbitals are called hybrid orbitals. The bonds formed by such orbitals are called hybrid bonds. The process of mixing of orbitals itself requires some energy. Thus, some additional energy, is needed for the hybridisation (mixing) of atomic orbitals.

Q. In neutral moleule  $XeO_{n_1}F_{n_2}$ , central atom has no lone pair and ratio

of  $\frac{n_2}{n_1}$  is two, then which of the following orbitals does not participate in

bonding ( $n_1$  and  $n_2$  are natural numbers):

A.  $d_{x^2}$ 

 $B. p_r$ 

C.  $d_{x^2-y^2}$ 

D. None of these

### Answer: D



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character in its bond pair:

energetics of hybridization. It will occupy a stereochemically inactive sorbital, and the bonding will be through p-orbitals and bond angles will be nearly  $90^{\circ}$  if the electronegativity of the surrounding atoms is  $\leq 2.5$ . Q. In which of the following molecule central atom has higher % s-

50. Drago suggested an emprical rule which is compatible with the

A.  $AsH_3$ 

 $\operatorname{B.} \operatorname{GeH}_4$ 

 $\mathsf{C}.\,P_4$ 

D.  $H_2Se$ 

### **Answer: B**



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**51.** Drago suggested an emprical rule which is compatible with the energetics of hybridization. It will occupy a stereochemically inactive sorbital, and the bonding will be through p-orbitals and bond angles will be nearly  $90^\circ$  if the electronegativity of the surrounding atoms is  $\leq 2.5$ . Q. Correct order of bond angle is :

Q. Correct order of boild aligie is

A. 
$$PH_4^{\,+}>OF_2>SF_2>SbH_3>H_2Te$$

B. 
$$OF_2>SF_2>PH_4^{\ +}>SbH_3>H_2Te$$

C. 
$$PH_4^{\,+}\,>SF_2>OF_2>SbH_3>H_2Te$$

D. 
$$SF_2 > OF_2 > P{H_4}^+ > SbH_3 > H_2Te$$

#### **Answer: A**



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**52.** According to hybridisation theory, the % s-character in sp,  $sp^2$  and  $sp^3$  hybrid orbitals is 50, 33.3 and 25 respectively, but this is not true for all the species. When  $\theta$  is the bond angle between equivalent hybrid orbitals then % s and p-character in hybrid orbitals (when only s and p-orbitals are involved in hybridisation) can be calculated by the following formula:

$$\cos\theta = \frac{S}{S-1} = \frac{P-1}{P}$$

Q. Two elements X and Y combined together to form a covalent compound. If % p-character is found to be 80% in a orbital then the hybridised state of central atom X for the orbital is :

A.  $sp^2$ 

 $\mathsf{B}.\,sp^3$ 

 $\mathsf{C}.\,sp^4$ 

D.  $sp^5$ 

### **Answer: C**



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**53.** According to hybridisation theory, the % s-character in sp,  $sp^2$  and  $sp^3$  hybrid orbitals is 50, 33.3 and 25 respectively, but this is not true for all the species. When  $\theta$  is the bond angle between equivalent hybrid orbitals then % s and p-character in hybrid orbitals (when only s and p-orbitals are involved in hybridisation) can be calculated by the following formula:

$$\cos\theta = \frac{S}{S-1} = \frac{P-1}{P}$$

Q. Smallest  $O\widehat{S}O$  bond angle is found in :

A.  $SO_2F_2$ 

B.  $SO_2Cl_2$ 

$$\mathsf{C}.\,SO_2(CF_3)_2$$

D. 
$$SO_2(CH_3)_2$$

### **Answer: D**



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**54.** According to hybridisation theory, the % s-character in sp,  $sp^2$  and  $sp^3$  hybrid orbitals is 50, 33.3 and 25 respectively, but this is not true for all the species. When  $\theta$  is the bond angle between equivalent hybrid orbitals then % s and p-character in hybrid orbitals (when only s and p-orbitals are involved in hybridisation) can be calculated by the following formula:

$$\cos\theta = \frac{S}{S-1} = \frac{P-1}{P}$$

Q. Correct order of P-P bond length in the following compound is :

A. 
$$P_2F_4 < P_2(CH_3)_4 < P_2(CF_3)_4 < P_2H_4$$

B. 
$$P_2F_4 < P_2(CF_3)_4 < P_2(CH_3)_4 < P_2H_4$$

C. 
$$P_2F_4 < P_2H_4 < P_2(CH_3)_{\scriptscriptstyle A} < P_2(CF_3)_{\scriptscriptstyle A}$$

D. 
$$P_2F_4 < P_2(CH_3)_4 < P_2H_4 < P_2(CF_3)_4$$

### **Answer: B**



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**55.**  $PCl_5$  is an example of a molecule having  $sp^3d$ -hybridisation. Three out of the orbitals in  $sp_xp_y$ -hybridisation while remaining two have  $p_xd_{x^2}$ -hybridisation. If P-atom is attached to substitutuents differ in electronegativity, as in  $PCl_xF_{5-x}$ , then it is has been experimently observed that the more electronegativity substituent occupies the axial position of t.b.p geometry.

Q. The incorrect statement regarding  $PCl_2F_2^-$  molecule will be:

A. given compound is polar

- B. both axial position occupied by F-atoms
- C. both Cl atoms present in equatorial position

D. one Cl atom present at axial and other Cl atom is present at equatorial position of geometry

### **Answer: D**



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**56.**  $PCl_5$  is an example of a molecule having  $sp^3d$ -hybridisation. Three out of the orbitals in  $sp_xp_y$ -hybridisation while remaining two have  $p_xd_{x^2}$ -hybridisation. If P-atom is attached to substitutuents differ in electronegativity, as in  $PCl_xF_{5-x}$ , then it is has been experimently observed that the more electronegativity substituent occupies the axial position of t.b.p geometry.

Q. The highest H-C-H bond angle present in :

A.  $CH_2F_2$ 

B.  $CH_4$ 

 $\mathsf{C}.\,CH_3F$ 

#### Answer: A



## **Watch Video Solution**

**57.** The first compound of the noble gasees was made in 1962. Barlett and Lohman had previously used the highly oxidizing compound platium hexaflouride to oxidize dioxygen.

$$O_2 + PtF_6 
ightarrow O_2^+ \left[PtF_6
ight]^-$$

The first ionization energy for  $O_2 o O_2$  is  $1165kJ \cdot mol^{-1}$ , which is almost as the value of  $1170kJ \cdot mol^{-1}$  for  $Xe o Xe^+$ . It was predicated that xenon should react with  $PtF_6$ . Experimental showed that when deep of red  $PtF_6$  vapour was mixed with an equal volume of Xe the gases combined immediately at room temperature to produce a solid.

$$Xe + PtF_6 \rightarrow Xe[PtF_6]$$

Q. IUPAC name of first xenon-compounds synthesized by secientist Barlett is:

A. xenonhexaflouroplatinate(IV)

B. xenonhexaflouroplatinate(V)

C. hexafluoroplatinum(V) xenon

D. xenoniumhexafluoroplatinum(V)

#### **Answer: B**



# **Watch Video Solution**

**58.** The first compound of the noble gases was made in 1962. Barlett and Lohman had previously used the highly oxidizing compound platium hexaflouride to oxidize dioxygen.

$$O_2 + PtF_6 
ightarrow O_2^+ \left[ PtF_6 
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 $Xe + PtF_6 
ightarrow Xe[PtF_6]$ 

Q. Noble gases are water insoluble, however their insolubility in water decreases down the group due to increase in :

A. dipole-dipole attraction

B. dipole-induced dipoles

C. instantaneous dipole-induced dipole attraction

D. None of these

#### **Answer: B**



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**59.** The first compound of the noble gasees was made in 1962. Barlett and Lohman had previously used the highly oxidizing compound platium hexaflouride to oxidize dioxygen.

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ightarrow O_2^+ [PtF_6]^-$$

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$$Xe + PtF_6 
ightarrow Xe[PtF_6]$$

Q. IUPAC name of first xenon-compounds synthesized by secientist Barlett

A.  $XeF_6$ 

is:

- B.  $SiF_6^{\,2\,-}$
- $\mathsf{C.}\,PCl_6^-$
- D.  $XeO_6^{4-}$

#### Answer: A



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formation of molecular orbital formed. Number of atomic orbitals overlapping together is equal to the molecule orbital formed. The two

60. According to MOT, two atomic orbitals overlap resulting in the

atomic orbital thus formed by LCAO (linear combination of atomic orbital) in the phase or in the different phase are known as bonding and antibonding molecular orbitals respectively. The energy of bonding molecular orbital is lower than that of the pure atomic orbitals by an amount  $\Delta$ . This known as the stabilization energy. The enerby of antibonding molecular orbital in increased by  $\Delta'$  (destabilisation energy).

Q. Which of the following statements is not correct regarding bonding molecular orbitals?

A.  $O_2$ 

B.  $O_2^{2-}$ 

 $\mathsf{C}.\,O_2^{\,+}$ 

D. None

#### **Answer: C**



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**61.** According to MOT, two atomic orbitals overlap resulting in the formation of molecular orbital formed. Number of atomic orbitals overlapping together is equal to the molecule orbital formed. The two atomic orbital thus formed by LCAO (linear combination of atomic orbital) in the phase or in the different phase are known as bonding and antibonding molecular orbitals respectively. The energy of bonding molecular orbital is lower than that of the pure atomic orbitals by an amount  $\Delta$ . This known as the stabilization energy. The enerby of antibonding molecular orbital in increased by  $\Delta'$  (destabilisation energy).

Q. Which among the following pairs contains both paramagnetic species.

- A.  $O_2^{2-}$  and  $N_2^{-}$
- $\mathsf{B}.\,O_2^- \;\;\mathrm{and}\;\; N_2$
- $C. O_2$  and  $N_2$
- $\mathsf{D}.\,O_2$  and  $N_2^-$

**62.** According to MOT, two atomic orbitals overlap resulting in the formation of molecular orbital formed. Number of atomic orbitals overlapping together is equal to the molecule orbital formed. The two atomic orbital thus formed by LCAO (linear combination of atomic orbital) in the phase or in the different phase are known as bonding and antibonding molecular orbitals respectively. The energy of bonding molecular orbital is lower than that of the pure atomic orbitals by an amount  $\Delta$ . This known as the stabilization energy. The enerby of antibonding molecular orbital in increased by  $\Delta'$  (destabilisation energy).

Q. Which of the following statement(s) is true:

A. Higher the bond order lesser the bond length.

B. Higher the bond order greater the bond length.

C. Higher the bond order lesser the bond energy.

D. Higher the bond order lesser the number of bonds

#### Answer: A



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**63.** According to MOT, two atomic orbitals overlap resulting in the formation of molecular orbital formed. Number of atomic orbitals overlapping together is equal to the molecule orbital formed. The two atomic orbital thus formed by LCAO (linear combination of atomic orbital) in the phase or in the different phase are known as bonding and antibonding molecular orbitals respectively. The energy of bonding molecular orbital is lower than that of the pure atomic orbitals by an amount  $\Delta$ . This known as the stabilization energy. The enerby of antibonding molecular orbital in increased by  $\Delta$ ' (destabilisation energy).

Q. How many nodal plane is present in  $\sigma_{s \text{ and } p}$  bonding molecular orbital ?

A. zero

C.2

D. 3

#### **Answer: A**



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**64.** According to MOT, two atomic orbitals overlap resulting in the formation of molecular orbital formed. Number of atomic orbitals overlapping together is equal to the molecule orbital formed. The two atomic orbital thus formed by LCAO (linear combination of atomic orbital) in the phase or in the different phase are known as bonding and antibonding molecular orbitals respectively. The energy of bonding molecular orbital is lower than that of the pure atomic orbitals by an amount  $\Delta$ . This known as the stabilization energy. The enerby of antibonding molecular orbital in increased by  $\Delta'$  (destabilisation energy).

Q. which of the following combination of orbitals is corrects?

$$\mathsf{D}^{(\mathsf{d})} \overset{\Diamond}{ } \stackrel{\Diamond}{-} \overset{\Diamond}{-} \overset{\Diamond}{ } \stackrel{\Diamond}{-} \overset{\Diamond}{-} \overset{\Diamond}{-$$

#### **Answer: C**



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**65.** According to MOT, two atomic orbitals overlap resulting in the formation of molecular orbital formed. Number of atomic orbitals overlapping together is equal to the molecule orbital formed. The two atomic orbital thus formed by LCAO (linear combination of atomic orbital) in the phase or in the different phase are known as bonding and antibonding molecular orbitals respectively. The energy of bonding molecular orbital is lower than that of the pure atomic orbitals by an amount  $\Delta$ . This known as the stabilization energy. The enerby of antibonding molecular orbital in increased by  $\Delta'$  (destabilisation

energy).

Q. Which of the following statements is not correct regarding bonding

Q. Which of the following statements is not correct regarding bonding molecular orbitals?

A. Bonding molecular orbitals possess energy than the atomic orbitals from which they are formed

B. Bonding molecular orbitals have low electron density the two nuclei

C. Electron in bonding molecular contributes to the attraction between atoms.

D. They are formed when the lobes of the conbining atomic orbitals have the same sign.

#### **Answer: B**



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**66.** According to MOT, two atomic orbitals overlap resulting in the formation of molecular orbital formed. Number of atomic orbitals

overlapping together is equal to the molecule orbital formed. The two atomic orbital thus formed by LCAO (linear combination of atomic orbital) in the phase or in the different phase are known as bonding and antibonding molecular orbitals respectively. The energy of bonding molecular orbital is lower than that of the pure atomic orbitals by an amount  $\Delta$ . This known as the stabilization energy. The enerby of antibonding molecular orbital in increased by  $\Delta'$  (destabilisation energy).

Q. If x-axis is the molecular axis, then  $\pi$ -moleulcar orbitals are formed by the overlap of :

- A.  $S+p_x$
- B.  $p_x+p_y$
- C.  $P_z+p_x$
- D.  $p_x+p_x$

#### Answer: C



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**67.** Polar covalent molecules exhibit dipole moment. Dipole moment is equal to the product of charge separation , q and the bond length d for the bond. Unit of dipole moment is debye. One debye is equal to  $10^{-18}$  esu cm.

Dipole moments is a vector quantity. It has both magnitude and direction. Hence, dipole moment of a molecule depends upon the relative orientation of the bond dipoles, but not on the polarity of bonds alone. A symmetrical structure shows zero dipole moment. Thus, dipole moment helps to predict the geometry of a molecules. Dipole moment values can be distinguish between cis- and trans- isomers, ortho, meta and pareforms of a substance, etc.

Q. Which is a polar molecule?

- A.  $XeF_4$
- $B.\,BF_3$
- C.  $I_2Cl_6$
- D.  $PCl_2F_3$



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**68.** Polar covalent molecules exhibit dipole moment. Dipole moment is equal to the product of charge separation , q and the bond length d for the bond. Unit of dipole moment is debye. One debye is equal to  $10^{-18}$  esu cm.

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Q. A diatomic molecule has a dipole moment of 1.2D. If the bond length is  $1.0 imes 10^{-8}$  cm, what fraction of charge does exist each atom?

- B.0.2
- C. 0.25
- D. 0.3

#### Answer: C



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**69.** Polar covalent molecules exhibit dipole moment. Dipole moment is equal to the product of charge separation , q and the bond length d for the bond. Unit of dipole moment is debye. One debye is equal to  $10^{-18}$  esu cm.

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forms of a substance, etc.

Q. Arrange the following compounds in increasing order of dipole moments, toluene (I), o-dichlorobenzene (II), m-dichlorobenzene (III) and p-dichlorobenzene (IV):

A. 
$$IV < I < II < III$$

$$\mathsf{B}.\,I < IV < II < III$$

$$\mathsf{C}.\,IV < I < III < II$$

$$\mathsf{D}.\,IV < II < I < III$$

### Answer: C



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**70.** Polar covalent molecules exhibit dipole moment. Dipole moment is equal to the product of charge separation , q and the bond length d for the bond. Unit of dipole moment is debye. One debye is equal to  $10^{-18}$  esu cm.

Dipole moments is a vector quantity. It has both magnitude and direction.

Hence, dipole moment of a molecule depends upon the relative orientation of the bond dipoles, but not on the polarity of bonds alone. A symmetrical structure shows zero dipole moment. Thus, dipole moment helps to predict the geometry of a molecules. Dipole moment values can be distinguish between cis- and trans- isomers, ortho, meta and pareforms of a substance, etc.

Q.  $\mu$  of the  $AX_4$  type of molecule is zero. The geometry of it can be :

A. tetrahedral

B. square planar

C. A or B

D. None of these

#### Answer: C



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71. Polar covalent molecules exhibit dipole moment. Dipole moment is equal to the product of charge separation, q and the bond length d for the bond. Unit of dipole moment is debye. One debye is equal to  $10^{-18}$  esu cm.

Dipole moments is a vector quantity. It has both magnitude and direction.

Hence, dipole moment of a molecule depends upon the relative orientation of the bond dipoles, but not on the polarity of bonds alone. A symmetrical structure shows zero dipole moment. Thus, dipole moment helps to predict the geometry of a molecules. Dipole moment values can be distinguish between cis- and trans- isomers, ortho, meta and pareforms of a substance, etc.

Q. Which of the following statement is correct regarding  $C_2H_2F_2$  molecule?

A. One isomer is polar, one is non-polar

B. Two isomers are polar, one is non-polar

C. Two isomers are planar, one is non-planar

D. Two isomers are only possible and planar

#### Answer: B



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72. In general boiling point of covalent compounds is affect due to increasing molecular weight and hydrogen bonding. Thus it is obversed that the boiling point of octanol is greater that water while that of methanol is lesser than water. The other properties which are considerably affected due to hydrogen bond formation are acidic nature, melting point, anamalous behaviour of water below  $4^{\circ}C$ , It is therefore concluded that hydrogen bonding palys a vutal role in explaning many observable facts.

Q. Which among the following has maximum boiling point?

A. 
$$CH_2-CH_2-(CH_2)_2-CH_2OH$$

B.  $CH_3-CH_2-CH_2-CH-CH_2-CH_3$ 

C.  $CH_3-CH_2-CH-CH_2-CH_3$ 
 $CH_3$ 

D.  $CH_3-CH_2-CH_2-CH_3$ 

#### Answer: A

73. In general boiling point of covalent compounds is affect due to increasing molecular weight and hydrogen bonding. Thus it is obversed that the boiling point of octanol is greater that water while that of methanol is lesser than water. The other properties which are considerably affected due to hydrogen bond formation are acidic nature, melting point, anamalous behaviour of water below  $4^{\circ}\,C$ ., It is therefore concluded that hydrogen bonding palys a vutal role in explaning many observable facts.

- Q. Which of the following statements is true?
  - A. The lattice structure of ice involves true covalent bond
  - B. The lattice structure of ice is the result of dipole-dipole interaction
  - C. The lattice structure of ice is the result of intra and inter-molecular hydrogen bond formation
  - D. The lattice structure of ice is only due to inter-molecular hydrogen

bonding

#### Answer: D



- **74.** In general boiling point of covalent compounds is affect due to increasing molecular weight and hydrogen bonding. Thus it is obversed that the boiling point of octanol is greater that water while that of methanol is lesser than water. The other properties which are considerably affected due to hydrogen bond formation are acidic nature, melting point, anamalous behaviour of water below  $4^{\circ}C$ ., It is therefore concluded that hydrogen bonding palys a vutal role in explaning many observable facts.
- Q.  $K_{a_1}$  of maleic acid is lesser than  $K_{a_1}$  of fumaric acid due to:
  - A. Intramolecular hydrogen bonding in the formed after one proton removal in fumaric acid
  - B. Intermolecular hydrogen bonding in the ion formed after one proton removal in maleic acid

C. Intramolecular hydrogen bonding in the ion formed after one proton removal in maleic acid

D. Intermolecular hydrogen bonding in the ion formed after one proton removal in fumaric acid

#### **Answer: C**



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**75.** The molecule in which an atom is associated with more than 8 electrons is known as hypervalent molecule and less than 8 electrons is known as hypovalent molecule. All hypervalent molecules must have  $p\pi-d\pi$  bonding but the molecules having back bonding back boding need not to have always  $p\pi-d\pi$  bondin.

Q.Which of the molecule is not hypovalent but complete its octet:

- A.  $AlI_3$
- B.  $AlCl_3$

- C.  $AlBr_3$
- D.  $AlF_3$

#### **Answer: C**



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**76.** The molecule in which an atom is associated with more than 8 electrons is known as hypervalent molecule and less than 8 electrons is known as hypovalent molecule. All hypervalent molecules must have  $p\pi-d\pi$  bonding but the molecules having back bonding back boding need not to have always  $p\pi-d\pi$  bondin.

Q. Which of the following is having complete octet:

- A.  $BeCl_2$  (dimer)
- B.  $BeH_2$  (dimer)
- C.  $BeH_2$  (s)
- D.  $BeCl_2$  (s)

#### **Answer: D**



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77. The molecule in which an atom is associated with more than 8 electrons is known as hypervalent molecule and less than 8 electrons is known as hypovalent molecule. All hypervalent molecules must have  $p\pi-d\pi$  bonding but the molecules having back bonding back boding need not to have always  $p\pi-d\pi$  bondin.

Q.Which of the following molecule is not having  $p\pi-d\pi$  bonding:

- A.  $SO_2$
- B.  $P_4O_{10}$
- $\mathsf{C}.\,PF_3$
- D.  $B_3N_3H_6$

#### **Answer: D**



## ONE OR MORE ANSWER IS/ARE CORRECT

1. Which is correct statement?

A. LiCl is more soluble in polar solvent (water ) than NaCl

B.  $K_{a_2}$  of fumaric acid is more than  $K_{a_2}$  of maleic acid

C. The O-O bond length in  $O_2[AsF_4]$  is shorter than  $KO_2$ 

D. In  $CF_2=C=CF_2$  molecule all the four flourine atoms are in the same plane

Answer: A::B::C



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**2.** Select correct statement(s) regarding  $\sigma$  and  $\pi$  bonds :

A.  $\sigma$ -bond lies on the line joining the nuclei of bonded atoms

B.  $\pi-$  electron cloud lies on either side to the line joining the nuclei

of bonded atoms

C.  $(2p\pi-3d\pi)\pi$ -bond is stronger than  $(2p\pi-2\pi)\pi$ -bond

 $\mbox{D.}\, \sigma\mbox{-bond}$  has primary effect to decide direction of covalent bond,

which  $\pi-$  bond has no primery effect in direction of bond.

#### Answer: A::B::C::D



**3.** Which of the following species is/are having  $\pi_{2p}$  as HOMO (highest occupied molecular orbital) ?

A. 
$$N_2^-$$

$$\operatorname{B.}O_2^{2\,+}$$

C. 
$$NO^+$$

D. 
$$B_2^+$$

## Answer: B::C::D



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4. Select correct order between given compounds.

A.  $COCl_2 > COF_2$  : XCX bond angle

B.  $NO_2F > NO_2Cl$  : ONO bond angle

C.  $SO_2F_2 < SOF_2$  : FSF bond angle

D.  $N_2F_2 < N_2(CH_3)_2$  : N-N bond length

#### Answer: A::B::D



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**5.** Select the correct statement(s) regarding  $BF_2NH_2$  molecule:

A. FBF bond angle  $\,< 120^{\circ}$ 

B. HNH bond angle  $\,>109^{\circ}\,28^{\,\prime}$ 

C. HNH bond angle  $\,< 109^{\circ}\,28$  '

D. FBF bond angle  $\,>120^\circ$ 

#### Answer: A::B



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- **6.** Correct statement (s) about dipole moment of  $R_3NO$  and  $R_3PO$  is /are:
  - A. dipole moment of  $R_3NO>\,$  dipole moment of  $R_3PO$
  - B. dipole moment of  $R_3NO<\,$  dipole moment of  $R_3PO$
  - C. Experimental dipole moment of  $R_3NO>\,$  Experimental dipole

moment of  $R_3PO$ 

D. Experimental dipole moment of  $R_3NO<\,$  Experimental dipole moment of  $R_3PO$ 

### Answer: A::D



**7.** In which of the following compounds observed bond angle is found to be greater than expected, but not due to back bonding.

- A.  $N(SiH_3)_3$
- B.  $N(CH_3)_3$
- $C.O(CH_3)_2$
- D.  $O(SiH_3)_2$

#### Answer: B::C



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**8.** Two compounds  $PX_2Y_3 \; ext{and} \; PX_3Y_2$  (Where P=phosphorous atom and

X, Y= monovalent atoms). If all 'X' atoms are replaced by 'Z' atoms and

electronegativity order is X>Y>Z. Then incorrect statement (s)

is/are:

A. The dipole moment of product obtained from  $PX_2Y_3$  is non-zero

B. The dipole moment of product obtained from  $PX_2Y_3$  is zero

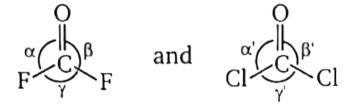
C. The dipole moment of product obtained from  $PX_{3}Y_{2}$  is zero

D. The dipole moment of product obtained from  $PX_3Y_2$  is non-zero

#### Answer: A::C



9. Corrrect order of bond angle in the given compounds is/are:



A. 
$$\gamma < \gamma$$

B. 
$$\gamma < \beta = \alpha$$

 $C. \alpha > \alpha'$ 

 $D.\beta > \beta$ 

Answer: A::B::C::D



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10. The correct statement(s) is /are:

A. Boiling point of m-hydroxybenzaldehyde is greater than o-

hydroxy benzalde hyde

B. Boiling point of  $CHCl_3$  is higher than  $CCl_4$ 

C. Melting point of  $BeCl_2$  is higher than  $BeF_2$ 

D. Boiling point of HF is greater than  $CH_3F$ 

Answer: A::B::D



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11. Consider of following reactions

$$CHF_{3}\stackrel{K_{a}}{\longrightarrow}CF_{3}^{\,-}+H^{\,+}$$

$$CHCl_3^- \stackrel{K_a^+}{\longrightarrow} CCl_3^- + H^+$$

Then regarding given reactions which of the following statement(s) is /are correct:

A. 
$$K_a > K^{\prime}_a$$

- B.  $CHF_3$  act as a stronger bronsted acid than  $CHCl_3$
- C.  $CCl_3^-$  is more stable than  $CF_3^{\,-}$
- D.  $CCl_3^-$  is weaker lewis base than  $CF_3^-$

#### **Answer: C::D**



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12. In which of the following molecules  $\mu$ (oberved) is found to be greater than  $\mu$  (theoretical):

A.  $POCl_3$ 

B. `##BLJ VKJ ORG CHE CO3 EO4 O13 OO1.png" width="30%">

 $\mathsf{C}.\,HNC$ 

D. `##BLJ VKJ ORG CHE CO3 EO4 O13 OO2.png" width="30%">

#### Answer: A::B::C::D



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**13.** Correct statement (s) regarding  $As(CH_3)F_2Cl_2$  molecule is/are:

A. Maximum three halogen atoms can lie in same plane

B. Both axial and equatorial plane may have equal number of atoms

C. As-Cl bond length is longer than As-F bond length .

D. Maximum five atoms can lie in equatorial plane.

# Answer: A::B::C::D



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**14.** Which of the following species is/are having 'N-N' bond order =2?

A.  $N_3^{\,-}$ 

B.  $N_2F_2$ 

 $\mathsf{C.}\,N_2O_4$ 

D.  $N_2O$ 

#### Answer: A::B



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**15.** Which of the following statements is correct?

A.  $ClF_3$  molecule is bent T shape

B. In  $SF_4$  molecule, F-S-F equatorial bond angle is  $103^\circ$  due to

lp-lp repulsion

C. In  $[Cl_4]^-$  molecular ion, Cl-I-Cl bond angle is  $90^\circ$ 

D. In  $OBr_2$ , the bond angle is less than  $OCl_2$ 

## Answer: A::B::C



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# 16. Which of the following species is/are not know?

A.  $FeI_3$ 

B.  $SH_6$ 

 $\mathsf{C}.\,PbI_4$ 

D.  $PI_5$ 

## Answer: A::B::C



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17. Select correct order between following compounds:

- A.  $NH_3>NF_3$ : bond angle
- B.  $NH_3>NF_3$  : dipole moment
- C.  $NH_3 > NF_3$  %s-character of lone pair
- D.  $NH_3>NF_3$  : reactivity towards lewis acid

Answer: A::B::D



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18. Which of the following is (are) V-shaped?

- A.  $S_3^{2\,-}$
- $\mathrm{B.}\,I_3^{\,-}$
- $\operatorname{C.}N_3^-$
- $\operatorname{D.}I_3^{\,+}$

### Answer: A::D



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19. Select correct order between given compounds.

- A.  $COCl_2 > COF_2$  :  $X\widehat{C}X$  bond angle
- B.  $NO_2F > NO_2Cl$  :  $\widehat{ONO}$  bond angle
- C.  $SO_2F_2 < SOF_2$  :  $F\widehat{S}F$  bond angle
- D.  $N_2F_2 < N_2(CH_3)_2$  : N-N bond length

#### Answer: A::B::D



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**20.** Which of the following equlibria would have highest and lowest value of  ${\cal K}_p$  at a commone temperature?

A.  $BeCO_3 
ightarrow BeO + CO_2$ 

B.  $CaCO_3 
ightarrow CaO + CO_2$ 

 $\mathsf{C}.\mathit{SrCO}_3 o \mathit{SrO} + \mathit{CO}_2$ 

D.  $BaCO_3 \rightarrow BaO + CO_2$ 

#### Answer: A::D



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21. Which of the following process is/are assciated with change of hybridisation of the underlined compounds?

A.  $Al(OH)_3$  pp. dissolved in NaOH

B.  $B_2H_6$  is dissolved in THF

C.  $SiF_4$  vapour is passed through liq. HF

D. solidifaction  $PCl_5$  vapour

# Answer: A::C::D

22. Which of the following are true?

A. van der waals forces are reponsible for the formation of molecular crytals

B. Branching lower the boiling points of isomeric organic due to decrease in van der waals forces

C. In graphite, van der waal forces act between the carbon layers

D. In diamond, van der waal forces act between the carbon layers

Answer: A::B::C



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23. Which of the following statement is incorrect?

A.  ${\cal O}_2$  is paramagnetic,  ${\cal O}_3$  is also paramagnetic

B.  $O_2$  is paramagnetic,  $O_2^{2-}$  is diamagnetic

C.  $B_2$  is paramagnetic,  $C_2$  is also paramagnetic

D. Different obervation is found in their bond length when

$$NO \rightarrow NO^+$$
 and  $CO \rightarrow CO^+$ 

#### **Answer: C**



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24. Which of the following statements are not correct?

A. All C-O bonds in  $CO_3^{2\,-}$  are equal but not in  $H_2CO_3$ 

B. All C-O bonds in  $HCO_2^-$  are equal but not in  $HCO_2H$ 

 ${
m C.}\ C-O$  bond length in  $HCO_2^-$  is longer than C-O bond length

in  $CO_3^{2-}$ 

D. C-O bond length in  $HCO_2^-$  and C-O bond length in  $CO_3^{2-}$ 

are equal

### **Answer: C::D**



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**25.** In the structure of  $H_2CSF_4$ , which of the following statement is/are correct?

A. Two  ${\cal C}-{\cal H}$  bonds are in the same plane of axial  $S-{\cal F}$  bonds

B. Two C-H bonds are in the same plane of equitorial S-F bonds

C. Total six atoms are in the plane

D. Equitorial S-F plane is perpendicular to the nodal plane of  $\pi$  — bonds

#### Answer: A::C::D



26. In which compound compounds vacant hybride orbitals take part in bonding:

A.  $B_2H_6$ 

B.  $Al_2Cl_6$ 

 $\mathsf{C}.\,C_2H_5Cl$ 

D.  $H_3BO_3$ 

### Answer: A::B



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A. Its molecule is linear

B. Symmetric N-O-N is a favoured structure as compared to

N-N-O skeleton

**27.** Which of the following is true for  $N_2O$ ?

C. Bond orders are fractional for  $N-N \ {
m and} \ N-O$  bonds

D. It is a neutral oxide

Answer: A::C::D



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**28.** Silane is more reactive than  $CH_4$  towards  $Nu^-$  substitution due to :

A. larger size of Si compared to C which facilitating the attack by nucleophile

B. Polarity of Si-H bonds is opposite to that of C-H bond

C. Availability of vacant 3d orbitals in case to form the reaction intermediate easily for nucleophilic attack

·

D. Si-H bond energy is lower than that of C-H bonds

Answer: A::B::C::D



**29.** Which of the following statement(s) is/are not correct for following compounds?

 $(I)SCl_2(OCH_3)_2$  and  $(II)SF_2(OCH_3)_2$ 

equitorial position in case of (II)

A.  $-OCH_3$  groups in both cases occupy the same position

B. Cl-atoms occupy equitorial position incase of (I) and F-atoms occupy

C. Cl-atoms occupy axial position in case of (II) & F-atoms occupy equitorial position in case of (II)

D. Cl and F- atoms occupy either axial or equitorial position in case of

(I) and (II) respectively

Answer: A::B::C



**30.** If  $N_B$  is the number of bonding electron and  $N_A$  is the number of antibonding electrons of a molecules. Then choose the incorrect statement(s) for the relationship ,  $N_B>N_A$ "

A. Molecule may be stable or unstable

B. Molecule may have any integral, frational or zero value of bond order

C. Molecule is only paramagnetic species

D. Molecule does not exist

## Answer: B::C::D



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**31.** Stepwise hydrolysis of  $P_4O_{10}$  takes place via formation of :

A. tetrametaphosphoric acid

B. tetrapolyphosphoric acid

C. pyrophosphoric acid

D. Orthophosphoric acid

Answer: A::B::C::D



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**32.** Select the correct statement(s) about the compound  $NO[BF_4]$  :

A. It has  $5\sigma \ {
m and} \ 2\pi \ {
m bond}$ 

B. Nitrogen-oxygen bond length is higher than nitric oxide (NO)

C. It is a diamagnetic species

D. B-F bond length in this compound is lower than in  $BF_3$ 

Answer: A::C



**33.** Which of the following molecule has as  $O_O$  bonds?

- A.  $H_2S_2O_8$
- $\mathsf{B.}\,H_2S_2O_7$
- C.  $H_2SO_5$
- D.  $H_2S_2O_6$

## Answer: A, C

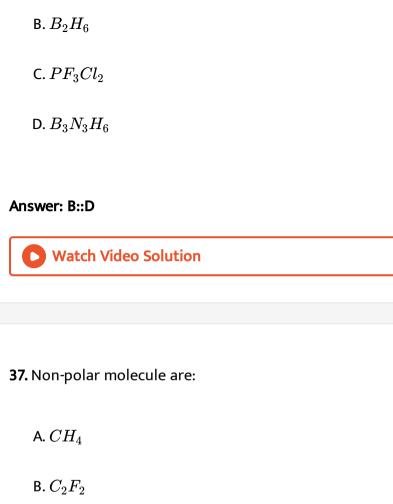


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**34.**  $CO_2$  molecule is not isostructural with :

- A.  $HgCl_2$
- $\mathsf{B.}\,SnCl_2$
- $\mathsf{C.}\,C_2H_2$
- D.  $NO_2$

Answer: B::D	
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<b>35.</b> Which of the following have a linear structure?	
A. $HgCl_2$	
B. $SnCl_2$	
C. $ICl_2^-$	
D. $CS_2$	
Answer: A::C::D	
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<b>36.</b> Which of the following compound (s) is/are non-polar?	
A. $NO_2$	



 $C. C_2F_4$ 

D.  $OF_2$ 

# Answer: A::B::C



**38.** Which of the following molecule species is/are having  $\pi_{2p}$  as H.O.M.O (highest occupied molecular orbital) :

- A.  $N_2^{\,-}$
- B.  $O_2^{2\,+}$
- $\mathsf{C.}\,NO^{\,+}$
- D.  $B_2^{\,+}$

# Answer: B::C::D



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**39.** Correct order of b.p.t is /are:

- A.  $H_2 < He$
- B.  $H_2 < D_2$
- $\mathsf{C.}\,H_2O < D_2O$
- D.  $NH_3 < SbH_3$

Answer: B::C::D



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**40.** Select correct statement(s) regarding  $\sigma$  and  $\pi$  bonds :

A.  $\sigma$ -bond lies on the line joining the nuclei of bonded atoms

B.  $\pi-$  electron cloud lies on either side to the line joining the nuclei of bonded atoms

C.  $(2p\pi-3d\pi)\pi$ -bond is stronger than  $(2p\pi-2\pi)\pi$ -bond

D.  $\sigma$ -bond has primary effect to decide direction of covalent bond, which  $\pi$  — bond has no primery effect in direction of bond.

Answer: A::B::C::D



**41.** Which of the following species is/are having  $\pi_{2p}$  as HOMO (highest occupied molecular orbital) ?

A. 
$$N_2^-$$

B.  $O_2^{2\,+}$ 

C. *NO* +

D.  $B_{2}^{+}$ 

# Answer: B::C::D



- - A.  $COCl_2 > COF_2$  : XCX bond angle

42. Select correct order between given compounds.

- B.  $NO_2F > NO_2Cl$  : ONO bond angle
- C.  $SO_2F_2 < SOF_2$  : FSF bond angle
- D.  $N_2F_2 < N_2(CH_3)_2$  : N-N bond length

## Answer: A::B::D



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**43.** Select the correct statement(s) regarding  $BF_2NH_2$  molecule:

A. FBF bond angle  $\,< 120^{\circ}$ 

B. HNH bond angle  $\,>109^{\circ}\,28\,{}^{\prime}$ 

C. HNH bond angle  $\,< 109^{\circ}\,28^{\,\prime}$ 

D. FBF bond angle  $\,>120^{\,\circ}$ 

#### Answer: A::B



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**44.** Correct statement (s) about dipole moment of  $R_3NO \ {
m and} \ R_3PO$  is

/are:

- A. dipole moment of  $R_3NO>\,$  dipole moment of  $R_3PO$
- B. dipole moment of  $R_3NO<\,$  dipole moment of  $R_3PO$
- C. Experimental dipole moment of  $R_3NO>$  Experimental dipole moment of  $R_3PO$
- D. Experimental dipole moment of  $R_3NO<\,$  Experimental dipole moment of  $R_3PO$

#### Answer: A::D



- **45.** In which of the following compounds observed bond angle is found to be greater than expected, but not due to back bonding.
  - A.  $N(SiH_3)_3$
  - B.  $N(CH_3)_3$
  - $\mathsf{C.}\,O(CH_3)_2$

D.  $O(SiH_3)_2$ 

Answer: B::C



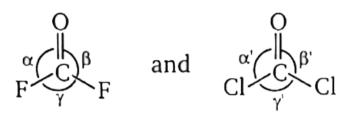
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- **46.** Two compounds  $PX_2Y_3$  and  $PX_3Y_2$  (Where P=phosphorous atom and X, Y= monovalent atoms). If all 'X' atoms are replaced by 'Z' atoms and electronegativity order is X>Y>Z. Then incorrect statement (s) is/are:
  - A. The dipole moment of product obtained from  $PX_2Y_3$  is non-zero
  - B. The dipole moment of product obtained from  $PX_2Y_3$  is zero
  - C. The dipole moment of product obtained from  $PX_3Y_2$  is zero
  - D. The dipole moment of product obtained from  $PX_3Y_2$  is non-zero

Answer: B::D



**47.** Corrrect order of bond angle in the given compounds is/are:



A. 
$$\gamma < \gamma$$
 '

B. 
$$\gamma < \beta = \alpha$$

$$\mathsf{C}.\, \alpha > \alpha'$$

D. 
$$\beta > \beta$$

Answer: A::B::C::D



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**48.** The correct statement(s) is /are :

A. Boiling point of m-hydroxybenzaldehyde is greater than o-

hydroxy benzalde hyde

B. Boiling point of  $CHCl_3$  is higher than  $\mathbb{C}l_4$ 

C. Melting point  $BeCl_2$  is higher than  $BeF_2$ 

D. Boiling point HF is greater than  $CH_3F$ 

### Answer: A::B::D



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**49.** Consider the following reversible chemical reactions:

$$A_2(g) + B_2(g) \stackrel{K_1}{\Longleftrightarrow} 2AB(g)....(1)$$

$$6AB(g) \stackrel{K_2}{\Longleftrightarrow} 3A_2(g) + 3B_2(g).....(2)$$

The relation between  $K_1$  and  $K_2$  is :

A. 
$$K_a > K^{\prime}_a$$

B.  $CHF_3$  act as a stronger bronsted acid than  $CHCl_3$ 

C.  $CCl_3^-$  is more stable than  $CF_3^-$ 

D.  $CCl_3^-$  is weaker lewis base than  $CF_3^-$ 

Answer: C::D



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**50.** In which of the following molecules  $\mu_{\rm exp}$  (observed dipole moment) is found to be greater than  $\mu_{\mu}$  (expected dipole moment)?

A.  $POCl_3$ 

В.

 $\mathsf{C}.\,HNC$ 

$$(d) \bigcirc F$$

D.

Answer: A::B::C::D



**51.** Correct statement (s) regarding  $As(CH_3)F_2Cl_2$  molecule is/are:

A. Maximum there halogen atoms can lie in same plane

B. Both axial and equatorial plane may have equal number of atoms

C. As-Cl bond length is longer than As-F bond length .

D. Maximum five atoms can lie in equatorial plane.

Answer: A::B::C::D



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**52.** Which of the following species are involved

A.  $N_3^-$ 

B.  $N_2F_2$ 

 $\mathsf{C.}\,N_2O_4$ 

D.  $N_2O$ 

#### Answer: A::B



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**53.** Which of the following statements is correct?

A.  $ClF_3$  molecule is bent 'T' shape

B. In  $SF_4$  molecule, F-S-F equatorial bond angle is  $103^\circ$  due to lp-lp repulsion

C. In  $[Cl_4]^-$  molecular ion, Cl-I-Cl bond angle is  $90^\circ$ 

D. In  $OBr_2$ , the bond angle is less than  $OCl_2$ 

#### Answer: A::C



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**54.** Which of the following species is/are not know?

A.  $FeI_3$ 

B.  $SH_6$ 

 $\mathsf{C}.\,PbI_4$ 

D.  $PI_5$ 

# Answer: A::B::C



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A.  $NH_3 > NF_3$ : bond angle

B.  $NH_3 > NF_3$ : dipole moment

C.  $NH_3 > NF_3$  %s-character of lone pair

D.  $NH_3>NF_3$  : reactivity towards lewis acid

55. Select correct order between following compounds:

# Answer: A::B::D



**56.** Which of the following is (are) V-shaped?

- A.  $S_3^{2\,-}$
- $\mathrm{B.}\,I_3^{\,-}$
- $\mathsf{C.}\,N_3^{\,-}$
- D.  $I_3^{\,+}$

#### Answer: A::D



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**57.** Which of the following equlibria would have highest and lowest value of  $K_p$  at a commone temperature?

A. 
$$BeCO_3 
ightarrow BeO + CO_2$$

B. 
$$CaCO_3 
ightarrow CaO + CO_2$$

C. 
$$SrCO_3 
ightarrow SrO + CO_2$$

D. 
$$BaCO_3 
ightarrow BaO + CO_2$$

#### Answer: A::D



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**58.** Which of the following process is/are assciated with change of hybridisation of the underlined compounds?

A.  $Al(OH)_3$  pp. dissolved in NaOH

B.  $B_2H_5$  is dissolved in THF

C.  $SiF_4$  vapour is passed through liq. HF

D. solidifaction  $PCl_5$  vapour

# Answer: A::C::D



**59.** Which of the following are true?

A. van der waals forces are reponsible for the formation of molecular crytals

B. Branching lower the boiling points of isomeric organic due to decrease in van der waals forces

C. In graphite, van der waal forces act between the carbon layers

D. In diamond, van der waal forces act between the carbon layers

### Answer: A::B::C



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**60.** Which of the following statement is incorrect?

A.  ${\cal O}_2$  is paramagnetic,  ${\cal O}_3$  is also paramagnetic

B.  $O_2$  is paramagnetic,  $O_2$  is diamagnetic

C.  $B_2$  is paramagnetic,  $C_2$  is also paramagnetic

D. Different obervation is found in their bond length when

$$NO o NO^+$$
 and  $CO o CO^+$ 

## Answer: A::B::D



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# 61. Which of the following statements are not correct?

A. All C-O bonds in  $CO_3^{2-}$  are equal but not in  $H_2CO_3$ 

B. All C-O bonds in  $HCO_2^-$  are equal but not in  $HCO_2H$ 

 ${
m C.}\ C-O$  bond length in  $HCO_2^-$  is longer than C-O bond length

in  $CO_3^{2-}$ 

D. C-O bond length in  $HCO_2^- \ {
m and} \ C-O$  bond length in  $CO_3^{2-}$ are equal

# Answer: C::D

**62.** In the structure of  $H_2CSF_4$ , which of the following statement is/are correct?

A. Two C-H bonds are in the same plane of axial S-F bonds

B. Two C-H bonds are in the same plane of equitorial S-F bonds

C. Total six atoms are in the plane

D. Equitorial S-F plane is perpendicular to the nodal plane of  $\pi$  — bonds

Answer: A::C::D



**63.** In which compound compounds vacant hybride orbitals take part in bonding:

- A.  $B_2H_6$
- B.  $Al_2Cl_6$
- $\mathsf{C}.\,C_2H_5Cl$ 
  - D.  $H_3BO_3$

## Answer: A::B



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**64.** Which of the following is true for  $N_2O$ ?

B. Symmetric N-O-N is a favoured structure as compared to

C. Bond orders are fractional for  $N-N \ {
m and} \ N-O$  bonds

- A. Its molecule is linear
- N-O-N skeleton
- D. It is a neutral oxide

Answer: A::C::D

**65.** Silane is more reactive than  $CH_4$  towards  $Nu^-$  substitution due to :

A. larger size of Si compared to C which facilitating the attack by nucleophile

B. Polarity of Si-H bonds is opposite to that of C-H bond

C. Availability of vacant 3d orbitals in case to form the reaction intermediate easily for nucleophilic attack

D. Si-H bond energy is lower than that of C-H bonds

Answer: A::B::C::D



**66.** Which of the following statement(s) is/are not correct for following compounds?

$$(I)SCl_2(OCH_3)_2$$
 and  $(II)SF_2(OCH_3)_2$ 

A.  $-OCH_3$  groups in both cases occupy the same position

B. Cl-atoms occupy equitorial position incase of (I) and F-atoms occupy

equitorial position in case of (II)

C. Cl-atoms occupy axial position in case of (II) & F-atoms occupy equitorial position in case of (II)

D. Cl and F- atoms occupy either axial or equitorial position in case of

(I) and (II) respectively

### Answer: A::B::C::D



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**67.** If  $N_B$  is the number of bonding electron and  $N_A$  is the number of antibonding electrons of a molecules. Then choose the incorrect statement(s) for the relationship ,  $N_B>N_A$ "

A. Molecule may be stable or unstable

B. Molecule may have any integral, frational or zero value of bond

order

C. Molecule is only paramagnetic species

D. Molecule does not exist

## Answer: B::C::D



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**68.** Stepwise hydrolysis of  $P_4O_{10}$  takes place via formation of :

A. tetrmetaphosphoric acid

B. tetrapolyphosphoric acid

C. pyrophosphoric acid

D. Orthophosphoric acid

Answer: A::B::C::D

**69.** Select the correct statement(s) about the compound  $NO[BF_4]$  :

A. It has  $5\sigma$  and  $2\pi$  bond

B. Nitrogen-oxygen bond length is higher than nitric oxide (NO)

C. It is a diamagnetic species

D. B-F bond length in this compound is lower than in  $BF_3$ 

# Answer: A::C



**70.** Which of the following molecule has as O-O bonds?

A.  $H_2S_2O_8$ 

 $\mathsf{B.}\,H_2S_2O_7$ 

 $\mathsf{C}.\,H_2SO_5$ 

D.	$H_2S_2O_6$

Answer: A::C



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71. Which of the following species in paramagnetic:

A.  $CN^{\,-}$ 

B.  $SnCl_2$ 

 $\mathsf{C}.\,C_2H_2$ 

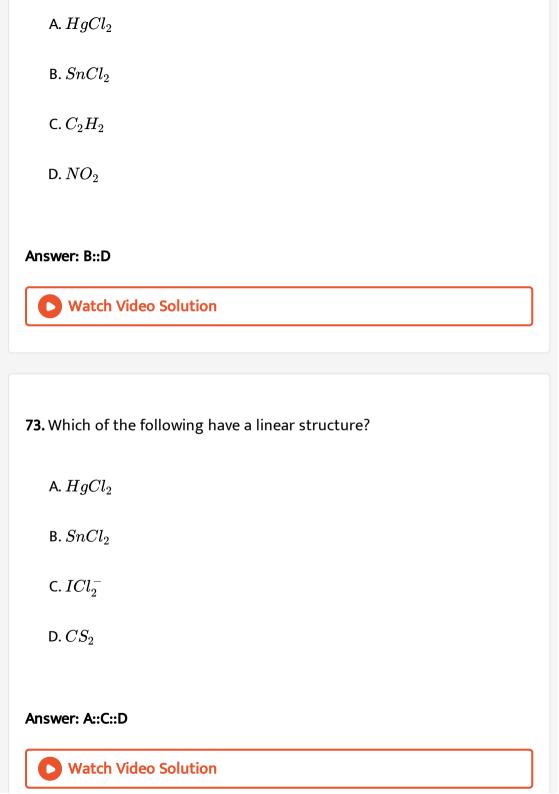
D.  $NO_2$ 

Answer: B::D



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**72.**  $CO_2$  molecule is not isostructural with :



<b>74.</b> Which of the following compound (s) is/are non-polar?
A. $NO_2$
B. $B_2H_6$
C. $PF_3Cl_2$
D. $B_3N_3H_6$
Answer: B::D
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<b>75.</b> Non-polar molecule are:
A. $CH_4$
B. $C_2F_2$
C. $C_2F_4$

D.  $OF_2$ 

Answer: A::B::C



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**76.** Which of the following molecule species is/are having  $\pi_{2p}$  as H.O.M.O (highest occupied molecular orbital) :

- A.  $N_2^{\,-}$
- B.  $O_2^{2\,+}$
- $\mathsf{C}.\,NO^+$
- D.  $B_2^+$

Answer: B::C::D



77. Correct order of b.p.t is /are:

A.  $H_2 < He$ 

 $\mathrm{B.}\,H_2 < D_2$ 

 $\mathsf{C.}\,H_2O < D_2O$ 

D.  $NH_3 < SbH_3$ 

#### Answer: B::C::D



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78. Incorrect order between following compounds is/are:

A.  $O(CH_3)_2 < O(SiH_3)_2$  : number of  $sp^3$ -hybride atoms

B.  $O(CH_3)_2 < O(SiH_3)_2$  : Bond angle w.r.t common atom

C.  $O(CH_3)_2 < O(SiH_3)_2$  :% s-character of hybrid orbital on central

atom

D.  $O(CH_3)_2 < O(SiH_3)_2$ : reactivity towards lewis acid

Answer: A::D



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#### MATCH THE COLUMN

1. Column-I and Column-II contains four entries each. Entries of Column -I are to be matched with some entries of Column-II. One or more than one entries of Column-I may have the matching with the same entries of Column-II.

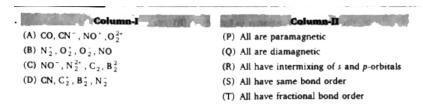
# Column-I

- (A) O(SiH<sub>3</sub>)<sub>2</sub>
- (B) Singlet CCl<sub>2</sub>
- (C) H<sub>4</sub>SiO<sub>4</sub>
- (D)  $B(NMe_2)_3$

#### Column-II

- (P) Hybridization of central atom does not change due to back bonding
- (Q) Bond angle of central atom increases due to combined effect of back bonding and steric factor
- (R) Anyone of  $t_{2g}$  d-orbital is involved in back
- bonding (S) Electron density on central
- decreases due to back bonding (T) Electron density on central
- increases due to back bonding







**3.** Column-I and Column-II contains four entries each. Entries of Column -I are to be matched with some entries of Column-II. One or more than one entries of Column-I may have the matching with the same entries of

#### Column-II.

#### Column-I (Reactions)

- (A) Formation of cation and anion by self ionization of iodine
- (B) Attack of hydroxide ion on boric acid
- (C)  $Ba^{2+}(aq) + SO_4^{2-}(aq) \longrightarrow BaSO_4 \downarrow \text{ (white ppt.)}$
- (D) MgSO<sub>4</sub>  $\xrightarrow{\text{T}>800^{\circ}\text{C}}$  MgO + SO<sub>2</sub> +  $\frac{1}{2}$ O<sub>2</sub>

# Column-II (Characteristics of final products)

- (P) Lone pair(s) is/are present at central atom
- (Q) d-orbital(s) involved in hybridization of central atom of either of product
- (R) d-orbital(s) not involved in hybridization of central atom of either of product
- (S)  $d\pi p\pi$  bond(s)
- (T) Planar covalent species is formed



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**4.** Column-I and Column-II contains four entries each. Entries of Column -I are to be matched with some entries of Column-II. One or more than one entries of Column-I may have the matching with the same entries of Column-II.

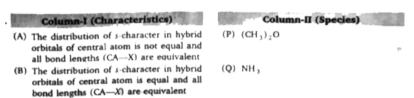
#### Column-I (Compounds)

- (A) H<sub>4</sub>SiO<sub>4</sub>
- (B)  $H_2SeO_4$
- (C) H<sub>3</sub>BO<sub>3</sub>
- (D) H<sub>2</sub>NBF<sub>2</sub>

#### Column-II (Characteristics)

- (P) Back bond
- (Q) Intermolecular hydrogen bond
- (R) Hypovalent compound
- (\$) Proton donor acid
- (T) Hypervalent compound







**6.** Column-I and Column-II contains four entries each. Entries of Column -I are to be matched with some entries of Column-II. One or more than one entries of Column-I may have the matching with the same entries of Column-II.

Column-I (Oxyacids)		Column-II (Oxidation state of S-atom)
(A) H <sub>2</sub> S <sub>4</sub> O <sub>6</sub>	(P)	+6
(B) H <sub>2</sub> S <sub>2</sub> O <sub>3</sub>	(Q)	+5
(C) H <sub>2</sub> SO <sub>5</sub>	(R)	0
(D) H <sub>2</sub> S <sub>2</sub> O <sub>5</sub>	(S)	-2
	(T)	+3



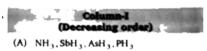
1000	Column-I	- Control K	Column-II
	NH <sub>4</sub> Cl	(P	Hydrogen bond
(B)	CuSO <sub>4</sub> ·5H <sub>2</sub> O	(Q	Co-ordinate bond
(C)	HNC	(R	Ionic bond
(D)	Liquid H <sub>2</sub> O <sub>2</sub>	(S)	Covalent bond



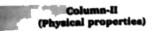
**8.** Column-I and Column-II contains four entries each. Entries of Column -I are to be matched with some entries of Column-II. One or more than one entries of Column-I may have the matching with the same entries of Column-II.







- (B) HI, HBr, HCl, HF
- (C) SnH4, GeH4, SiH4, CH4
- (D) H<sub>2</sub>O, H<sub>2</sub>Te, H<sub>2</sub>Se, H<sub>2</sub>S



- (P) Bond dipele moment
- (Q) Reducing property
- (R) Enthalpy of fusion
- (S) Boiling point



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10. Column-I and Column-II contains four entries each. Entries of Column -I are to be matched with some entries of Column-II. One or more than one entries of Column-I may have the matching with the same entries of Column-II.

#### Column-II (Bond angle) Column-I (Species) (A) NO; (P) 180° (B) NO. (O) 120° (C) NO<sub>2</sub> (R) 134° (D) NO; (S) 115° ) 109°



# (A) O(SiH<sub>3</sub>)<sub>2</sub> (P) (B) Singlet CCl<sub>2</sub> (Q) (C) H<sub>4</sub>SiO<sub>4</sub> (R) (D) B(NMe<sub>2</sub>)<sub>3</sub> (S)

# (P) Hybridization of central atom does not change due to back bonding (Q) Bond angle of central atom increases due to combined effect of back bonding and steric factor (R) Anyone of t<sub>2g</sub> d-orbital is involved in back bonding (S) Electron density on central atom decreases due to back bonding (T) Electron density on central atom increases due to back bonding



12. Column-I and Column-II contains four entries each. Entries of Column -I are to be matched with some entries of Column-II. One or more than one entries of Column-I may have the matching with the same entries of

#### Column-II.

#### Column-I

- (A) CO, CN -, NO ', O2+
- (B)  $N_2^+$ ,  $O_2^+$ ,  $O_2^-$ , NO
- (C) NO-, N2+, C2, B2
- (D) CN, C2, B2, N2

#### Column-II

- (P) All are paramagnetic
- (Q) All are diamagnetic
- (R) All have intermixing of s and p-orbitals
- (S) All have same bond order
- (T) All have fractional bond order



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**13.** Column-I and Column-II contains four entries each. Entries of Column -I are to be matched with some entries of Column-II. One or more than one entries of Column-I may have the matching with the same entries of Column-II.

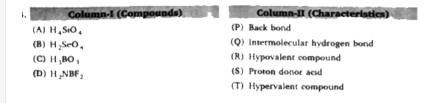
#### Column-I (Reactions)

- (A) Formation of cation and anion by self ionization of iodine
- (B) Attack of hydroxide ion on boric acid
- (C) Ba<sup>2\*</sup> (aq) + SO<sub>4</sub><sup>2-</sup> (aq)  $\longrightarrow$  BaSO<sub>4</sub>  $\downarrow$  (white ppt.)
- (D) MgSO<sub>4</sub>  $\xrightarrow{T>800^{\circ}C}$  MgO + SO<sub>2</sub> +  $\frac{1}{2}$ O<sub>2</sub>

# Column-II (Characteristics of final products)

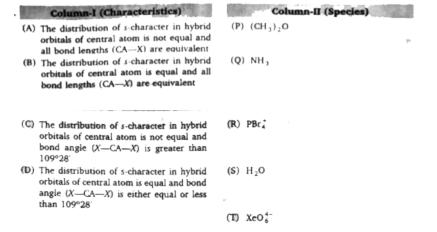
- (P) Lone pair(s) is/are present at central
- (Q) d-orbital(s) involved in hybridization of central atom of either of product
- (R) d-orbital(s) not involved in hybridization of central atom of either of product
- (S) dπ pπ bond(s)
- (T) Planar covalent species is formed







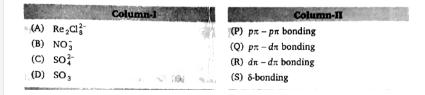
**15.** Column-I and Column-II contains four entries each. Entries of Column -I are to be matched with some entries of Column-II. One or more than one entries of Column-I may have the matching with the same entries of Column-II.





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**16.** Column-I and Column -II contains four entries each. Entries of Column-I are to be matched with some entries of Column-II. One or more than one entries of Column-I may have the matching with the same entries of Column-II.





Column-I (Oxyacids)		Column-II (Oxidation state of S-ato	m)
(A) H <sub>2</sub> S <sub>4</sub> O <sub>6</sub>	(P)	+6	
(B) $H_2S_2O_3$	(Q)	+5	
(C) H <sub>2</sub> SO <sub>5</sub>	(R)	0	
(D) $H_2S_2O_5$	(S)	-2	
	(T)	+3	



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**18.** Column-I and Column-II contains four entries each. Entries of Column -I are to be matched with some entries of Column-II. One or more than one entries of Column-I may have the matching with the same entries of Column-II.

	Column-I	K	Column-II
(A)	NH ₄Cl	(1	) Hydrogen bond
(B)	CuSO 4 - 5H 2O	((	() Co-ordinate bond
(C)	HNC	(F	() Ionic bond
(D)	Liquid H <sub>2</sub> O <sub>2</sub>	(5	6) Covalent bond



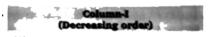




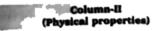
**20.** Column-I and Column-II contains four entries each. Entries of Column - I are to be matched with some entries of Column-II. One or more than one entries of Column-I may have the matching with the same entries of Column-II.

# Column-II (Pair of species) (A) PCI<sub>3</sub>F<sub>2</sub>.PCI<sub>2</sub>F<sub>3</sub> (B) BF<sub>3</sub> and BCI<sub>3</sub> (C) CO<sub>2</sub> and CN<sub>2</sub><sup>-2</sup> (D) C<sub>6</sub>H<sub>6</sub> and B<sub>3</sub>N<sub>3</sub>H<sub>6</sub> (Column-II (Identical Property in pairs of species) (P) Hybridisation of central atom (Q) Shape of molecule/ion (R) µ (dipole moment) (S) Total number of electrons





- (A) NH3, SbH3, AsH3, PH3
- (B) HI, HBr, HCl, HF
- (C) SnH4, GeH4, SiH4, CH4
- (D) H<sub>2</sub>O, H<sub>2</sub>Te, H<sub>2</sub>Se, H<sub>2</sub>S



- (P) Bond dipele moment
- (Q) Reducing property
- (R) Enthalpy of fusion
- (S) Boiling point



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#### **ASSERTION-REASON TYPE QUESTIONS**

**1.** Assertion :  $C_3O_2$  is non-polar molecule.

Reason: Terminal 'pi-bonds of the molecule are lying in different planes.

A. If assertion is true but the reason is false

B. If assertion is false but reason is true

C. If both assertion and reason are true and the reason is the correct explanation of assertion

D. If both assertion and reason are true and the reason is not the correct explanation of assertion

#### **Answer: B**



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**2.** Assertion:- If  $d_{x^2y^2}$  and py orbitals come close together along z-axis, then can form  $\pi$  — bond by sideways overlapping.

Reason:- Both orbitals do not have electron density along z-axis.

- A. If assertion is true but the reason is false
- B. If assertion is false but reason is true
- C. If both assertion and reason are true and the reason is the correct explanation of assertion

D. If both assertion and reason are true and the reason is not the correct explanation of assertion

#### **Answer: B**



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**3.** Assertion :  $BF_3$  undergoes in partial hydrolysis

Reason : Due to strong back bonding in  $BF_3$  only two flouride groups have come out on nucleaophili attack by  $H_2O$ .

- A. If assertion is true but the reason is false
- B. If assertion is false but reason is true
- C. If both assertion and reason are true and the reason is the correct
  - explanation of assertion
- D. If both assertion and reason are true and the reason is not the correct explanation of assertion

#### **Answer: A**



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- **4.** Assertion : The central catbon atom in  $F_2C=C=CF_2$  and both carbo atoms in  $F_2B-C=C-BF_2$  are sp-hybridized. Reasom molecules are planar.
  - A. If assertion is true but the reason is false
  - B. If assertion is false but reason is true
  - C. If both assertion and reason are true and the reason is the correct explanation of assertion
  - D. If both assertion and reason are true and the reason is not the correct explanation of assertion

#### **Answer: A**



**5.** Assertion : Formation of  ${PH_4^+}$  ion is relatively difficult in comparison

to  $NH_4^+$  ion.

Reason : Lone pair of phosphorus atom in  $PH_3$  resides in stereochemically inactive pure s-orbital.

A. If assertion is true but the reason is false

B. If assertion is false but reason is true

C. If both assertion and reason are true and the reason is the correct

explanation of assertion

D. If both assertion and reason are true and the reason is not the

correct explanation of assertion

#### **Answer: C**



**6.** Assertion : Bond dissociation energy of B-F bond in  $BF_3$  molecule is lower than C-F bond in  $CF_4$  molecule.

Reason: Atomic size of B-atom is larger than that of C-atom.

A. If assertion is true but the reason is false

B. If assertion is false but reason is true

C. If both assertion and reason are true and the reason is the correct

explanation of assertion

D. If both assertion and reason are true and the reason is not the correct explanation of assertion

#### **Answer: B**



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**7.** Assertion :  $PF_3$  is stronger lewis base than  $PH_3$ .

Reason : I.p. of P-atom in  $PF_3$  molecule is present in  $\mathit{sp}^3$ -hybrid orbital,

whereas l.p. of P-atom in  $PH_{\mathrm{3}}$  is present in almost pure s-orbital.

A. If assertion is true but the reason is false

B. If assertion is false but reason is true

C. If both assertion and reason are true and the reason is the correct explanation of assertion

D. If both assertion and reason are true and the reason is not the correct explanation of assertion

#### **Answer: B**



8. Assertion: NiO is less than CaO.

Reason : $Ni^{2+}$  is pseudo noble gas configuration cation whereas  $Ca^{2+}$  is noble gas configuration cation.

A. If assertion is true but the reason is false

- B. If assertion is false but reason is true
- C. If both assertion and reason are true and the reason is the correct explanation of assertion
- D. If both assertion and reason are true and the reason is not the correct explanation of assertion

#### **Answer: A**



- **9.** Assertion: When two gaseous OF molecules are allowed to cool, then they undergo dimerisation through O-atom.
- Reason: Dimerr form of OF molecule (i.e,  ${\cal O}_2 {\cal F}_2$ ) is having one peroxy linkage in its structure.
  - A. If assertion is true but the reason is false
  - B. If assertion is false but reason is true

C. If both assertion and reason are true and the reason is the correct

explanation of assertion

D. If both assertion and reason are true and the reason is not the correct explanation of assertion

#### **Answer: A**



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**10.** Assertion : bond dissociation energy of N-F bond in  $NF_3$  molecule is lower than that of in  $NCl_3$  molecule.

Reason : Interelectronic repulsion exists between small size N and F atoms N-F bond of  $N\!F_3$  molecule.

A. If assertion is true but the reason is false

B. If assertion is false but reason is true

C. If both assertion and reason are true and the reason is the correct

explanation of assertion

D. If both assertion and reason are true and the reason is not the correct explanation of assertion

#### **Answer: B**



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**11.** Assertion: If  $d_{x^2-y^2}$  and  $p_y$  orbitals come close together along z-axis, then they can form  $\pi$  — bond by sideways overlapping.

Reason: Both orbitals do not have electron density along z-axis.

- A. If assertion is true but the reason is false
- B. If assertion is false but reason is true
- C. If both assertion and reason are true and the reason is the correct
- D. If both assertion and reason are true and the reason is not the
  - correct explanation of assertion

explanation of assertion

#### **Answer: B**



**12.** Assertion : HOF bond angle is higher than HOCl.

Reason: Oxygen is more electronegative than chlorine.

A. If assertion is true but the reason is false

B. If assertion is false but reason is true

C. If both assertion and reason are true and the reason is the correct explanation of assertion

D. If both assertion and reason are true and the reason is not the correct explanation of assertion

#### **Answer: B**



13. Assertion: NaCl is more ionic than Nal.

Reason: Chlorine is more electronegative than iodine.

A. If assertion is true but the reason is false

B. If assertion is false but reason is true

explanation of assertion

C. If both assertion and reason are true and the reason is the correct

D. If both assertion and reason are true and the reason is not the correct explanation of assertion

#### **Answer: D**



**14.** Assertion :  $PbI_4$  doesn't exist and converts into  $PbI_2$  and  $I_2$  spontaneously at room temperature but  $PbCl_4$  needs heatin to convert into  $PbCl_2$  and  $Cl_2$ .

Reason :  $Pb^{2+}$  is more stable than  $Pb^{4+}$  due to inert pair effect.

- A. If assertion is true but the reason is false
- B. If assertion is false but reason is true
- C. If both assertion and reason are true and the reason is the correct explanation of assertion
- D. If both assertion and reason are true and the reason is not the correct explanation of assertion

#### **Answer: D**



- **15.** Statement-1 : Dipole moment of  $NF_3$  is less than that of  $NH_3$ .
- Statement-2 : Polarity of N-F bond is less than that of N-H bond.
  - A. If assertion is true but the reason is false
  - B. If assertion is false but reason is true

C. If both assertion and reason are true and the reason is the correct explanation of assertion

D. If both assertion and reason are true and the reason is not the correct explanation of assertion

#### Answer: A



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**16.** Assertion: Solubility of n-orbital of n-orbital in water decrease with increase in molecular weight.

Reason: The hydrophobic nature of alkyl chain increase.

- A. If assertion is true but the reason is false
- B. If assertion is false but reason is true

explanation of assertion

C. If both assertion and reason are true and the reason is the correct

D. If both assertion and reason are true and the reason is not the correct explanation of assertion

#### **Answer: C**



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**17.** Assertion : The unpaired electron of  $CH_3^{ullet}$  free radical occupies poribtal.

 ${\sf Reason}: CH_3^{\:\bullet\:} \; {\sf possesses} \; sp^2 \text{-hybridisation}.$ 

- A. If assertion is true but the reason is false
- B. If assertion is false but reason is true
- C. If both assertion and reason are true and the reason is the correct
- explanation of assertion
- D. If both assertion and reason are true and the reason is not the correct explanation of assertion

#### **Answer: C**



**18.** Assertion: Nitrogen is unreactive at room temperature but becomes reactive at elevated temperature (on heating or in the presence of catalysts).

Reason: In nitrogen molecule, there is extensive delocalisation of electrons

- A. If assertion is true but the reason is false
- B. If assertion is false but reason is true
- C. If both assertion and reason are true and the reason is the correct explanation of assertion
- D. If both assertion and reason are true and the reason is not the correct explanation of assertion

#### Answer: A

**19.** Assertion: The p-isomer of dichlorobenzene has higher melting point than o- and m-isomer.

Reason: p-isomer is symmetrical and thus shows more closely packed strucure.

A. If assertion is true but the reason is false

B. If assertion is false but reason is true

C. If both assertion and reason are true and the reason is the correct

explanation of assertion

D. If both assertion and reason are true and the reason is not the correct explanation of assertion

**Answer: C** 



**20.** Assertion (A):  $Na_2SO_4$  is soluble in water while  $BaSO_4$  is insoluble.

Reason (R ): Latice enthalpy of  $BaSO_4$  exceeds its hydration enthalpy.

A. If assertion is true but the reason is false

B. If assertion is false but reason is true

C. If both assertion and reason are true and the reason is the correct

explanation of assertion

D. If both assertion and reason are true and the reason is not the correct explanation of assertion

#### **Answer: C**



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**21.** Statement 1:  $N_2$  and  $NO^+$  are both dia-magnetic.

Statement 2:  $NO^+$  is isoelectronic with  $N_2$ 

A. If assertion is true but the reason is false

- B. If assertion is false but reason is true
- C. If both assertion and reason are true and the reason is the correct
  - explanation of assertion
- D. If both assertion and reason are true and the reason is not the correct explanation of assertion

#### Answer: D



- **22.** Assertion: Bond order can assume any value number including zero.
- Reason :Higher the bond order ,shorter is bond length and greater is bond energy.
  - A. If assertion is true but the reason is false
  - B. If assertion is false but reason is true
  - C. If both assertion and reason are true and the reason is the correct
    - explanation of assertion

D. If both assertion and reason are true and the reason is not the correct explanation of assertion

#### **Answer: D**



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**23.** Assertion:  $C_3O_2$  has linear structure.

Reason: Each carbon atom in  $C_3 O_2$  in  $\mathit{sp}$  -hydridized.

- A. If assertion is true but the reason is false
- B. If assertion is false but reason is true
- C. If both assertion and reason are true and the reason is the correct
  - explanation of assertion
- D. If both assertion and reason are true and the reason is not the correct explanation of assertion

## Answer: C

**24.** Assertion In case of  $(CH_3)_3N$  geometry is pyramidal but in case of  $(SiH_3)_3N$  it is planar

Reasoning The maximum covalency of  $\operatorname{Si}$  is  $\operatorname{six}$  but that of C is four .

A. If assertion is true but the reason is false

B. If assertion is false but reason is true

C. If both assertion and reason are true and the reason is the correct

D. If both assertion and reason are true and the reason is not the correct explanation of assertion

**Answer: C** 



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explanation of assertion

**25.** Assertion: Superoxides of alkali metals are paramagnetic.

Reason: Superoxides contain  $O_2^-$  ion which has one unpaired electron.

A. If assertion is true but the reason is false

B. If assertion is false but reason is true

C. If both assertion and reason are true and the reason is the correct

explanation of assertion

D. If both assertion and reason are true and the reason is not the correct explanation of assertion

# **Answer: C**



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**26.** Assertion : The  $HF_2^-$  ion exists in the solid state & also in liquid state but not in aqueous state.

Reason : The magniture of hydrogen bonds among HF molecules is weaker than that in between  $HF \ {
m and} \ H_2O.$ 

A. If assertion is true but the reason is false

B. If assertion is false but reason is true

explanation of assertion

C. If both assertion and reason are true and the reason is the correct

D. If both assertion and reason are true and the reason is not the correct explanation of assertion

# **Answer: C**



**27.** Assertion: If  $d_{x^2-y^2}$  and  $p_y$  orbitals come close together along z-axis, then they can form  $\pi$  — bond by sideways overlapping.

Reason: Both orbitals do not have electron density along z-axis.

A. If assertion is true but the reason is false

B. If assertion is false but reason is true

C. If both assertion and reason are true and the reason is the correct

D. If both assertion and reason are true and the reason is not the

#### **Answer: B**



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explanation of assertion

correct explanation of assertion

**28.** Assertion : The H-bond present in  $NH_3$  dissolved in water is best

represented by :  $H- \underset{|}{N}-H-- \underset{|}{O}-H$  and not by

Reason : The  ${\cal O}-{\cal H}$  bond polarity is more compared to that  ${\cal N}-{\cal H}$  bond.

A. If assertion is true but the reason is false

- B. If assertion is false but reason is true
- C. If both assertion and reason are true and the reason is the correct explanation of assertion
- D. If both assertion and reason are true and the reason is not the correct explanation of assertion

#### **Answer: B**



**29.** Assertion :  $C_3O_2$  is non-polar molecule.

Reason: Terminal `pi-bonds of the molecule are lying in different planes.

- A. If assertion is true but the reason is false
- B. If assertion is false but reason is true

explanation of assertion

C. If both assertion and reason are true and the reason is the correct

D. If both assertion and reason are true and the reason is not the correct explanation of assertion

#### **Answer: B**



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**30.** Assertion : If  $d_{x^2-y^2}$  and  $p_y$  orbitals come close together along z-axis, they can form  $\pi$  — bond by sideways overlapping.

Reason: Both oritals do not electron density along z-axis.

- A. If assertion is true but the reason is false
- B. If assertion is false but reason is true
- C. If both assertion and reason are true and the reason is the correct
- D. If both assertion and reason are true and the reason is not the
  - correct explanation of assertion

explanation of assertion

#### **Answer: B**



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**31.** Assertion :  $BF_3$  undergoes in partial hydrolysis

Reason : Due to strong back bonding in  $BF_3$  only two flouride groups have come out on nucleaophili attack by  ${\cal H}_2O$ .

A. If assertion is true but the reason is false

B. If assertion is false but reason is true

C. If both assertion and reason are true and the reason is the correct explanation of assertion

D. If both assertion and reason are true and the reason is not the correct explanation of assertion

# **Answer: A**



**32.** Assertion : The central catbon atom in  $F_2C=C=CF_2$  and both carbo atoms in  $F_2B-C\stackrel{-}{=}C-BF_2$  are sp-hybridized.

Reasom molecules are planar.

- A. If assertion is true but the reason is false
- B. If assertion is false but reason is true
- C. If both assertion and reason are true and the reason is the correct
  - explanation of assertion
- D. If both assertion and reason are true and the reason is not the correct explanation of assertion

# **Answer: A**



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**33.** Assertion : Formation of  $PH_4^+$  ion is relatively difficult in comparison to  $NH_4^+$  ion.

Reason : Lone pair of phosphorus atom in  $PH_3$  resides in stereochemically inactive pure s-orbital.

A. If assertion is true but the reason is false

B. If assertion is false but reason is true

explanation of assertion

C. If both assertion and reason are true and the reason is the correct

D. If both assertion and reason are true and the reason is not the correct explanation of assertion

# **Answer: C**



**34.** Assertion : Bond dissociation energy of B-F bond in  $BF_3$  molecule

is lower than C-F bond in  $CF_4$  molecule.

A. If assertion is true but the reason is false

Reason: Atomic size of B-atom is larger than that of C-atom.

B. If assertion is false but reason is true

C. If both assertion and reason are true and the reason is the correct

explanation of assertion

D. If both assertion and reason are true and the reason is not the correct explanation of assertion

#### Answer: B



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**35.** Assertion :  $PF_3$  is stronger lewis base than  $PH_3$ .

Reason : I.p. of P-atom in  $PF_3$  molecule is present in  $sp^3$ -hybrid orbital, whereas l.p. of P-atom in  $PH_3$  is present in almost pure s-orbital.

A. If assertion is true but the reason is false

B. If assertion is false but reason is true

C. If both assertion and reason are true and the reason is the correct

explanation of assertion

D. If both assertion and reason are true and the reason is not the correct explanation of assertion

#### **Answer: B**



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36. Assertion: NiO is less than CaO.

Reason  $:Ni^{2+}$  is pseudo noble gas configuration cation whereas  $Ca^{2+}$  is noble gas configuration cation.

- A. If assertion is true but the reason is false
- B. If assertion is false but reason is true
- C. If both assertion and reason are true and the reason is the correct explanation of assertion
- D. If both assertion and reason are true and the reason is not the correct explanation of assertion

#### Answer: A



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**37.** Assertion : When two gaseous OF molecules are allowed to cool, than they undergoes dimerisation through O-atom.

Reason : Dimer form of OF molecule (i.e.,  ${\cal O}_2 {\cal F}_2$ ) is having one peroxy linkage in its structure.

A. If assertion is true but the reason is false

B. If assertion is false but reason is true

C. If both assertion and reason are true and the reason is the correct

explanation of assertion

D. If both assertion and reason are true and the reason is not the correct explanation of assertion

# Answer: A



**38.** Assertion : bond dissociation energy of N-F bond in  $NF_3$  molecule is lower than that of in  $NCl_3$  molecule.

Reason : Interelectronic repulsion exists between small size N and F atoms N-F bond of  $NF_2$  molecule.

A. If assertion is true but the reason is false

B. If assertion is false but reason is true

C. If both assertion and reason are true and the reason is the correct explanation of assertion

D. If both assertion and reason are true and the reason is not the correct explanation of assertion

# **Answer: B**



**39.** Assertion: If  $d_{x^2-y^2}$  and  $p_y$  orbitals come close together along z-axis,

then they can form  $\pi-\,$  bond by sideways overlapping.

Reason: Both orbitals do not have electron density along z-axis.

A. If assertion is true but the reason is false

B. If assertion is false but reason is true

C. If both assertion and reason are true and the reason is the correct

explanation of assertion

D. If both assertion and reason are true and the reason is not the correct explanation of assertion

# **Answer: B**



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**40.** Assertion :  $H_2S$  is stronger acid than  $PH_3$ .

Reason : S is more electronegative than P, conjugate base $HS^-$  is

more stable than  $H_2P^{\,-}$  .

A. If assertion is true but the reason is false

B. If assertion is false but reason is true

C. If both assertion and reason are true and the reason is the correct explanation of assertion

D. If both assertion and reason are true and the reason is not the correct explanation of assertion

# **Answer: B**



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**41.** Assertion: NaCl is more ionic than Nal.

Reason: Chlorine is more electronegative than iodine.

A. If assertion is true but the reason is false

B. If assertion is false but reason is true

C. If both assertion and reason are true and the reason is the correct

explanation of assertion

D. If both assertion and reason are true and the reason is not the correct explanation of assertion

#### Answer: D



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**42.** Assertion :  $PbI_4$  doesn't exist and converts into  $PbI_2$  and  $I_2$  spontaneously at room temperature but  $PbCl_4$  needs heatin to convert into  $PbCl_2$  and  $Cl_2$ .

Reason :  $Pb^{2+}$  is more stable than  $Pb^{4+}$  due to inert pair effect.

A. If assertion is true but the reason is false

B. If assertion is false but reason is true

C. If both assertion and reason are true and the reason is the correct

explanation of assertion

D. If both assertion and reason are true and the reason is not the correct explanation of assertion

#### **Answer: D**



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**43.** Assertion : Dipole moment of  $NF_3$  is less than that of  $NH_3$ .

Reason : polarity of N-F bond is less than that of N-H bond.

A. If assertion is true but the reason is false

B. If assertion is false but reason is true

C. If both assertion and reason are true and the reason is the correct

explanation of assertion

D. If both assertion and reason are true and the reason is not the correct explanation of assertion

# Answer: A

44. Assertion: Solubility of n-orbital of n-orbital in water decrease with increase in molecular weight.

Reason: The hydrophobic nature of alkyl chain increase.

A. If assertion is true but the reason is false

B. If assertion is false but reason is true

explanation of assertion

C. If both assertion and reason are true and the reason is the correct

D. If both assertion and reason are true and the reason is not the correct explanation of assertion

Answer: C



**45.** Assertion : The unpaired electron of  $CH_3^{\bullet}$  free radical occupies poribtal.

 ${\sf Reason}: CH_3^{\:\bullet\:} \; {\sf possesses} \; sp^2 \hbox{-hybridisation}.$ 

A. If assertion is true but the reason is false

B. If assertion is false but reason is true

C. If both assertion and reason are true and the reason is the correct explanation of assertion

D. If both assertion and reason are true and the reason is not the correct explanation of assertion

# **Answer: C**



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46. Assertion: Nitrogen in inactive at room temperature but becomes reactive at elevated temperature (on heating or in the presence of catalyst).

Reason: In nitrogen molecule, there is delocalization electrons.

A. If assertion is true but the reason is false

B. If assertion is false but reason is true

C. If both assertion and reason are true and the reason is the correct explanation of assertion

D. If both assertion and reason are true and the reason is not the correct explanation of assertion

# **Answer: A**



**47.** Assertion The p - isomer of dichlorobenzene has higher m.p than o-

Reason p-isomer is symmetrical and thus shows more closely packed structure.

- A. If assertion is true but the reason is false
- B. If assertion is false but reason is true
- C. If both assertion and reason are true and the reason is the correct explanation of assertion
- D. If both assertion and reason are true and the reason is not the correct explanation of assertion

# **Answer: C**



- **48.** Assertion (A):  $Na_2SO_4$  is soluble in water while  $BaSO_4$  is insoluble.
- Reason (R ): Latice enthalpy of  $BaSO_4$  exceeds its hydration enthalpy.
  - A. If assertion is true but the reason is false
  - B. If assertion is false but reason is true

C. If both assertion and reason are true and the reason is the correct explanation of assertion

D. If both assertion and reason are true and the reason is not the correct explanation of assertion

#### **Answer: C**



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**49.** Assertion :  $N_2$  and  $NO^+$  both are diamagnetic substances.

Reason :  $NO^+$  is isoelectronic with  $N_2$ .

A. If assertion is true but the reason is false

B. If assertion is false but reason is true

C. If both assertion and reason are true and the reason is the correct explanation of assertion

D. If both assertion and reason are true and the reason is not the correct explanation of assertion

#### **Answer: D**



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**50.** Assertion: Bond order can assume any value number including zero.

Reason :Higher the bond order ,shorter is bond length and greater is bond energy.

- A. If assertion is true but the reason is false
- B. If assertion is false but reason is true
- C. If both assertion and reason are true and the reason is the correct
  - explanation of assertion
- D. If both assertion and reason are true and the reason is not the correct explanation of assertion

#### **Answer: D**



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**51.** Assertion:  $C_3O_2$  has linear structure.

Reason: Each carbon atom in  $C_3O_2$  in  $\mathit{sp}$  -hydridized.

A. If assertion is true but the reason is false

B. If assertion is false but reason is true

C. If both assertion and reason are true and the reason is the correct explanation of assertion

D. If both assertion and reason are true and the reason is not the correct explanation of assertion

# **Answer: C**



**52.** Assertion : H-bonding occurs in  ${\cal H}_2{\cal O}$  due to larger size of O-atom.

Reason: The size of O-atom is larger than H-atom.

A. If assertion is true but the reason is false

B. If assertion is false but reason is true

C. If both assertion and reason are true and the reason is the correct

explanation of assertion

D. If both assertion and reason are true and the reason is not the correct explanation of assertion

# Answer: A



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**53.** Assertion :  $(CH_3)_3N$  geometry is pyramidal but in case  $(SiH_3)_3N$  it is planar.

Reason: The maximum covalency of Si is six but that of C is four.

A. If assertion is true but the reason is false

B. If assertion is false but reason is true

C. If both assertion and reason are true and the reason is the correct

explanation of assertion

D. If both assertion and reason are true and the reason is not the correct explanation of assertion

#### **Answer: C**



**54.** Assertion: Supeoxides of alkali matels are paramagnetic.

Reason : Superoxides conatain the ion  ${\cal O}_2^-$  which has one unpaired electron in its anti-bonding molecule orbital.

A. If assertion is true but the reason is false

B. If assertion is false but reason is true

C. If both assertion and reason are true and the reason is the correct

explanation of assertion

D. If both assertion and reason are true and the reason is not the correct explanation of assertion

#### **Answer: C**



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**55.** Assertion : The  $HF_2^-$  ion exists in the solid state & also in liquid state but not in aqueous state.

Reason : The magniture of hydrogen bonds among HF molecules is weaker than that in between HF and  $H_2O$ .

- A. If assertion is true but the reason is false
- B. If assertion is false but reason is true
- C. If both assertion and reason are true and the reason is the correct
- D. If both assertion and reason are true and the reason is not the
  - correct explanation of assertion

explanation of assertion

#### **Answer: C**



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**56.** Assertion : If  $d_{x^2-y^2}$  and  $p_y$  orbitals come close together along z-axis, they can form  $\pi$  — bond by sideways overlapping.

Reason: Both oritals do not electron density along z-axis.

A. If assertion is true but the reason is false

B. If assertion is false but reason is true

C. If both assertion and reason are true and the reason is the correct explanation of assertion

D. If both assertion and reason are true and the reason is not the correct explanation of assertion

# **Answer: B**



**57.** Assertion : The H-bond present in  $NH_{\rm 3}$  dissolved in water is best

represented by :  $H- \underset{|}{N}- H- - \underset{|}{O}- H$  and not by

Reason : The  ${\cal O}-{\cal H}$  bond polarity is more compared to that  ${\cal N}-{\cal H}$  bond.

A. If assertion is true but the reason is false

B. If assertion is false but reason is true

C. If both assertion and reason are true and the reason is the correct

explanation of assertion

D. If both assertion and reason are true and the reason is not the

correct explanation of assertion

# **Answer: B**



# SUBJECTIVE PROBLEMS

1. There are two groups of compounds A and B. Groups A contains three compounds  $Px_4, Qy_3, Rz_2$ . Groups B also contains three compounds  $Sx_4, Ty_3, Uz_2$ . Hybridization of each central atom of group A compounds is same as that of iodine in  $IBrCl^-$  while in group B compounds it is same as that of iodine  $IBrCl^+$ . Substituents X, Y and Z exhibit covalency of one in ground state. Then find the value of x/y.

Where, x and y are total number of lone pair present at central atoms of compounds of group A and B respectively.



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**2.** Consider the following three compounds  $(i)AX_{2n}^{n-}$ ,  $(ii)AX_{3n}$  and (ii) $AX_{4n}^{n+}$ , where central atom A is 15th group element and their maximum covalency is 3n. If total number of proton in surrounding atom X is n and value of n is one, then calculate value of  $x^3+y^2+z^2$  . (where x, y and z are total number of lone pair at central atom in compounds (i), (ii) and (iii) respectively.



**3.** When  $B_2H_4$  is allowed to react with following lewis bases, then how many given lewis bases form adduct through symmetrical cleavage of  $B_2H_6$ )?

 $NH_3,\,MeNH_3$ , pyridine, CO, T.H.F.,  $PH_3,\,PF_3,\,Me_3N,\,Me_3NH$ 



**4.** Consider the following elements A, B, C and D and their outer electronic configurtions are  $ns^2np^1$ ,  $ns^2np^3$ ,  $ns^2np^4$  and  $ns^2np^5$  respectively. Element E also has same outer electronic configuration like D but shows only single oxidation state(-1). If element A, B, C and D belong to same period as that of sodium. Consider the following compounds. (i) $CE_4$  (ii) $BD_2E_3$  (iii) $DE_3$  (iv) $DE_2$  (v) $BD_3E_2$  (vi) $C_2E_2$  (vii)DE (viii) $A_2D_6$ .

Then calculate the value of x+y, (where x and y are total number of polar and non-polar compounds).



- **5.** Consider following four compounds:
- (i)  $C_x O_y$
- (ii)  $C_x O_{y+1}$
- (iii)  $C_{x\,+\,2}O_{y\,+\,1}$  and (iv)  $C_{x\,+\,11}O_{y\,+\,8}$ ,
- if "x=y=1", then calculate the vauue of |p-q|, where p and q are total number of  $sp^2$  and hybridized carbon atoms respectively in given four compounds.



**6.** Calculate expression (x+y+z) for diatomic molecules.

where x=Total number of singly occupied molecular orbital (SOMO) in  ${\cal O}_2$ . y=Total number of singly occupied molecular orbital (SOMO) in  ${\cal B}_2$ 

z=Total number of singly occupied molecular orbital (SOMO) in NO.

**7.** If Hund rule violate, then find the total number of species among following which whill be dimagnetic:

$$B_2, O_2, N_2^-, C_2, NO, OF, N_2^{2-}, BN$$



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# 8. Consider the following table

Compounds (X are monovalent surrounding atoms)		Central atoms(A to B) belong to group	Characteristics of compounds	Number of lone pair(s) at central atom
(i)	AX 2	16	Planar and polar	m <sub>1</sub>
(ii)	BX n2	15	Trigonal pyramidal	m <sub>2</sub>
(iii)	cx.,	14	Zero dipole moment	m 3
(iv)	DX n4	13	All X — D — X bond angle are 120°	m <sub>4</sub>

Than calculate value of experssion 
$$\left|rac{n_1+n_2+n_3+n_4}{m_1+m_2+m_3+m_4}
ight|^2$$
 .



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**9.** Total number of species among following, in which bond angle is equal to or less than  $109^{\circ}\,28'$  and also they act as lewis base :

 $NH_3, NMe_3, O(SiH_3)_2, ICl_4^-, XeO_3, BF_2Cl, SiF_4, AsH_3, SO_2F_2$ 



**10.** Total number of unpaired electrons(s) present in both cationic and anionic part of compound  $O_2[PtF_6]$ .



**11.** Total number of species which has/ have symmetrical electronic distribution in their HOMO and also paramagnetic.

$$N_2^+, O_2^{2-}, C_2, O_2, B_2, C_2^{2-}, N_2^{2-}.$$



of effective back bond.

12. Total number of molecules, in which each covalent bond is comprised

$$BF_3, N(SiH_3)_3, PF_3, POF_3, B(OH)_3, O(SiH_3)_2, BFH_2, BF_2(NH_2), N_3$$
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**13.** Total number of angle in  $SeCl_4$  which are less than  $90^{\circ}$ .



\_\_\_\_



**14.** Consider the following species  $O(Me)_2, N(SiH_3)_3, CO, O(SiH_3)_2, \mathbb{C}l_2(\text{singlet}), \mathbb{C}l_3^-, H_4SiO_4, OCl_2, Me$ 

. Then calculate the number of species which have  $(p_\pi - p_x)$  back bond

or  $(p_\pi - d_\pi$  back bond.

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**15.** Total number of molecules which can form H-bond among themselves.

 $SiH_3OH, HCN, B(OMe)_3, NHMe_2, CH_2CONH_2, HCHO, HCOOH, N$ 



16. Consider two covalent compounds  $AL_{n_1}$  and  $BL_{n_2}$ , if central atom (A) of first compounds has total six electron pairs and central atom (B) of second compound contains total five electron pairs in its valence shell and both compounds are planar and non-polar then calculate value of experssion  $(n_1 - n_2)^2$ .

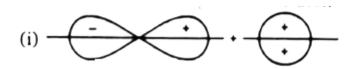
[Where  $n_1 \; {
m and} \; n_2$  are number of monovalent surrounding atom (L)]



**17.** Calculate the I-I distance in (Å) for given compound  $H_2C_2I_2$  if C-I bond length is 2.35Å.  $(\sin 60^\circ = 0.866)$ 



**18.** There are some arrangements of atomic orbitals which are given below:



Then calculate the value of "QtimesR-P" where P, Q and R are no. of arrangements which give bonding molecular orbitals (positive overlap), antibonding molecular orbitals (negative overlap) and non-bonding molecular orbitals (zero overlap) respectively.



**19.** Number of hybrid orbital C atoms which have 33% p-character in  $C(CN)_{{\scriptscriptstyle A}}.$ 



**20.** Max. no. of equal P-O bonds in  $P_2O_7^{4-}$  ion is :

## 21. Consider the following species:

- (i)  $CH_3^+$
- (ii)  $(C_3H_5)_3Al$
- (iii) HCHO
- (iv)  $CH_4$
- (v)  $\left(C_2H_5
  ight)_3N$
- (vI)  $TiCl_4$
- (vii)  $CO_2$
- (viii)  $SiCl_4$
- (ix)  $BF_3$

the find out total number of species which can act as Lewis acid.



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**22.** Calculate value of |x-y|, here x and y are the total number of bonds in benzene and benzyne respectively which are formed by overlapping of

hybridized orbitals.



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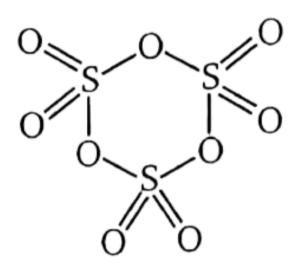
23. Consider the following compounds:

(i) 
$$IF_5$$
 (ii)  $ClI_4^-$  (iii)  $XeO_2F_2$  (iv)  $NH_2^-$  (v)  $BCl_3$  (vi)  $BeCl_2$  (vii)  $AsCl_4^+$  (viii)

$$B(OH)_3$$
 (ix) $NO_2^-$  (x) $ClO_2^+$ 

Then, calculate value of "x+y-z", here x, y and z are total number of compounds in given compounds in which central atom used their all three p-orbitals, only two p-orbitals and only one p-orbital in hybridisation respectively:





Calculate value of p+q, here p and q are total number of  $d\pi-p\pi$  bonds and total number of  $sp^3$  hybridised atoms respectively in given molecule.



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**25.** Consider the following orbitals (i) $3p_x$  (ii) $4d_{z^2}$  (iii) $3d_{x^2-y^2}$  (iv) $3d_{yz}$ 

Then, calculate value of "x+y-z" here x is total number of gerade orbital and y is total number of ungerade orbitals and z is total number of axial orbitals in given above orbitals.

## 26. Consider the following oxyanions:

$$PO_4^{3-}, P_2O_6^{4-}, SO_4^{2-}, MnO_4^-, CrO_4^{2-}, S_2O_5^{2-}, S_2O_7^{2-}$$

and find the value of R+O-P

where P-number of oxy anions having three equivalent X-O bonds per

Q=number of oxy anions having two equivalent X-O bonds per central atom.

R=Number of oxy anions having four equivalent X-O bonds per central atom.



central atom

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**27.** Consider the following three compounds  $(i)AX_{2n}^{n-}$ , (ii) $AX_{3n}$  and (ii)  $AX_{4n}^{n+}$ , where central atom A is 15th group element and their maximum covalency is 3n. If total number of proton in surrounding atom X is n and value of n is one, then calculate value of  $x^3+y^2+z^2$  . (where x, y and z are total number of lone pair at central atom in compounds (i), (ii) and (iii) respectively.



28. Consider the following compounds

(1) $H_3CF$  (2) $H_2CF_2$  (3) $CH_4$  (4) $H_3CCF_3$  (5) $CH_3CH_2+$  (6) $C_2H_4$  and calculate value of Y+X, (where X is the total number of compounds which have H-C-H bond angles equal to  $109^\circ\,28'$  and Y is the total number of compounds which have H-C-H bond angles greater than  $109^\circ\,28'$  and less than  $120^\circ$ 



**29.** There are some species given below. (i) $O_2^+$  (ii)CO (iii) $B_2$  (iv) $O_2^-$  (v)  $NO^+$  (vi) $He_2^+$  (vii) $C_2^{2+}$  (vii) $CN^-$  (ix) $N_2^-$ 

total number of species which have fractional bond order.



**30.** Following compounds A and B have similar structure with delocalization of  $\pi$ -electrons system.  $(A)(PNCl_2)_{\mathbf{x}/2}$  (B) $(CH)_x$  If value of x is 6, then calculate value of "P+Q", where 'P' is total no. of  $\sigma$ -bonds in compounds A and B and 'Q' is total no. of  $\pi$  bonds in compounds A and B.



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**31.** The hybridization of central atoms of compounds A, B, C and D are  $sp^3d$ ,  $sp^3$ ,  $sp^2$  and sp respectively. If compounds A and D have same shape like  $I_3^-$  and compounds B and C have same shape like water structure. Then calculate value of "P+Q+R+S", where P, Q, R, and S are number of lone pairs on central atoms of compounds A, B, C and D respectively.



**32.** In compound  $PCl_xF_{5-x}$ , possible values of x are 0 to 5, then calculate value of  $x_1+x_2+x_3$  (where  $x_1,x_2$  and  $x_3$ ... are possible values of x, with zero dipole moment for given compound).



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**33.** There are two groups of compounds A and B. Groups A contains three compounds  $Px_4$ ,  $Qy_3$ ,  $Rz_2$ . Groups B also contains three compounds  $Sx_4$ ,  $Ty_3$ ,  $Uz_2$ . Hybridization of each central atom of group A compounds is same as that of iodine in  $IBrCl^-$  while in group B compounds it is same as that of iodine  $IBrCl^+$ . Substituents X, Y and Z exhibit covalency of one in ground state. Then find the value of x/y. Where, x and y are total number of lone pair present at central atoms of



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compounds of group A and B respectively.

**34.** Consider the following three compounds  $(i)AX_{2n}^{n-}$ ,  $(ii)AX_{3n}$  and  $(ii)AX_{4n}^{n+}$ , where central atom A is 15th group element and their maximum covalency is 3n. If total number of proton in surrounding atom X is n and value of n is one, then calculate value of  $x^3+y^2+z^2$ . (where x, y and z are total number of lone pair at central atom in compounds (i), (ii) and (iii) respectively.



**35.** When  $B_2H_4$  is allowed to react with following lewis bases, then how many given lewis bases form adduct through symmetrical cleavage of  $B_2H_6$ )?

 $NH_3,\,MeNH_3$ , pyridine, CO, T.H.F.,  $PH_3,\,PF_3,\,Me_3N,\,Me_3NH$ 



**36.** Consider the following elements A, B, C and D and their outer electronic configurtions are  $ns^2np^1, ns^2np^3, ns^2np^4$  and  $ns^2np^5$ 

respectively. Element E also has same outer electronic configuration like D but shows only single oxidation state(-1). If element A, B , C and D belong to same period as that of sodium. Consider the following compounds.  $(i)CE_4 \ (ii)BD_2E_3 \ (iii)DE_3 \ (iv)DE_2 \ (v)BD_3E_2 \ (vi)C_2E_2 \ (vii)DE \ (viii)A_2D_6.$  Then calculate the value of x+y, (where x and y are total number of polar and non-polar compounds).



**View Text Solution** 

**37.** Consider following four compounds:

- (i)  $C_x O_y$
- (ii)  $C_x O_{y+1}$
- (iii)  $C_{x+2}O_{y+1}$  and (iv)  $C_{x+11}O_{y+8}$ ,

if "x=y=1", then calculate the value of |p-q|, where p and q are total number of  $sp^2$  and hybridized carbon atoms respectively in given four compounds.



**38.** Calculate expression (x+y+z) for diatomic molecules.

where x=Total number of singly occupied molecular orbital (SOMO) in  $O_2$ . y=Total number of singly occupied molecular orbital (SOMO) in  $B_2$ 

z=Total number of singly occupied molecular orbital (SOMO) in NO.



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**39.** If Hund rule violate, then find the total number of species among following which whill be dimagnetic:

$$B_2, O_2, N_2^-, C_2, NO, OF, N_2^{2-}, BN$$



## 40. Consider the following table

Compounds (X are monovalent surrounding atoms)		Central atoms(A to D) belong to group	Characteristics of compounds	Number of lone pair(s) at central atom
(i)	AX 3	16	Planar and polar	m ,
(ii)	BX n2	15	Trigonal pyramidal	m 2
(iii)	ex.	14	Zero dipole moment	m 3
(iv)	DX n4	13	All X — D — X bond angle are 120°	m <sub>4</sub>

Than calculate value of experssion  $\left| \frac{n_1 + n_2 + n_3 + n_4}{m_1 + m_2 + m_3 + m_4} \right|^2$ .

**41.** Total number of species among following, in which bond angle is equal to or less than  $109^{\circ}\,28$  and also they act as lewis base :

 $NH_3, NMe_3, O(SiH_3)_2, ICl_4^-, XeO_3, BF_2Cl, SiF_4, AsH_3, SO_2F_2$ 



**42.** Total number of unpaired electrons(s) present in both cationic and anionic part of compound  $O_2[PtF_6]$ .



**43.** Total number of species which has/ have symmetrical electronic distribution in their HOMO and also paramagnetic.

$$N_2^+, O_2^{2-}, C_2, O_2, B_2, C_2^{2-}, N_2^{2-}.$$



44. Total number of molecules, in which each covalent bond is comprisedof effective back bond.

 $BF_3, N(SiH_3)_3, PF_3, , POF_3, B(OH)_3, O(SiH_3)_2, BFH_2, BF_2(NH_2), N_3, D(SiH_3)_3, D(SiH_3)_4, D(SiH_3)_5, D(SiH_3)_6, D(SiH_3)_6$ 





**46.** Consider the following species  $O(Me)_2, N(SiH_3)_3, CO, O(SiH_3)_2, \mathbb{C}l_2(\mathrm{singlet}), \mathbb{C}l_3^-, H_4SiO_4, OCl_2, Me$ 

. Then calculate the number of species which have  $(p_\pi-p_x)$  back bond or  $(p_\pi-d_\pi$  back bond.

**45.** Total number of angle in  $SeCl_4$  which are less than  $90^{\circ}$ .



**47.** Total number of molecules which can form H-bond among themselves.



**48.** Consider two covalent compounds  $AL_{n_1}$  and  $BL_{n_2}$ , if central atom (A) of first compounds has total six electron pairs and central atom (B) of

second compound contains total five electron pairs in its valence shell and both compounds are planar and non-polar then calculate value of experssion  $(n_1-n_2)^2$ .

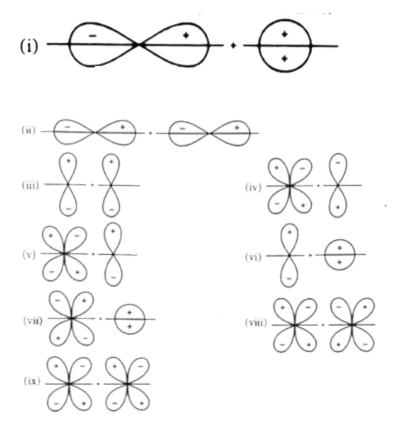
[Where  $n_1 \; {
m and} \; n_2$  are number of monovalent surrounding atom (L)]



**49.** Calculate the I-I distance in (Å) for given compound  $H_2C_2I_2$  if C-I bond length is 2.35Å.  $(\sin 60^\circ = 0.866)$ 



**50.** There are some arrangements of atomic orbitals which are given below:



Then

calculate the value of "`QtimesR-P'" where P, Q and R are no. of arrangements which give bonding molecular orbitals (positive overlap), antibonding molecular orbitals (negative overlap) and non-bonding molecular orbitals (zero overlap) respectively.



**51.** Number of hybrid orbital C atoms which have 33% p-character in  $C(CN)_4.$ 



**52.** Max. no. of equal P-O bonds in  $P_2O_7^{4-}$  ion is :



**53.** Consider the following species:

(i)  $CH_3^+$ 

- (ii)  $(C_3H_5)_3Al$
- (iii) HCHO

(iv)  $CH_4$ 

(v)  $(C_2H_5)_3N$ 

(vI)  $TiCl_4$ 

(vii)  $CO_2$ 

(viii)  $SiCl_4$ 

(ix)  $BF_3$ 

the find out total number of species which can act as Lewis acid.



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**54.** Calculate value of |x-y|, here x and y are the total number of bonds in benzene and benzyne respectively which are formed by overlapping of hybridized orbitals.



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**55.** Consider the following compounds:

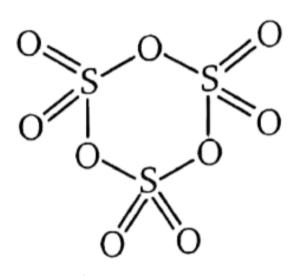
$$(i)IF_5 \hspace{0.5cm} (ii)ClI_4^- \hspace{0.5cm} (iii)XeO_2F_2 \hspace{0.5cm} (iv)NH_2^-$$

$$(v)BCl_3 \hspace{0.5cm} (vi)BeCl_2 \hspace{0.5cm} (vii)AsCl_4^+ \hspace{0.5cm} (viii)B(OH)_3$$

(ix) 
$$NO_2^ (x)ClO_2^+$$

Then calculate value of "x+y-z", here, x,y and z are total number of compounds in given compounds in which central atom used their all three p-orbitals, only two p-orbitals and only one p-orbital in hybridisation respectively.

**56.** Consider the following



Calculate value of p+q, here p and q are total number of  $d\pi-p\pi$  bonds and total number of  $sp^3$  hybridised atoms respectively in given molecule.



**57.** Consider the following orbitals (i) $3p_x$  (ii) $4d_{z^2}$  (iii) $3d_{x^2-y^2}$  (iv) $3d_{yz}$ 

Then, calculate value of "x+y-z" here  ${\sf x}$  is total number of gerade

orbital and y is total number of ungerade orbitals and z is total number of axial orbitals in given above orbitals.



and find the value of R+Q-P

**58.** Consider the following oxyanions:

$$PO_4^{3-}, P_2O_6^{4-}, SO_4^{2-}, MnO_4^-, CrO_4^{2-}, S_2O_5^{2-}, S_2O_7^{2-}$$

where P-number of oxy anions having three equivalent X-O bonds per central atom

Q=number of oxy anions having two equivalent X-O bonds per central atom.

R=Number of oxy anions having four equivalent X-O bonds per central atom.



**59.** Consider the following three compounds  $(i)AX_{2n}^{n-}$ , (ii) $AX_{3n}$  and (ii)  $AX_{4n}^{n+}$ , where central atom A is 15th group element and their maximum

covalency is 3n. If total number of proton in surrounding atom X is n and value of n is one, then calculate value of  $x^3+y^2+z^2$ . (where x, y and z are total number of lone pair at central atom in compounds (i), (ii) and



**60.** Consider the following compounds

(iii) respectively.

(1) $H_3CF$  (2) $H_2CF_2$  (3) $CH_4$  (4) $H_3CCF_3$  (5) $CH_3CH_3$  (6) $C_2H_4$  and calculate value of Y+X, (where X is the total number of compounds which have H-C-H bond angles equal to  $109^\circ 28'$  and Y is the total number of compounds which have H-C-H bond angles greater than  $109^\circ 28'$  and less than  $120^\circ$ 



**61.** There are some species given below. (i) $O_2^+$  (ii)CO (iii) $B_2$  (iv) $O_2^-$  (v)  $NO^+$  (vi) $He_2^+$  (vii) $C_2^{2+}$  (vii) $CN^-$  (ix) $N_2^-$ 

total number of species which have fractional bond order.

**62.** Following compounds A and B have similar structure with delocalization of  $\pi$ -electrons system.  $(A)(PNCl_2)_{\mathbf{x}/2}$  (B) $(CH)_x$  If value of x is 6, then calculate value of "P+Q", where 'P' is total no. of  $\sigma$ -bonds in compounds A and B and 'Q' is total no. of  $\pi$  bonds in compounds A and B.



**63.** The hybridization of central atoms of compounds A, B, C and D are  $sp^3d$ ,  $sp^3$ ,  $sp^2$  and sp respectively. If compounds A and D have same shape like  $I_3^-$  and compounds B and C have same shape like water structure. Then calculate value of "P+Q+R+S", where P, Q, R, and S are number of lone pairs on central atoms of compounds A, B, C and D respectively.



**64.** In compound  $PCl_xF_{5-x}$ , possible values of x are 0 to 5, then calculate value of  $x_1+x_2+x_3$  (where  $x_1,x_2$  and  $x_3$ ... are possible values of x, with zero dipole moment for given compound).

