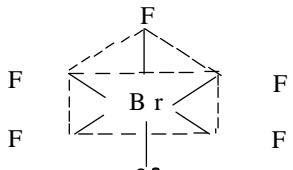


GET EQUIPPED FOR IIT – JEE Advanced

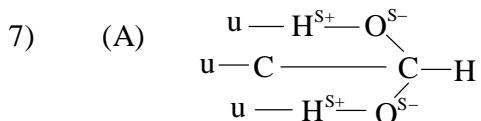
Only one option correct

- 1) (B) 4 B.P and Zero L.P is SiF_4 .
- 2) (D) ICl_2^+ has 2 B.P and 2 L.P.
 Sp^3 hybridization and bent shape.
- 3) (C) Sp^2 hybridization and trigonal planar geometry.
- 4) (D) BrF_5 : 5 B.P + 1 L.P
 Sp^3d^2 hybridization and square pyramidal geometry.
 8 F – Por – F angles at 90° .



- 5) (C) ClO_4^- : 4 B.P + Zero L.P Sp^3 hybridization.

- 6) (D)



- 8) Bond order of $\text{O}_2^{2-} = 1$; $\text{O}_2 = 2$
 $\text{O}_2^- = 1.5$; $\text{O}_2^+ = 2.5$

Hence, Stability $\text{O}_2^+ > \text{O}_2 > \text{O}_2^- > \text{O}_2^{2-}$; Ans : (B)

- 17) $\text{NO}_2^+ \Rightarrow \text{Sp}$; $\text{NO}_3^- \Rightarrow \text{Sp}^2$; $\text{NH}_4^+ \Rightarrow \text{Sp}^3$ (3)

- 18) (B) $\text{NH}_3 \Rightarrow \text{Sp}^3$; $[\text{PHl}_4]^{2-} \Rightarrow \text{dSp}^2$ (Square planar)
 $\text{PCl}_5 \Rightarrow \text{Sp}^3\text{d}$; $\text{BCl}_3 \Rightarrow \text{sp}^2$.

- 19) (A) All are 14 e^- squares with B.O = 3.

- 20) (A) The e^- density lies above and below the inter-nuclear / molecular axis.

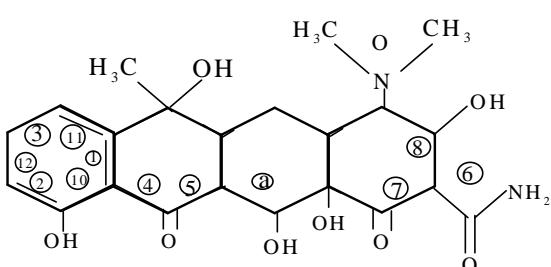
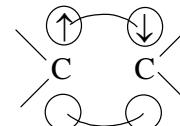
- 21) (B)

- 22) (C) $\text{O}_2^- : \sigma / s^2 \sigma^* / s^2 \sigma 2s^2 \sigma 2p_z^2 z 2p_n^2 z^y 2p_n^2 \sigma^* 2p_z z 2py^2 z^y 2py^1$

- 23) (B) More than number of H – atoms, more is the H – Bonding hence, Due to higher intermolecular forces of attraction, higher is the M. pt.

- 24) (A) Lead Oxide PbO_2 .

- 25) (D) 12



- 26) (C)

- 27) (C) Dipole Moment of 

$$= \sqrt{3} \times 1.5 \text{ D} \quad \left(\sqrt{p_1^2 + p_2^2 + 2p_1 p_2 \cos 60^\circ} \right)$$

$$= 2.6 \text{ D.} \quad P_1 = P_2.$$

- 28) (D) O – H bond is more polar than N – H bond. & SO + on H is more in O – H bond. Also N is better donor than O

- 29) (C) 

Comprehension Type

Passage 1:

- 1) (D) Both F – atoms are present in axial positions.

2) (A) uF_3 has Sp^3d hybridization ; 3 B. P + 2 L.P

3) (D) PI_5 and $[\text{PBr}_6]^e$ do not exist due to steric factors. In PH_5 , Because of its low electronegative H – atom cannot effect the contraction of dz^2 orbital to form Sp^3 hybridized P = atom.

4) (B) $[\text{PBr}_6]^e$ does not exist since P cannot accommodate 6 Br – atoms around it due to steric hindrance.

5) (D) ${}^{30}\text{P}({}^{35}\text{Cl})_5 + {}^{30}\text{P}({}^{37}\text{Cl})_5 + {}^{31}\text{P}({}^{35}\text{Cl})_5 + {}^{31}\text{P}({}^{37}\text{Cl})_5 = 4$

i) ${}^{31}\text{P}({}^{35}\text{Cl})_4({}^{37}\text{Cl})_4 \rightarrow {}^{31}\text{P}({}^{35}\text{Cl})_{ax}({}^{37}\text{Cl})_{eq}$
 $= 2$ forms
 $\rightarrow {}^{31}\text{P}({}^{35}\text{Cl} {}^{37}\text{Cl}_3)_{eq}({}^{37}\text{Cl})_{ax}$

ii) ${}^{31}\text{P}({}^{35}\text{Cl})_2({}^{37}\text{Cl})_3 \rightarrow {}^{31}\text{P}_{ax}({}^{35}\text{Cl})_2({}^{37}\text{Cl})_3$
 $3 = \rightarrow {}^{31}\text{P}({}^{35}\text{Cl} {}^{37}\text{Cl})_{ax}({}^{35}\text{Cl} {}^{37}\text{Cl}_2)_{eq}$
 $\rightarrow {}^{31}\text{P}({}^{37}\text{Cl})_{ax}({}^{35}\text{Cl}_2 {}^{37}\text{Cl})_{eq}$

iii) ${}^{31}\text{P}({}^{35}\text{Cl})_3({}^{37}\text{Cl})_2 = 3$ forms ; same as (ii)

iv) ${}^{31}\text{P}({}^{35}\text{Cl})_4({}^{37}\text{Cl})_1 = 2$ forms ; same as (ii)

Similarly with ${}^{30}\text{P}$ – atom as central Atom, 10 forms will be there. Total = 24.

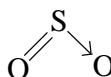
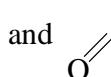
6) SF₄ has 4 G.P + 1 L.P ; Hence (C).

7) (C) IF₂⁻ (Sp³d) has 2B.P + 3 L.P

8) (C) In Sp³d hybridized, dz² orbital is utilized.

Passage 2:

1) A) i) I₃⁺ : 2B.P + 2L.P; Sp³ hybridization

ii) SO₂ can be represented as  and  . It is considered that S being in 3rd period can utilize its d – orbitals to form p_z – dx bonds with O atoms.

iii) XeF₂ : 3 L.P and 2 B.P \Rightarrow Sp³d hybridization.

CO₂ : Sp hybridization \Rightarrow 2 B.P

iv) SF₄ : Sp³d hybridization \Rightarrow 4 B.P + 1 L.P

ICl₃ : Sp³d hybridization \Rightarrow 3 B.P + 2 L.P

2) I₃⁻ : L.P = 3; B.P = 2 Ratio \Rightarrow 1.5

XeF₄ : L.P = 2 ; B.P = 4 Ratio \Rightarrow 0.5

3) (A) NO₂⁺ \Rightarrow linear \Rightarrow 180°

NO₂ (odd e⁻ system) \Rightarrow bent \Rightarrow 134°

NO₂ (1 L.P) \Rightarrow bent \Rightarrow 115°

4) (B) Bond angle PF₃ > PCl₃ < PBr₃ < PI₃.

Due to dx – px bonds in PF₃, double bond character develops increase in B.P – B.P repulsions lead to increase in Bond Angle.

5) (D) B.O of O₂⁺ = 2.5; O₂⁻ = 1.5; O₂ = 2, O₂⁺² = 3; O₂²⁻ = 1

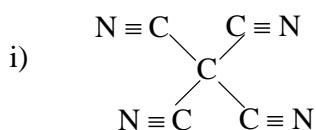
Bond length O₂²⁻ > O₂⁻ > O₂ > O₂⁺² > O₂⁺

6) (D) B.O of N₂⁺ = 2.5

B.O of N₂⁻ = 2.5; e⁻s occupy non – bonding Molecular orbital's.

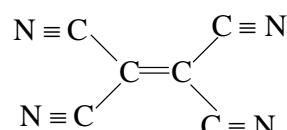
\therefore bond strength of N₂⁻ < Bond strength of N₂⁺

7) (C)



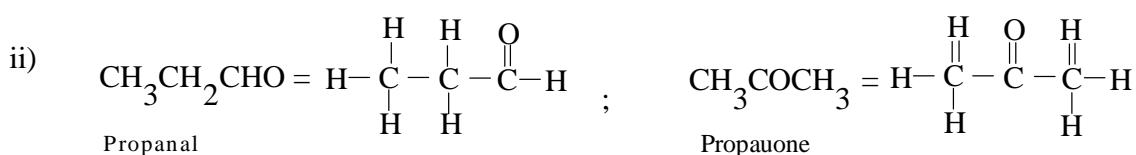
σ bonds = 8

π bonds = 8



σ bonds = 9

π bonds = 9



- ii) XeF_4 is square planar in shape.
- SF_4 has See – saw Shape.
- iv) Due to smaller orbital size, effective overlapping take place in u – u bond.

Passage 3:

- 1) (D)
- 2) (A) In HF_2^- H bonding exists between HF and I^- .
- 3) (B)

Passage 4:

- 1) A) BeCl_2 has maximum covalent character because of smallest size of cation (Be^{+2})
- 2) D) I^- is the largest Anion.
It has maximum polar ability.
- 3) C) Highest positive charge and smallest size of cation (Al^{+3}), hence maximum polarization and covalent character.
- 4) B) Liu has maximum covalent character, hence it will be most soluble in non – polar solvent ether.
- 5) D) CaI_2 has maximum covalent character hence least M.pt...

Passage 5:

- 1) A) and B) These are planar molecules. XeF_4 has square planar shape.
 XeF_4 has trigonal planar shape.

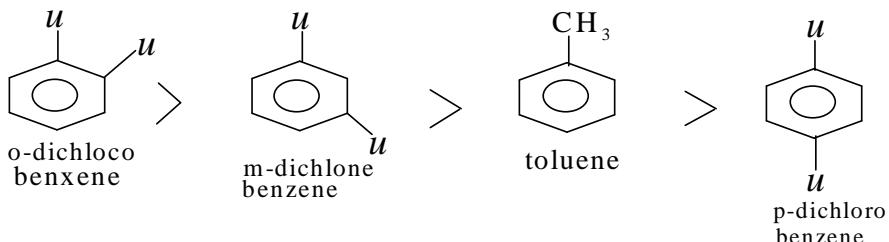
- 2) $p = q \times l$

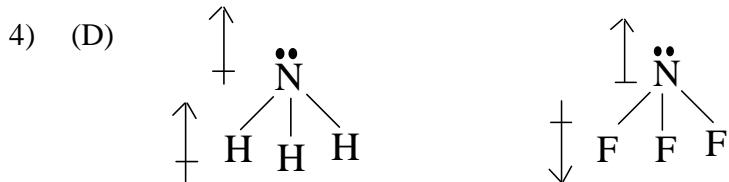
$$q = \frac{p}{l} = \frac{1.2 \times 10^{-18} \text{ esu cm}}{10^{-8} \text{ cm}}$$

$$q = 1.2 \times 10^{-10} \text{ esu.}$$

$$\text{Fraction of change} = \frac{1.2 \times 10^{-10}}{4.8 \times 10^{-10}} = \frac{1}{4} = 0.25.$$

- 3) (C)





5) % Ionic character = $\frac{1.5 \times 10^{-29} \text{ C.m}}{1.6 \times 10^{-19} \text{ C} \times 150 \times 10^{-12} \text{ m}} \times 100 = 62.5\%$

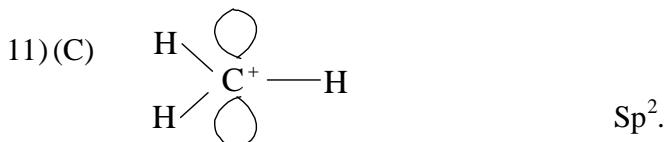
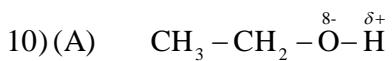
Passage 6:

- 1) B
- 2) A
- 3) A.

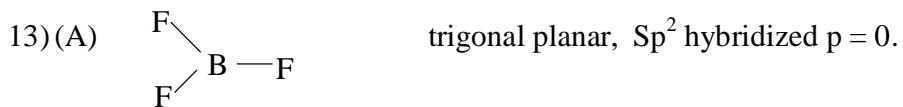
Window To IIT – JEE Advance

C. Multiple Choice Questions with ONE correct answer :

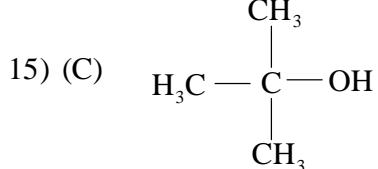
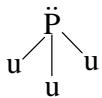
- 1) (C) Sp^2 hybridized, $p = 0$.
- 2) (A) CO and $\text{CN}^- \Rightarrow 14 e^-$ systems.
- 3) (A) CO_2 . Sp hybridized $\text{O}=\text{C}=\text{O}$
- 4) (B) $\text{CCl}_4 \Rightarrow \text{Sp}^3$ hybridized with tetrahedral geometry.
- 5) (B)
- 6) (D)
- 7) (A) $\text{NO} \Rightarrow 15 e^-$ system ; has 1 unpaired e^- .
 $\text{O}_2 \Rightarrow 16 e^-$ system ; has 2 unpaired e^- s.
- 8) (C) More than bond forms stronger H – Bonds.
- 9) (C)



12)(A)

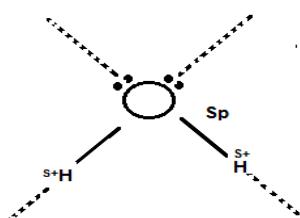


14) (A) PCl_3 has 3 B.P + 1 L.P



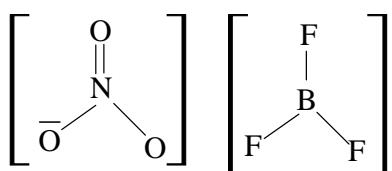
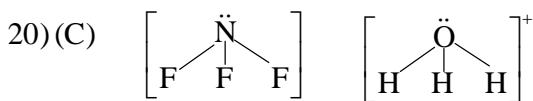
16) (A) 2 B.P + 2 L. P ; Sp^3 hybridized.

17) (B)



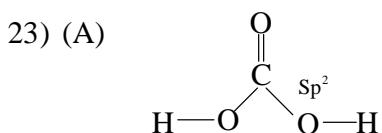
18) (C) Least oxidation state, least polarization, least covalent character.

19) (D) $\text{O}_2 : \sigma / s^2 \quad \sigma^x / s^2 \quad \sigma 2s^2 \quad \sigma 2s^2 \quad \sigma 2\text{pz}^2 \pi 2\text{pn}^2 \quad \pi^x 2\text{pn}^1 \quad \sigma^x 2\text{pz} \quad 2\text{py}^2 \quad \pi^x 2\text{py}^1$



21) (B) $\text{Ca}^{+2} [\text{C} \equiv \text{C}]^{2-} \quad 1\sigma + 2\pi$

22) (C) $\text{O}_2^- \Rightarrow$ superoxide ion.



24) (C)

25)(D)

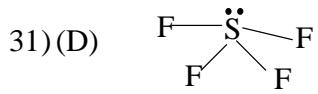
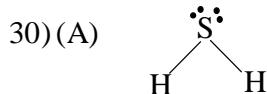
26)(B)

27)(A)

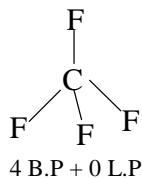
28)(C)

29)(D) B.O of CO = 3 ; CO₂ = 2 ; CO₃²⁻ = 1.5.

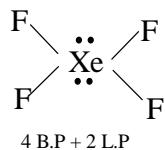
∴ Bond Length : CO₃²⁻ > CO₂ > CO.



32) 4 BP + 1 L.P

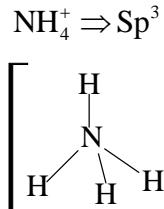
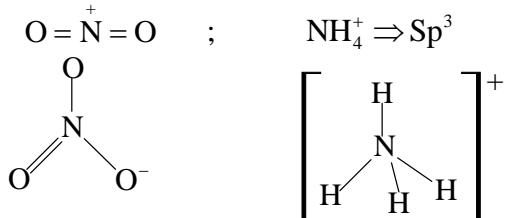


4 B.P + 0 L.P



4 B.P + 2 L.P

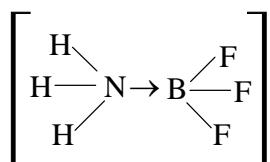
33)(B) $\text{NO}_2^+ \Rightarrow \text{Sp}$
 $\text{NO}_3^- \Rightarrow \text{Sp}^2$



34)(A)

35)(B)

36)(A)



37)(C)

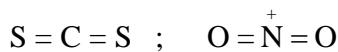
38) O₂⁺ ⇒ B.O is 25 ; (B)

39)(D) I₃⁻ has 3 L.P, ; XeF₄ has 2 L.P
SF₄ has 1 L.P, ; ClO₃⁻ has 1 L.P

D. Multiple Choice Questions with ONE or MORE THAN ONE correct answer :

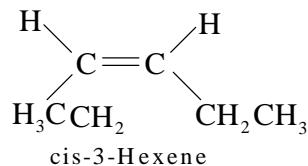
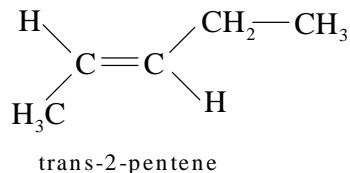
(1) (A) and (C). $\text{Cl}-\text{Hg}-\text{Cl}$; $\text{H}-\text{C} \equiv \text{C}-\text{H}$

(2) (B), (C), (D).



(3) A, (C) CN^- and NO^+ has B.O = 3.

(4) B, C.



(5) (D) H_3O^+ , NH_3 and CH_3^\ominus have Sp^3 hybridization.

(6) (B)

(7) (A)

(8) (B)

(9) (B), (D)

(10) B, C, D

(11) A, B, D

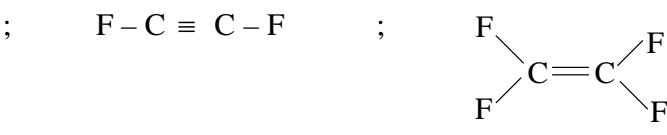
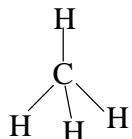
(12) A, B, C. HF and H_2O have abnormally high B.pt due to H – Bonds

(13) A, C, D $\text{Cl}-\text{Hg}-\text{Cl}$; $[\text{u}-\overset{\cdot}{\underset{\cdot}{\text{I}}}-\text{u}]^\ominus$; $\text{S}=\text{C}=\text{S}$

14) B, D

15) A, D

16) A, B, C

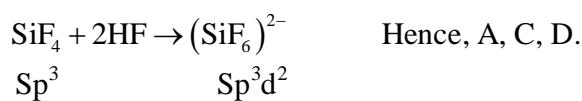
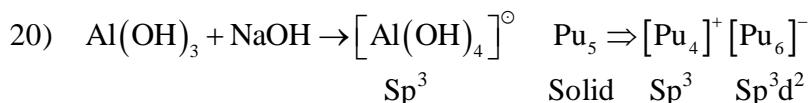


17) A, B, C, D.

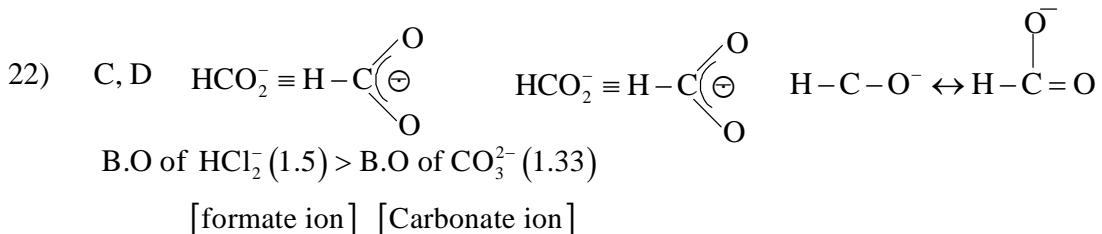
18) B, C $\text{SF}_4 \Rightarrow 4 \text{ B.P} + 1 \text{ L.P}$

$\text{PF}_5 \Rightarrow 5 \text{ B.P} + \text{Zero L.P.}$

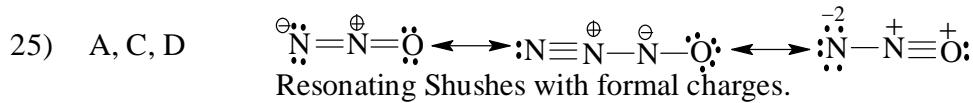
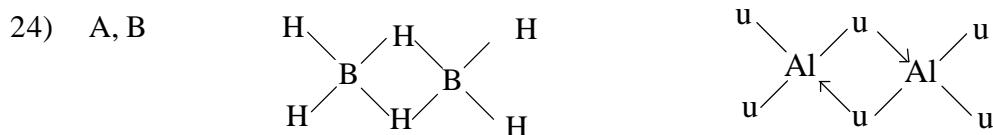
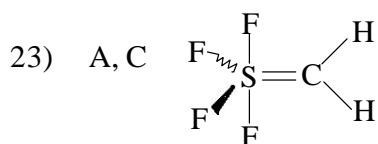
19) A, C



21) A, C ; Only O₂ and B₂ are paramagnetic.



∴ Bond length of HCO₂⁻ < Bond length of CO₃²⁻



26) A, B, C, D.

27) A, B, C, D

More electronegative element in each case, i.e. O and F occupy axial position in I & II respectively.