Chemical Bonding And Molecular Structure

Question1

Identify the correct answer.

[NEET 2024]

Options:

A.

Three resonance structures can be drawn for ozone

В.

 ${\rm BF}_3$ has non-zero dipole moment

C.

Dipole moment of NF_3 is greater than that of NH_3

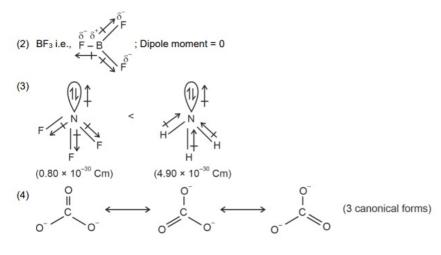
D.

Three canonical forms can be drawn for CO $_3\ ^{2-}\text{ion}$

Answer: D

Solution:

(1) In ozone; there are two resonating structures.



Question2

Which of the following molecules has "NON ZERO" dipole moment value?

[NEET 2024 Re]

- A.
- CCl_4
- B.
- HI
- C.

 CO_2

D.

 BF_3

Answer: B

Solution:

Dipole moment of a molecule depends both on shape and bond dipole.

Molecule	Shape Cl	μ(Debye)
CCl4		0
HI	H-1	0.38
CO ₂	O = C = O	0
BF3		0

Question3

Identify the incorrect statement about PCl₅.

[NEET 2024 Re]

Options:

A.

 PCl_5 possesses two different Cl - P - Cl bond angles

B.

All five P – Cl bonds are identical in length

C.

PCl 5 exhibits $sp^3 d$ hybridisation

D.

 PCl_5 consists of five P - Cl (sigma) bonds

Answer: B

Solution:



It is sp^3d hybridised with axial to equatorial angle of 90° and equatorial bond angles of 120°. It has five P - Cl sigma bonds. Axial bonds are longer than equatorial bonds.

Question4

Match List-I with List-II:

>>

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List-II

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List-I

	(Molecule)		(Bond enthalpy (kJ mol ⁻¹))
Α.	HCl	I.	435.8
В.	N ₂	II.	498
C.	H ₂	III.	946.0
D.	0 ₂	IV.	431.0

Choose the correct answer from the options given below:

[NEET 2024 Re]

Options:

Α.

A-III, B-IV, C-I, D-II

В.

A-IV, B-I, C-III, D-II

C.

A-IV, B-III, C-II, D-I

D.

A-IV, B-III, C-I, D-II

Answer: D

Solution:

(Molecule)	(Bond enthalpy (kJ mol ⁻¹))
HCl	431.0
N ₂	946.0
H ₂	435.8
O ₂	498

Question5

List I (Molecule)		List II (Number and types of bond/s between two carbon atoms)	
А.	ethane	I.	one $\alpha\text{-bond}$ and two $\pi\text{-bonds}$
В.	ethene	II.	two π-bonds
С.	carbon molecule, C ₂	III.	one σ-bond
D.	ethyne	IV.	one σ -bond and one π -bond

Choose the correct answer from the options given below:

[NEET 2024]

Options:

```
A.
A-I, B-IV, C-II, D-III
B.
A-IV, B-III, C-II, D-I
C.
A-III, B-IV, C-II, D-I
D.
```

A-III, B-IV, C-I, D-II

Answer: C

Solution:

A.		III.	one (C – C) σ bond
В.	Ethene H > c = c < H	IV.	one (C – C) σ and one (C – C) π bond
C.	<i>C</i> ₂	II.	two (C – C) π bonds
D.	Ethyne $H - C = C - H$	I.	two (C – C) π bonds and one (C – C) σ bond

Question6

Given below are two statements:

Statement I: The boiling point of hydrides of Group 16 elements follow the order $H_2O > H_2Te > H_2Se > H_2S$.

Statement II: On the basis of molecular mass, H_2O is expected to have lower boiling point than the other members of the group but due to the presence of extensive H - bonding in H_2O , it has higher boiling point.

In the light of the above statements, choose the correct answer from the options given below:

[NEET 2024]

Options:

A.

Both Statement I and Statement II are true

В.

Both Statement I and Statement II are false

C.

Statement I is true but Statement II is false

D.

Statement I is false but Statement II is true

Answer: A

Solution:

 $\label{eq:statement I} \textbf{Statement I} is correct, because boiling point of hydrides of group 16 follows the order H_2O > H_2Te > H_2Se > H_2S \; .$

Statement II due to intermolecular H -bonding H_2O shows higher boiling point than respective hydrides of group 16.

(Both Statement are true)

Order from H_2Te to H_2S is due to decreasing molar mass.

Question7



Intramolecular hydrogen bonding is present in

[NEET 2024]

Options:

A.

B.

C.

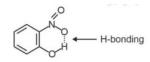
D.

HF

Answer: A

Solution:

In o-nitrophenol intramolecular H-bonding is present.



Question8

Match List I with List II.

	List I (Compound)		List II (Shape/geometry)
А.	NH ₃	I.	Trigonal Pyramidal
В.	BrF ₅	II.	Square Planar
C.	XeF ₄	III.	Octahedral
D.	SF ₆	IV.	Square Pyramidal

Choose the correct answer from the options given below:

[NEET 2024]

```
Options:
A.
A-I, B-IV, C-II, D-III
B.
A-II, B-IV, C-III, D-I
C.
A-III, B-IV, C-I, D-II
D.
```

A-II, B-III, C-IV, D-I

Answer: A

Solution:

NH₃ \Rightarrow sp³ hybridised with 1 lone pair. Structure will be Trigonal Pyramidal. BrF₅ \Rightarrow sp³d² hybridised with 1 lone pair. Structure will be Square Pyramidal. XeF₄ \Rightarrow sp³d² with two lone pairs. Structure will be Square Planar. SF₆ \Rightarrow sp³d² with no lone pair. Structure will be Octahedral. A-I, B-IV, C-II, D-III

Question9

The correct order of energies of molecular orbitals of N_2 molecule, is

[NEET 2023]

Options:

A.

 $\sigma 1s < \sigma * 1s < \sigma 2s < \sigma * 2s < \sigma 2p_z < (\pi 2p_x = \pi 2p_y) < (\pi * 2p_x = \pi * 2p_y) < \sigma * 2p_z < \sigma 2p$

В.

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

C.

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

D.

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

Answer: D

Solution:

For molecules like B_2 , C_2 , N_2 etc. the increasing order of energies of various molecular orbitals is $\sigma 1s < \sigma * 1s < \sigma 2s < \sigma * 2s < (\pi 2p_x = \pi 2p_y) < \sigma 2p_z < (\pi * 2p_x = \pi * 2p_y) < \sigma * 2p_z$

Question10

Amongst the following the total number of species NOT having eight electrons around central atom in its outermost shell, is

NH₃, AlCl₃, BeCl₂, CCl₄, PCl₅ : **[NEET 2023]**

Options:



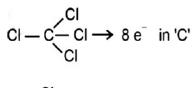
- A.
- 2
- B.
- В
- 4
- ſ
- C.
- 1
- D.
- 3

Answer: D

Solution:

 $\begin{array}{c}
\stackrel{\text{in}}{H} \rightarrow 8 e^{-} \text{ in nitrogen} \\
\stackrel{\text{in}}{H} H \\
\stackrel{\text{in}}{H} H \\
\stackrel{\text{cl}}{H} \rightarrow 6 e^{-} \text{ in Al}
\end{array}$

$$CI - Be - CI \rightarrow 4 e^{-}$$
 in Be



$$\begin{array}{c} CI \\ | \ CI \\ CI - P \\ | \ CI \\ CI \\ CI \end{array} \rightarrow 10 e^{-} \text{ in 'P'}$$

Question11

The correct order of dipole moments for molecules NH_3 , H_2S , CH_4 and HF, is:

[NEET 2023 mpr]

Options:

```
A.

CH_4 > H_2S > NH_3 > HF

B.

H_2S > NH_3 > HF > CH_4

C.

NH_3 > HF > CH_4 > H_2S

D.
```

 $\mathrm{HF} > \mathrm{NH}_3 > \mathrm{H}_2\mathrm{S} > \mathrm{CH}_4$

Answer: D

Solution:

 $\mathrm{HF} > \mathrm{NH}_{3} > \mathrm{H}_{2}\mathrm{S} > \underset{\mathrm{Non-polar}}{\mathrm{CH}_{4}}$

Question12

Which one of the following represents all isoelectronic species?

[NEET 2023 mpr]

Options:

A.

Na⁺ , Cl⁻, O⁻, NO⁺

В.

 $\mathrm{N_2O},\,\mathrm{N_2O_4},\,\mathrm{NO^+}$, NO

C.

Na⁺, Mg²⁺, O⁻, F⁻

D.

 Ca^{2+} , Ar, K^+ , Cl^-

Answer: D

Solution:

Total numbers electrons are same

Ca⁺², Ar, K⁺, Cl⁻ \rightarrow 20 electrons

Question13

Which one of the following statements is incorrect related to Molecular Orbital Theory?

[NEET 2023 mpr]

Options:

A.

The π^* antibonding molecular orbital has a node between the nuclei.

B.

In the formation of bonding molecular orbital, the two electron waves of the bonding atoms reinforce each other.

C.

Molecular orbitals obtained from $2P_{\rm x}$ and $2P_{\rm y}$ orbitals are symmetrical around the bond axis.

D.

A $\pi\mbox{-}bonding$ molecular orbital has larger electron density above and below the internuclear axis.

Answer: C

Solution:





In the formation of BMO, the two electron waves of the bonding atoms reinforce each other due to constructive interference.

Molecular orbitals obtained from $2P_x$ and $2P_y$ orbitals are 'unsymmetrical' around bond axis.

Question14

Given below are two statements:

Statement I : Hydrated chlorides and bromides of Ca,Sr and Ba on heating undergo hydrolysis.

Statement II :

Hydrated chlorides and bromides of Be and Mg on heating undergo dehydration. In the light of the above statements, choose the correct answer from the options given below :

[NEET 2023 mpr]

Options:

A.

Statement-I is correct but Statement-II is false.

В.

Statement-I is incorrect but Statement-II is true.

C.

Both Statement-I and Statement-II are true.

D.

Both Statement-I and Statement-II are false.

Answer: D

Solution:

Hydrated chlorides and Bromides of Ca, Sr and Ba are Ionic so undergo dehydration after heating. Hydrated chlorides and Bromides of Be and Mg are covalent so undergo hydrolysis on Heating.

Question15

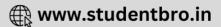
List - I (molecules)	List - II(shape)
(a) NH ₃	(i) square pyramidal
(b) ClF ₃	(ii) trigonal bipyramidal
(c) PCl ₅	(iii) trigonal pyramidal
(d) BrF ₅	(iv) T-shape

Choose the correct answer from the options given below : [NEET Re-2022]

Options:

A. (a) - (iii), (b) - (iv), (c) - (i), (d) - (ii)

B. (a) - (ii), (b) - (iii), (c) - (iv), (d) - (i)



C. (a) - (iii), (b) - (iv), (c) - (ii), (d) - (i)

D. (a) - (iv), (b) - (iii), (c) - (i), (d) - (ii)

Answer: C

Solution:

 $\mathrm{NH}_3 \rightarrow$ Trigonal pyramidal

 $\mathrm{ClF}_3 {\rightarrow} \text{T-shape}$

 $\text{PCl}_5 {\rightarrow} \text{Trigonal bipyramidal}$

 $\mathrm{BrF}_5 {\rightarrow} \text{ Square pyramidal}$

Question16

The correct order of bond angles in the following compounds/ species is : [NEET Re-2022]

Options:

A. $CO_2 < NH_3 < H_2O < NH_4^+$ B. $H_2O < NH_3 < NH_4^+ < CO_2$ C. $H_2O < NH_4^+ < NH_3 < CO_2$ D. $H_2O < NH_4^+ = NH_3 < CO_2$

Answer: B

Solution:

 $\begin{array}{l} \mathrm{CO}_{2} \rightarrow 180^{\circ} \\ \overset{+}{\mathrm{NH}_{4}} \rightarrow 109.5^{\circ} \\ \overset{-}{\mathrm{NHH}_{3}} \rightarrow 107^{\circ} \ (\ \mathrm{N} \ \mathrm{atom \ has \ lone \ pair}) \\ \overset{-}{\mathrm{H}_{2}\mathrm{O}} : \rightarrow 104.5^{\circ} \ (\mathrm{oxygen \ atom \ has \ two \ lone \ pairs)} \end{array}$

Due to lone pair-lone pair repulsions the bond angle in water decreases more.

Question17

Four gas cylinders containing He, N₂, CO₂ and NH₃ gases separately are gradually cooled from a temperature of 500K. Which gas will liquify first ? (Given T_c in K – He : 5.3, N₂ : 126, CO₂ : 304.1 and .NH₃ : 405.5) [NEET Re-2022]

Options:

- A. NH_3
- B. He
- C. N₂
- D. CO_2

Solution:

Ease of Liquefaction < Critical Temperature

So, NH3 will liquify first.

Question18

A vessel contains 3.2g of dioxygen gas at STP (273.15 K and 1 atm pressure). The gas is now transferred to another vessel at constant temperature, where pressure becomes one third of the original pressure. The volume of new vessel in L is: (Given - molar volume at STP is 22.4L) [NEET Re-2022]

Options:

A. 67.2

B. 6.72

C. 2.24

D. 22.4

Answer: B

Solution:

Moles of oxygen $= \frac{3.2}{32} = 10^{-1}$ mole Volume at STP $= 10^{-1} \times 22.4 = 2.24$ L $P_1 = 1$ atm $V_1 = 2.24$ L $P_2 = \frac{1}{3}$ atm $V_2 = ?$ $P_1V_1 = P_2V_2$ $1 \times 2.24L = \frac{1}{3} \times V_2$

 $V_2 = 3 \times 2.24 L = 6.72 L$

Question19

Amongst the following which one will have maximum 'lone pair - lone pair' electron repulsions? [NEET-2022]

Options:

A. Cl F $_3$

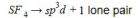
B. I F $_5$

C. SF $_4$

D. X eF $_2$

Answer: D

Solution:



$$F = F$$

$$F =$$

 $ClF_3 \rightarrow sp^3d + 2$ lone pair

$$F_{F_{5}} \rightarrow sp^{3}d^{2} + 1 \text{ lone pair}$$

XeF₂ having maximum lone pairs, so, it has maximum 'lone pair-lone pair' electron repulsions.

Question20

Which amongst the following is incorrect statement? [NEET-2022]

Options:

A. The bond orders of $\mathrm{O_2}^+,\,\mathrm{O_2},\,\mathrm{O_2}^-\text{and}\,\mathrm{O_2}^{2-}$ are 2.5, 2, 1.5 and 1 , respectively

B. C_2 molecule has four electrons in its two degenerate π molecular orbitals

C. H $_2^+$ ion has one electron

D. O_2^+ ion is diamagnetic

Answer: D

Solution:

Solution:

Due to one unpaired electron in $\pi^* 2p$ molecular orbital, O_2^{-1} is a paramagnetic ion.

Question21

BF $_3$ is planar and electron deficient compound. Hybridization and number of electrons around the central atom, respectively are : [NEET 2021]

Options:

A. sp^3 and 4

```
B. sp^3 and 6
```

```
C. sp^2 and 6
```

D. sp^2 and 8

Answer: C

Solution:

- Number of electrons around boron atom is 6.
- Hybridization of B is sp^2 .
- Shape is trigonal planar.

Question22

Match List-I with List-II.

List-I	List-II
(a) <i>PCl</i> 5	(i) Square pyramidaL
(b) <i>SF</i> ₆	(ii) Trigonal planar
(C) BrF 5	(iii) Octahedral
(d) <i>BF</i> 3	(iv) Trigonal bipyramidal

Choose the correct answer from the options given below. [NEET 2021]

Options:

A. (a)-(iv), (b)-(iii), (c)-(i), (d)-(ii)

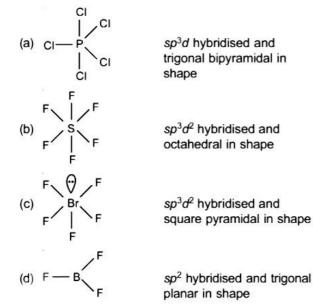
B. (a)-(ii), (b)-(iii), (c)-(iv), (d)-(i)

C. (a)-(iii), (b)-(i), (c)-(iv), (d)-(ii)

D. (a)-(iv), (b)-(iii), (c)-(ii), (d)-(i)

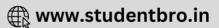
Answer: A

Solution:



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Question23

The correct sequence of bond enthalpy of 'C—X' bond is : [NEET 2021]

Options:

A. $CH_{3}-F < CH_{3}-Cl < CH_{3}-Br < CH_{3}-I$ B. $CH_{3}-F > CH_{3}-Cl > CH_{3}-Br > CH_{3}-I$ C. $CH_{3}-F < CH_{3}-Cl > CH_{3}-Br > CH_{3}-I$ D. $CH_{3}-Cl > CH_{3}-F > CH_{3}-Br > CH_{3}-I$

Answer: B

Solution:

Solution:

The size of halogen atom increases from F to I hence bond length from C-F to C-I increases \therefore Bond enthalpy from CH $_3$ – F to CH $_3$ – I decreases

C – X Bond	Bond dissociationenthalpies/kJ mol^{-1}
CH 3 - F	452
CH ₃ – Cl	351
CH ₃ -Br	293
CH 3 - I	234

Question24

Which of the following molecules is non-polar innature? [NEET 2021]

Options:

A. POCl₃

B. CH ₂O

C. SbCl₅

D. N O_2

Answer: C

Solution:

Solution: Net vector summation of bond moments will be zero so $SbCl_5$ is a non-polar molecule.

Question25

Which of the following set of molecules will have zero dipole moment? [2020]

Options:

A. Boron trifluoride, hydrogen fluoride, carbon dioxide, 1,3-dichlorobenzene

B. Nitrogen trifluoride, beryllium difluoride, water, 1,3-dichlorobenzene

C. Boron trifluoride, beryllium difluoride, carbon dioxide, 1,4-dichlorobenzene

D. Ammonia, beryllium difluoride, water, 1,4-dichlorobenzene

Answer: C

Solution:

Solution: (c) BF $_3$, BeF $_2$, CO $_2$ and 1, 4-dichlorobenzene all are symmetrical molecules.

Question26

Identify a molecule which does not exist. [2020]

Options:

A. Li_2

B. C₂

 $C.O_2$

D. H e_2

Answer: D

Solution:

(d) For He molecule, Electronic configuration is $\sigma 1s^2$, $\sigma^* 1s^2$ Bond order = $\frac{1}{2}(N_b - N_a) = \frac{1}{2}(2 - 2) = 0$ since, bond order of H e_2 is zero, so it does not exist.

Question27

Which of the following diatomic molecular species has only π bonds according to Molecular Orbital Theory? (NEET 2019)

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Options: A. Be₂ B. O₂ C. N₂ D. C₂ **Answer: D Solution:**

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Be₂(8) : K K σ(2s)²σ^{*}(2s)² O₂(16) : K K σ(2s)²σ^{*}(2s)²σ(2p_z)²π(2p_x)²π(2p_y)² π^{*}(2p_x)¹π^{*}(2p_y)¹ N₂(14) : K K σ(2s)²σ^{*}(2s)²π(2p_x)²π(2p_y)²σ(2p_z)² C₂(12) : K K σ(2s)²σ^{*}(2s)²π(2p_x)²π(2p_y)² Therefore, C₂ contains 2π bonds as it has electrons in two pi-molecular orbitals.

Question28

Which of the following is paramagnetic? (Odisha NEET 2019)

Options:

A. N $_2$

B. H₂

C. Li₂

 $D. O_2$

Answer: D

Solution:

$$\begin{split} N_{2}(14) &: K K \sigma 2 s^{2} \sigma^{*} 2 s^{2} \pi 2 p_{x}^{2} = \pi 2 p_{y}^{2} \sigma 2 p_{z}^{2} \\ H_{2}(2) &: \sigma 1 s^{2} \\ Li_{2}(6) &: \sigma 1 s^{2} \sigma^{*} 1 s^{2} \sigma 2 s^{2} \\ O_{2}(16) &: \sigma 1 s^{2} \sigma^{*} 1 s^{2} \sigma 2 s^{2} \sigma^{*} 2 s^{2} \sigma 2 p_{z}^{2} \pi 2 p_{x}^{2} = \pi 2 p_{y}^{2} \\ \pi^{*} 2 p_{x}^{-1} &= \pi^{*} 2 p_{y}^{-1} \end{split}$$

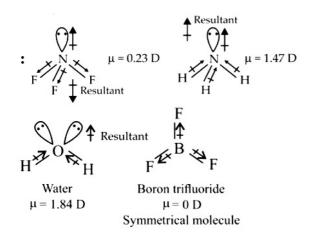
Question29

Which of the following is the correct order of dipole moment? (Odisha NEET 2019)

Options:

A. $NH_{3} < BF_{3} < NF_{3} < H_{2}O$ B. $BF_{3} < NF_{3} < NH_{3} < H_{2}O$ C. $BF_{3} < NH_{3} < NF_{3} < H_{2}O$ D. $H_{2}O < NF_{3} < NH_{3} < BF_{3}$ Answer: B

Solution:



Question30

Consider the following species : CN^+ , CN^- NO and CN. Which one of these will have the highest bond order? (NEET 2018)

Options:

A. NO

B. CN⁻

C. CN $^+$

D. CN

Answer: B

Solution:

Solution: NO: $(\sigma 1s)^2$, $(\sigma^* 1s)^2$, $(\sigma 2s)^2$, $(\sigma^* 2s)^2$, $(\sigma 2p_z)^2$, $(\pi 2p_x)^2 = (\pi 2p_y)^2$, $(\pi^* 2p_x)^1 = (\pi^* 2p_y)^0$ B. O. $= \frac{10-5}{2} = 2.5$ CN⁻: $(\sigma 1s)^2$, $(\sigma^* 1s)^2$, $(\sigma 2s)^2$, $(\sigma^* 2s)^2$, $(\pi 2p_x)^2 = (\pi 2p_y)^2$, $(\sigma 2p_z)^2$ B.O. $= \frac{10-4}{2} = 3$ CN: $(\sigma 1s)^2$, $(\sigma^* 1s)^2$, $(\sigma 2s)^2$, $(\sigma^* 2s)^2$, $(\pi 2p_x)^2 = (\pi 2p_y)^2$, $(\sigma 2p_z)^1$ B.O. $= \frac{9-4}{2} = 2.5$ CN⁺: $(\sigma 1s)^2$, $(\sigma^* 1s)^2$, $(\sigma 2s)^2$, $(\sigma^* 2s)^2$, $(\pi 2p_x)^2 = (\pi 2p_y)^2$ B. O. $= \frac{8-4}{2} = 2$ Hence, CN⁻ has highest bond order.

Question31

In the structure of Cl F $_{\rm 3},$ the number of lone pairs of electrons on central atom 'Cl is (NEET 2018)

Options:

- A. one
- B. two
- C. four
- D. three.

Answer: B

Solution:

Hence, Cl has 2 lone pairs of electrons.

Question32

Which of the following pairs of compounds is isoelectronic and isostructural? (NEET 2017)

Options:

A. T el $_{\rm 2}$, X eF $_{\rm 2}$

B. I $\mathrm{Br_2}^-$, X eF $_2$

C. IF $_3$,X eF $_2$

D. BeCl 2, X eF 2

E. None of the above

Answer: E

Solution:

Species	No.of electrons	Structure
Tel ₂	158	Bent
XeF ₂	72	Linear
IBr ₂	124	Linear
XeF ₂	72	Linear
IF ₃	80	T-Shaped
XeF ₂	72	Linear
BeCl ₂	38	Linear
XeF ₂	72	Linear

Note : In this question, in place of isoelectronic there should be same number of valence electrons.

Question33

The species, having bond angles of 120° is (NEET 2017)

Options:

A. Cl F $_3$

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B. N Cl₃

C. BCl₃

D. PH ₃

Answer: C

Solution:

Solution:

 BCl_3 -Trigonal planar, sp^2 -hybridised, 120° angle.

Question34

Which one of the following pairs of species have the same bond order? (NEET 2017)

Options:

A. O_2 , N O^+

B. CN ⁻, CO

C. N₂, O₂⁻

D. CO, NO

Answer: B

Solution:

Molecular orbital electronic configurations and bond order values are : $O_2 : \sigma 1s^2$, $\sigma^* 1s^2$, $\sigma 2s^2$, $\sigma^* 2s^2$, $\sigma 2p_z^2$, $\pi 2p_x^2 = \pi 2p_y^2 \pi^* 2p_x^{-1} = \pi^* 2p_y^{-1}$ B.O. $= \frac{1}{2}(N_b - N_a) = \frac{1}{2}(10 - 6) = 2$ NO⁺ : $\sigma 1s^2$, $\sigma^* 1s^2$, $\sigma 2s^2$, $\sigma^* 2s^2$, $\sigma 2p_z^2$, $\pi 2p_x^2 = \pi 2p_y^2$ B.O. $= \frac{1}{2}(10 - 4) = 3$ CN⁻ : $\sigma 1s^2$, $\sigma^* 1s^2$, $\sigma 2s^2$, $\sigma^* 2s^2$, $\pi 2p_x^2 = \pi 2p_y^2$, $\sigma 2p_z^2$ B.O. $= \frac{1}{2}(10 - 4) = 3$ CO : $\sigma 1s^2$, $\sigma^* 1s^2$, $\sigma 2s^2$, $\sigma^* 2s^2$, $\pi 2p_x^2 = \pi 2p_y^2$, $\sigma 2p_z^2$ B.O. $= \frac{1}{2}(10 - 4) = 3$ N₂ : $\sigma 1s^2$, $\sigma^* 1s^2$, $\sigma 2s^2$, $\sigma^* 2s^2$, $\pi 2p_x^2 = \pi 2p_y^2$, $\sigma 2p_z^2$ B.O. $= \frac{1}{2}(10 - 4) = 3$ O₂⁻ = $\sigma 1s^2$, $\sigma^* 1s^2$, $\sigma 2s^2$, $\sigma^* 2s^2$, $\sigma 2p_z^2$, $\pi 2p_x^2 = \pi 2p_y^2\pi^* 2p_x^2 = \pi^* 2p_y^1$ B.O. $= \frac{1}{2}(10 - 4) = 3$ O₂⁻ = $\sigma 1s^2$, $\sigma^* 1s^2$, $\sigma 2s^2$, $\sigma^* 2s^2$, $\sigma 2p_z^2$, $\pi 2p_x^2 = \pi 2p_y^2\pi^* 2p_x^2 = \pi^* 2p_y^1$ B.O. $= \frac{1}{2}(10 - 7) = 1.5$ NO : $\sigma 1s^2$, $\sigma^* 1s^2$, $\sigma 2s^2$, $\sigma^* 2s^2$, $\sigma 2p_z^2$, $\pi 2p_x^2 = \pi 2p_y^2\pi^* 2p_x^{-1}$ B.O. $= \frac{1}{2}(10 - 5) = 2.5$

Question35

Which one of the following compounds shows the presence of intramolecular hydrogen bond? (NEET-II 2016)

Options:

- A. H $_2O_2$
- B. HCN
- C. Cellulose
- D. Concentrated acetic acid

Answer: C

Solution:

H 202, H CN and conc. CH 3COOH form intermolecular hydrogen bonding while cellulose has intramolecular hydrogen bonding.

Question36

The hybridizations of atomic orbitals of nitrogen in $\rm N\,O_2^{-+},\,N\,O_3^{--}$ and $\rm N\,H_4^{++}$ respectively are (NEET-II 2016)

Options:

A. sp, sp^3 and sp^2

B. sp^2 , sp^3 and sp

C. sp, sp^2 and sp^3

D. $\mbox{sp}^2\mbox{, sp}$ and \mbox{sp}^3

Answer: C

Solution:

Solution: $X = \frac{1}{2}(VE + MA - c + a)$ For NO₂⁺, $X = \frac{1}{2}(5 + 0 - 1) = 2$ i.e., sp hybridisation For NO₃⁻, $X = \frac{1}{2}(5 + 0 + 1) = 3$ i.e., sp² hybridisation For NH₄⁺, $X = \frac{1}{2}(5 + 4 - 1) = 4$ i.e., sp³ hybridisation

Question37

Which of the following pairs of ions is isoelectronic and isostructural? (NEET-II 2016)

Options:

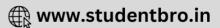
A. CO_3^{2-} , N O_3^{-} B. $Cl O_3^{-}$, CO_3^{2-} C. SO_3^{2-} , N O_3^{-} D. $Cl O_3^{-}$, SO_3^{2-} Answer: D

Solution:

(a) CO_3^{2-} : 6 + 24 + 2 = 32; sp²; trigonal planar

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 $NO_3^-: 7 + 24 + 1 = 32$; sp²; trigonal planar Hence, these are isoelectronic as well as isostructural. (b) $CIO_3^-: 17 + 24 + 1 = 42$; sp³, trigonal pyramidal $CO_3^{2^-}: 6 + 24 + 2 = 32$; sp², trigonal planar Hence, these are neither isoelectronic nor isostructural. (c) $SO_3^{2^-}: 16 + 24 + 2 = 42$; sp³, trigonal pyramidal $NO_3^-: 7 + 24 + 1 = 32$; sp², trigonal planar Theseare neither isoelectronic nor isostructural (d) $CIO_3^-: 17 + 24 + 1 = 42$; sp³, trigonal pyramidal $SO_3^{2^-}: 16 + 24 + 2 = 42$; sp³, trigonal pyramidal Hence, these are isoelectronic as well as isostructural.

Question38

The correct geometry and hybridization for X eF $_{\rm 4}$ are (NEET - II 2016)

Options:

A. octahedral, sp^3d^2

B. trigonal bipyramidal, sp^3d

C. planar triangle, sp^3d^3

D. square planar, sp 3 d 2

Answer: A

Solution:

sp³d² hybridisation
(octahedral geometry, square
 planar shape)

Question39

Among the following, which one is a wrong statement? (NEET-II 2016)

Options:

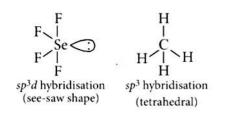
A. PH 5 and BiCl 5 do not exist.

- B. $p\pi d\pi$ bonds are present in SO₂.
- C. SeF $_{\rm 4}$ and CH $_{\rm 4}$ have same shape.

D. I_3^+ has bent geometry.

Answer: C

Solution:



Question40

Consider the molecules CH $_4$, N H $_3$ and H $_2O$ Which of the given statements is false? (NEET- I 2016)

Options:

A. The H - O - H bond angle in H ₂O is smaller than the H - N - H bond angle in N H ₃.

B. The H - C - H bond angle in CH₄, is larger than the H - N - H bond angle in N H₃

C. The H - C - H bond angle in CH $_4$, the H - N - H bond angle in N H $_3$, and the H - O - H bond angle in H $_2$ O are all greater than 90°.

D. The H $\,$ – O – H $\,$ bond angle in H $_2 \rm O$ is larger than the H $\,$ – C – H $\,$ bond angle in CH $_4$

Answer: D

Solution:

C 109°28′ V N 107°

Question41

Predict the correct order among the following : (NEET - I 2016)

Options:

A. bond pair-bond pair > lone pair - bond pair lone pair - lone pair

B. lone pair - bond pair > bond pair - bond pair > lone pair - lone pair

C. lone pair - lone pair > lone pair - bond pair bond pair - bond pair

D. lone pair - lone pair > bond pair - bond pair > lone pair - bond pair

Answer: C

Solution:

Solution:

According to VSEPR theory, the repulsive forces between lone pair and lone pair are greater than between lone pair and bond pair which are further greater than bond pair and bond pair.

Question42

Decreasing order of stability of $\rm O_2$, $\rm O_2^{-}, \, O_2^{+}$ and $\rm O_2^{\ 2^{-}}$ is (2015)

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Options:

A. $O_2^{2-} > O_2^{-} > O_2 > O_2^{+}$ B. $O_2 > O_2^{+} > O_2^{2-} > O_2^{-}$ C. $O_2^{-} > O_2^{2-} > O_2^{+} > O_2$ D. $O_2^{+} > O_2 > O_2^{-} > O_2^{2-}$

Answer: D

Solution:

Solution: $O_2(16) : K K \sigma 2s^2 \sigma * 2s^2 \sigma 2p_z^2 \pi 2p_x^2 = \pi 2p_y^2 \pi * 2p_x^2 = \pi * 2p_y^2$ Bond order $= \frac{1}{2}(8-6) = 1$ $O_2^{-}(17) : K K \sigma 2s^2 \sigma * 2s^2 \sigma 2p_z^2 \pi 2p_x^2 = \pi 2p_y^2 \pi * 2p_x^2 = \pi * 2p_y^2$ Bond order $= \frac{1}{2}(8-5) = 1.5$ $O_2^{+}(15) : K K \sigma 2s^2 \sigma * 2s^2 \sigma 2p_z^2 \pi 2p_x^2 = \pi 2p_y^2 \pi * 2p_x^1$ Bond order $= \frac{1}{3}(8-3) = 2.5$ As, bond order « stability The decreasing order of stability is $O_2^{+} > O_2 > O_2^{-} > O_2^{2^{-}}$

Question43

In which of the following pairs, both the species are not isostructural? (2015)

Options:

A. Diamond, Silicon carbide

B. NH₃, PH₃

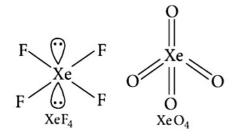
C. XeF_4 , XeO_4

D. SiCl₄, PCl₄

Answer: C

Solution:

In diamond and silicon carbide, central atom is sp^3 hybridised and hence, both are isostructural. N H ₃ and PH ₃, both are pyramidal and central atom in both cases is sp^3 hybridised SiCl ₄ and PCl ₄⁺, both are tetrahedral and central atom in both cases is sp^3 hybridised In X eF ₄, X e is sp^3d^2 hybridised and structure is square planar while in X eO₄, X e is sp^3 hybridised and structure is tetrahedral



Question44

Maximum bond angle at nitrogen is present in which of the following?

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(2015 Cancelled)

Options:

A. NO_2^+

B. NO_3^-

 $\mathrm{C.}~\mathrm{NO}_2$

D. NO_2^-

Answer: A

Solution:

Species	NO3 ⁻	<i>NO</i> ₂	NO ₂ -	NO2 ⁺
Hybridisation	sp ²	sp ²	sp ²	sp(linear)
Bond angle	120°	134°	115°	180°

So, N $\mathrm{O_2}^+$ has maximum bond angle

Question45

Which of the following options represents the correct bond order? (2015)

Options:

A. $O_2^- > O_2 < O_2^+$ B. $O_2^- < O_2 > O_2^+$ C. $O_2^- > O_2 > O_2^+$ D. $O_2^- < O_2 < O_2^+$ Answer: D

Solution:

 $O_2^- < O_2^- < O_2^+$ B.O: 1.5 2.0 2.5

Question46

The correct bond order in the following species is (2015 Cancelled)

Options:

A. $O_2^+ < O_2^- < O_2^{2+}$ B. $O_2^- < O_2^+ < O_2^{2+}$ C. $O_2^{2+} < O_2^+ < O_2^-$



D. $O_2^{2+} < O_2^{-} < O_2^{+}$

Answer: B

Solution:

 $O_2^- < O_2 < O_2^+ < O_2^{2+}$ B.O:1.5 2.0 2.5 3.0

Question47

Which of the following pairs of ions are isoelectronic and isostructural? (2015)

Options:

A. SO₃²⁻, NO₃⁻

B. ClO₃⁻, SO₃²⁻

C. CO_3^{2-} , SO_3^{2-}

D. ClO₃⁻, CO₃²⁻

Answer: B

Solution:

Species	Hybridisation	Shape	No. of <i>e</i> s
SO3 ²⁻	sp ³	Pyramidal	42
ClO ₃	sp ³	Pyramidal	42
CO3 ²⁻	sp ²	Triangularplanar	32
NO3	sp ²	Triangular planar	32

Question48

Which one of the following species has plane triangular shape? (2014)

Options:

A. N $_3$

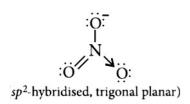
B. N O₃⁻

 $\mathrm{C.~N~O_2}^-$

D. CO_2

Answer: B

Solution:



Question49

Which of the following molecules has the maximum dipole moment? (2014)

Options:

A. CO_2

B. CH $_4$

C. N H ₃

D. N F ₃

Answer: C

Solution:

In N H $_3$, H is less electronegative than N and hence dipole moment of each N—H bond is towards N and create high net dipole moment whereas in N F $_3$, F is more electronegative than N, the dipole moment of each N—F bond is opposite to that of lone pair, hence reducing the net dipole moment.

Question50

Dipole-induced dipole interactions are present in which of the following pairs? (NEET 2013)

Options:

A. HCl and He atoms

B. SiF_4 and He atoms

C. H₂O and alcohol

D. Cl₂ and CCl₄

Answer: A

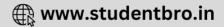
Solution:

HCl is polar ($\mu{\ne}0)$ and He is non-polar ($\mu{=}0)$ gives dipole-induced dipole interaction.

Dipole -induced dipole forces /interactions are those which take place between a polar and a non polar molecule or with an atom. H_2O and Alcohol both are polar, thus this is not an example of Dipole-Induce dipole. While HCl is a polar while helium is non-polar, thus it is an example of Dipole induce dipole. SiF₄ and He atom are both non polar. Cl₂ and CCl₄ are both non-polar molecule. (In CCl₄ chlorine atoms are bonded symmetrically with the carbon central atom thus it is non-polar similarly SiF₄ is also non-polar)

Question51

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Which of the following is paramagnetic? (NEET 2013)

Options:

- A. CN⁻
- $B. NO^+$
- C. CO
- $D. O_2^{-}$

Answer: D

Solution:

 $O_2^{-}(17)$ superoxide has one unpaired electron. $\sigma 1s^2\sigma * 1s^2\sigma 2s^2\sigma * 2s^2\sigma 2p_z^{-2}\pi 2p_x^{-2} = \pi 2p_y^{-2}\pi * 2p_x^{-2} = \pi * p_y^{-1}$

Question52

Which of the following is a polar molecule? (NEET 2013)

Options:

A. SiF_4

B. XeF_4

C. BF₃

D. SF_4

Answer: D

Solution:

 SF_4 has sp^2d --hybridisation and see-saw shape with (4bp+1lp)

$$F \\ S \\ F \\ F$$

and resultant $\mu \neq 0$

Question53

```
XeF<sub>2</sub> is isostructural with (NEET 2013)
```

Options:

A. $SbCl_3$

B. $BaCl_2$

C. TeF $_2$

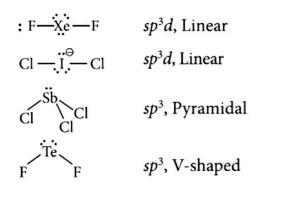
D. ICl_2^-

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Answer: D

Solution:



Question54

Which of the following is electron-deficient? (NEET 2013)

Options:

A. (BH ₃)₂

B. PH₃

C. (CH₃)₂

D. $(SiH_3)_2$

Answer: A

Solution:

Solution: Boron hydrides are electron deficient compounds.

Question55

The pair of species that has the same bond order in the following is (Karnataka NEET 2013)

Options:

A. CO, N O^+

B. N O⁻, CN ⁻

C. O₂, N $_2$

D. O₂, B₂

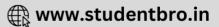
Answer: A

Solution:

```
\begin{array}{l} CO=6+8=14 \text{ electrons} \\ N \ O^{+}=7+8-1=14 \text{ electrons} \\ \text{Electronic configuration of } N \ O^{+}: \\ \sigma 1s^{2}\sigma^{*}1s^{2}\sigma 2s^{2}\sigma^{*}2s^{2}\sigma 2p_{z}^{2}\pi 2p_{x}^{2}\pi 2p_{y}^{2} \end{array}
```

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Electronic configuration of CO : $\sigma 1 s^2 \sigma^* 1 s^2 \sigma 2 s^2 \sigma^{\overline{*}} 2 s^2 \pi 2 p_x^{-2} \pi 2 p_y^{-2} \sigma 2 p_z^{-2}$ So, both have bond order = $\frac{10-4}{2} = 3$

Question56

The outer orbitals of C in ethene molecule can be considered to be hybridized to give three equivalent sp² orbitals. The total number of sigma (σ) and pi (π) bonds in ethene molecule is (Karnataka NEET 2013)

Options:

A. 3 sigma (σ) and 2 pi (π) bonds

B. 4sigma(σ) and 1pi(π) bonds

C. 5 sigma (σ) and 1 pi (п) bonds

D. 1 sigma (σ) and 2 pi (π) bonds.

Answer: C

Solution:

 $\frac{H \sigma}{H \sigma} C \frac{\pi}{\sigma} C \frac{\sigma}{\sigma} H$ 5 σ -bond and 1 π -bond

Question57

In which of the following pairs both the species have sp³ hybridization? (Karnataka NEET 2013)

Options:

A. SiF 4, BeH 2

B. N F ₃, H ₂O

C. N F $_3$, BF $_3$

D. H_2S , BF₃

Answer: B

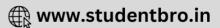
Solution:

N F $_3$ and H $_2$ O are sp 3 -hybridised.

Question58

In which of the following ionization processes the bond energy increases and the magnetic behaviour changes from paramagnetic to diamagnetic? (Karnataka NEET 2013)

Options:



A. $O_2 \rightarrow O_2^+$ B. $C_2 \rightarrow C_2^+$ C. N O \rightarrow N O⁺ D. N₂ \rightarrow N₂⁺

Answer: C

Solution:

Molecular orbital configuration of $O_2 : \sigma 1s^2 \sigma^* 1s^2 \sigma 2s^2 \sigma^* 2s^2 \sigma 2p_z^2 \pi 2p_x^2 \pi 2p_y^2 \pi^* 2p_x^1 \pi^* 2p_y^1$ ⇒ Paramagnetic Bond order = $\frac{10-6}{2} = 2$ $O_2^+: \sigma 1s^2 \sigma^* 1s^2 \sigma 2s^2 \sigma^* 2s^2 \sigma 2p_z^2 \pi 2p_x^2 \pi 2p_y^2 \pi^* 2p_x^1$ ⇒ Paramagnetic Bond order = $\frac{10-5}{2} = 2.5$ $C_2: \sigma 1s^2 \sigma^* 1s^2 \sigma 2s^2 \sigma^* 2s^2 \pi 2p_x^2 \pi 2p_y^2$ ⇒ Diamagnetic Bond order $= \frac{8-4}{2} = 2$ $C_2^+: \sigma 1s^2 \sigma^* 1s^2 \sigma 2s^2 \sigma^* 2s^2 \pi 2p_x^2 \pi 2p_y^1$ ⇒ Paramagnetic Bond order = $\frac{7-4}{2} = 1.5$ $N O: \sigma 1 s^{2} \sigma^{*} 1 s^{2} \sigma 2 s^{2} \sigma^{*} 2 s^{2} \sigma 2 p_{z}^{2} \pi 2 p_{x}^{2} \pi 2 p_{y}^{2} \pi^{*} 2 p_{x}^{-1}$ ⇒ Paramagnetic Bond order = $\frac{10-5}{2} = 2.5$ N O⁺ : $\sigma 1 s^2 \sigma^* 1 s^2 \sigma 2 s^2 \sigma^* 2 s^2 \sigma 2 p_z^2 \pi 2 p_x^2 \pi 2 p_y^2$ Bond order = $\frac{10-4}{2} = 3$ N₂: $\sigma 1s^2 \sigma^* 1s^2 \sigma 2s^2 \sigma^* 2s^2 \pi 2p_x^2 \pi 2p_y^2 \sigma 2p_z^2$ ⇒Diamagnetic ⇒ Diamagnetic Bond order $=\frac{10-4}{2}=3$ N₂⁺: $\sigma 1s^2 \sigma^* 1s^2 \sigma^2 2s^2 \sigma^* 2s^2 \pi 2p_x^2 \pi 2p_y^2 \sigma 2p_z^1$ ⇒ Paramagnetic Bond order = $\frac{9-4}{2} = 2.5$

Question59

Four diatomic species are listed below. Identify the correct order in which the bond order is increasing in them (2008, 2012 M)

Options:

A. N O < $O_2^- < C_2^{2-} < H e_2^+$ B. $O_2^- < N O < C_2^{2-} < H e_2^+$ C. $C_2^{2-} < H e_2^+ < O_2^- < N O$ D. $H e_2^+ < O_2^- < N O < C_2^{2-}$ Answer: D

Solution:

Diatomic species	Bond order
NO	2.5
<i>O</i> ₂ ⁻	1.5
C ₂ ²⁻	3.0
He_2^+	0.5

Thus increasing order : H $e_2^+ < O_2^- < N O < C_2^{2-}$

Question60

During change of O_2 to O_2^- ion, the electron adds on which one of the following orbitals? (Mains 2012)

Options:

A. π^* orbital

B. π orbital

C. σ^* orbital

D. σ orbital

Answer: A

Solution:

Electronic configuration of O₂ $\sigma(1s)^2$, $\sigma * (1s)^2 \sigma(2s)^2 \sigma * (2s)^2 \sigma(2p_z)^2 \pi (2p_x)^2 \pi (2p_x)^2 \pi (2p_y)^2 \pi * (2p_x)^1 \pi * (2p_y)^1 \pi (2p_y)^2 \pi (2p_x)^2 \pi (2p_x)^$

Thus the incoming electron will enter in π * 2px to form O_2^{-}

Question61

The pair of species with the same bond order is (2012)

Options:

A. O_2^{2-} , B_2

B. O_2^{+} , N O^{+}

C. NO , CO

D. N $_2$, O $_2$

Answer: A

Solution:

```
\begin{array}{ll} \mathrm{O}^{2^-} \rightarrow 1 & \mathrm{B_2} \rightarrow 1 \\ \mathrm{O}^{2^-} \rightarrow 2.5 & \mathrm{N} \ \mathrm{O}^+ \rightarrow 3 \\ \mathrm{N} \ \mathrm{O} \rightarrow 2.5 & \mathrm{CO} \rightarrow 3 \end{array}
```

Question62

Which of the following species contains three bond pairs and one lone pair around the central atom? (2012)

Options:

A. H $_2$ O

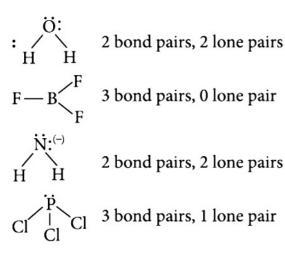
B. BF ₃

 $C. N H_2^{-}$

D. PCl₃

Answer: D

Solution:



Question63

Bond order of 1.5 is shown by (2012)

Options:

A. O_2^{+}

B. 0₂⁻

C. O₂²⁻

D. 0₂

Answer: B

Solution:

 $\begin{array}{l} \mbox{Configuration of } O_2 \\ \sigma 1 s^2 \sigma * 1 s^2 \sigma 2 s^2 \sigma * 2 s^2 \sigma 2 p_z^{-2} \pi p_x^{-2} \pi 2 p_y^{-2} \pi * 2 p_x^{-1} \pi * 2 p_y^{-1} \\ \mbox{Bond order } = \frac{No. \ of \ e^{-in \ bonding \ M.O - No. \ of \ e^{-in \ antibonding \ M.O}}{2} \ . \end{array}$

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Bond order of $O_2^{+} = \frac{10-5}{2} = 2.5$ Bond order of $O_2^{-} = \frac{10-7}{2} = 1.5$ Bond order of $O_2^{2-} = \frac{10-8}{2} = 1.0$ Bond order of $O_2 = \frac{10-6}{2} = 2$

Question64

Which one of the following pairs is isostructural (i.e., having the same shape and hybridization)? (2012)

Options:

A. [BCl₃ and BrCl₃]

B. [NH_3 and NO_3]

C. [NF₃ and BF₃]

D. [BF_4^- and NH_4^+]

Answer: D

Solution:

BCl₃ \Rightarrow sp², trigonal planar BrCl₃ \Rightarrow sp³d, T-shaped N H₃ \Rightarrow sp³, pyramidal N O₃⁻ \Rightarrow sp², trigonal planar N F₃ \Rightarrow sp³, pyramidal BF₃ \Rightarrow sp², trigonal planar BF₄⁻ \Rightarrow sp³, tetrahedral N H₄⁺ \Rightarrow sp³, tetrahedral

Question65

The pairs of species of oxygen and their magnetic behaviour are noted below. Which of the following presents the correct description? (2011 Mains)

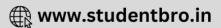
Options:

A. O ₂ ⁻ , O ₂ ²⁻	-	Both diamagnetic
B. O ⁺ , O ₂ ^{2–}	-	Both paramagnetic
C. O_2^{+} , O_2^{-}	-	Both paramagnetic
D. O, O ₂ ^{2–}	-	Both paramagnetic
Answer: C		

Solution:

 $\mathrm{O_2}^+$ and $\mathrm{O_2}$ are paramagnetic in nature as they contain one and two unpaired electrons respectively.

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Question66

Which of the following structures is the most preferred and hence of lowest energy for SO_3 ?

(Mains 2011)

Options:

В.

C.

D.

Answer: D

Solution:

No. Nas maximum number of covalent bonds.

Question67

The correct order of increasing bond length of C - H, C - O, C - C and C=C is (2011)

Options:

A. C-H < C=C < C-O < C-C B. C-C < C=C < C-O < C-H C. C-O < C-H < C-C < C=C D. C-H < C-O < C-C < C=C

Answer: A

Solution:

Increasing order of bond length is C - H < C = C < C - O < C - C107 pm 134pm 141 pm 154 pm

Question68

Which of the two ions from the list given below that have the geometry that is explained by the same hybridization of orbitals, NO_2^- , NO_3^- , NH_2^- , NH_4^+ , SCN⁻? (2011)

Options:

A. N O_2^- and N O_3^-

B. N H $_4^+$ and N O $_3^-$

C. SCN $^-$ and N H $_2^-$

D. N O_2^- and N H $_2^-$

Answer: A

Solution:

lons	Hybridisation
NO2 ⁻	sp ²
NO3 ⁻	sp ²
NH2 ⁻	sp ³
NH 4 +	sp ³
SCN ⁻	sp

Question69

Which of the following has the minimum bond length ? (2011)

Options:

A. 0₂⁺

B. O₂⁻

C. O₂²⁻

 $D.~O_2$

Answer: A

Solution:

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Electronic configuration $O_{2}: K K (\sigma 2s)^{2} (\sigma * 2s)^{2} (\sigma 2p_{z})^{2} (\pi 2p_{x})^{2} (\pi * 2p_{x})^{1} (\pi * 2p_{y})^{1}$ Bond order $= \frac{1}{2}(8 - 4) = 2$ $O_{2}^{+} \text{ Bond order } = \frac{1}{2}(8 - 3) = 2\frac{1}{2}$ $O_{2}^{-} \text{ Bond order } = \frac{1}{2}(8 - 5) = 1\frac{1}{2}$ $O_{2}^{2-} \text{ Bond order } = \frac{1}{2}(8 - 6) = 1$ As bond order increases, bond length decreases.

Question70

Some of the properties of the two species, N O_3^- and H $_3O^+$ are described below. Which one of them is correct? (2010 Mains)

Options:

A. Dissimilar in hybridization for the central atom with different structures

B. Isostructural with same hybridization for the central atom

C. Isostructural with different hybridization for the central atom

D. Similar in hybridization for the central atom with different structures

Answer: A

Solution:

No. of electron pairs at the central atom = No. of atoms bonded to it +1/ 2[Group number of central atom - Valency of the central atom ± no. of electrons] No. of electron pairs at the central atom in N O_3^- = $3 + \frac{1}{2}[5 - 6 + 1] = 3$ (sp² hybridisation) No. of electron pairs at the central atom in H $_3O^+$ $3 + \frac{1}{2}[6 - 3 - 1] = 4$ (sp³ hybridisation)

Question71

In which of the following molecules the central atom does not have ${\rm sp}^3$ hybridization ? (2010 Mains)

Options:

A. CH_4

B. SF₄

C. BF $_4^-$

D. N H $_4^+$

Answer: B

Solution:

For neutral molecules,No. of electron pairs = No. of atoms bonded to it + 1/2[Gp.no of central atom-Valency of central atom] \therefore For CH $_4$ no of e⁻ pairs = $4 + \frac{1}{2}[4 - 4] = 4(sp^3)$

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For SF $_4$ no of e^- pairs $= 4 + \frac{1}{2}[6 - 4] = 5(sp^3d)$ For ions, Nof electron pairs = NO of atoms bonded to it $+1/2[G.P. no. of central atom - Valency of central atom <math>\pm$ No of electrons] \therefore For BF $_4^-$, no of e^- pairs $= 4 + \frac{1}{2}[3 - 4 + 1] = 4(sp^3)$ hybridisation For N H $_4^+$, no of e^- pairs $= 4 + \frac{1}{2}[5 - 4 - 1] = 4(sp^3)$ hybridisation

Question72

In which one of the following species the central atom has the type of hybridization which is not the same as that present in the other three ? (2010)

Options:

A. SF $_4$

B. I₃⁻

C. SbCl $_5^{2-}$

D. PCl₅

Answer: C

Solution:

Hybridisation of the central atom can be calculated as:

 $H = \frac{1}{2}[(No. of valence electrons in the central atom) + (No. of monovalent atoms around central atom) - (Charge on cation) + (Charge on anion)]$ After calculation of hybridisation of the central atom we find that all the given species except [SbCl₅]²⁻ have central atom with sp³d (corresponding H =5) hybridization, In [SbCl₅₄]²⁻,Sb is sp³d² hybridized.

Question73

Which one of the following species does not exist under normal conditions? (2010)

Options:

A. Be_2^+

B. Be₂

C. B_2

D. Li₂

Answer: B

Solution:

Be₂ does not exist Be₂ has an electronic configuration :σ1s²σ * 1s²σ * 2s² ∴ Bond order = $\frac{4-4}{2} = 0$ Thus,Be₂ does not exist

Question74

In which of the following pairs of molecules/ions, the central atoms have sp^2 hybridisation? (2010)

Options:

A. N $\mathrm{O_2}^-$ and N H $_3$

B. BF $_{\rm 3}$ and N $\rm O_2^-$

C. N H $_2^{-}$ and H $_2O$

D. BF $_{\rm 3}$ and N H $_{\rm 2}^-$

Answer: B

Solution:

Solution:

```
The hybridisation of the central atom can be calculated as

H = \frac{1}{2}[(No. of electrons in valence shell of atom) + (No. of monovalent atoms around central atom) - (Charge on cation) + (Charge on anion)]

For

BF <sub>3</sub>, H = <math>\frac{1}{2}[(3) + (3) - (0) + (0)] = 3

\Rightarrow sp^2 hybridisation

For

N O<sub>2</sub><sup>-</sup>, H = \frac{1}{2}[(5) + (0) - (0) + (1)] = 3

\Rightarrow sp^2 hybridisation
```

Question75

In which of the following molecules/ions BF $_3$, N O $_2^-$, N H $_2^-$ and H $_2$ O, the central atom is sp² hybridised ? (2009)

Options:

A. N H $_2^-$ and H $_2\mathrm{O}$

B. N O_2^- and H $_2O$

C. BF $_3$ and N O $_2^-$

D. N O_2^- and N H $_2^-$

Answer: C

Solution:

 $BF_{3} \rightarrow sp^{2}$ $NO_{2}^{-} \rightarrow sp^{2}$ $NH_{2}^{-} \rightarrow sp^{3}$ $H_{2}O \rightarrow sp^{3}$

Question76

According to MO theory which of the lists ranks the nitrogen species in terms of increasing bond order? (2009)

Options:

A. $N_2^{2-} < N_2^{-} < N_2$ B. $N_2 < N_2^{2-} < N_2^{-}$ C. $N_2^{-} < N_2^{2-} < N_2$ D. $N_2^{-} < N_2 < N_2^{2-}$

Answer: A

Solution:

According to MOT, the molecular orbital electronic configuration of N₂: $(\sigma 1s)^2(\sigma * 1s)^2(\sigma 2s)^2(\sigma * 2s^2)(\pi 2p_x)^2(\pi 2p_y)^2(\sigma 2p_z)$ $\therefore B \cdot O = \frac{10-4}{2} = 3$ N₂⁻: $(\sigma 1s)^2(\sigma * 1s)^2(\sigma 2s)^2(\sigma * 2s)^2(\pi 2p_x)^2(\pi 2p_y)^2(\sigma 2p_z)^2(\pi * 2p_x)^2$ $\therefore B > O = \frac{10-5}{2} = 2.5$ N₂²⁻ = $(\sigma 1s)^2(\sigma * 1s)^2(\sigma 2s)^2(\sigma * 2s)^2(\pi 2p_x)^2(\pi 2p_y)^2(\sigma 2p_z)^2(\pi * 2p_x)^2(\pi * 2p_z)^2$ $\therefore B \cdot O = \frac{10-6}{2} = 2.$ Hence the order : N₂²⁻ < N₂²⁻ < N₂

Question77

What is the dominant intermolecular force or bond that must be overcome in converting liquid CH $_3$ OH to a gas ? (2009)

Options:

- A. Dipole-dipole interaction
- B. Covalent bonds
- C. London dispersion force
- D. Hydrogen bonding

Answer: D

Solution:

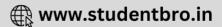
Methanol can undergo intermolecular association through H-bonding as the - OH group in alcohols is highly polarised. CH_3 CH_3 CH_3 CH_3

--- $\stackrel{1}{O}$ - H --- $\stackrel{1}{O}$ - H --- $\stackrel{1}{O}$ - H ---As a result, in order to convert liquid CH ₃OH to gaseous sian the strong hydrogen bonds must be broken.

Question78

The correct order of increasing bond angles in the following triatomic species is (2008)

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Options:

A. $N O_2^+ < N O_2 < N O_2^-$ B. $N O_2^+ < N O_2^- < N O_2$ C. $N O_2^- < N O_2^+ < N O_2$ D. $N O_2^- < N O_2 < N O_2^+$

Answer: D

Solution:

Solution: Structures of N O_2^- , N O_2^- and N O_2^+ The correct order of increasing bond angles in the following triatomic species is N $O_2^- < N O_2 < N O_2^+$

Question79

Four diatomic species are listed below in different sequences. Which of these presents the correct order of their increasing bond order? (2008)

Options:

```
A. C_2^{2-} < H e_2^+ < N O < O_2^-
B. H e_2^+ < O_2^- < N O < C_2^{2-}
C. O_2^- < N O < C_2^{2-} < H e_2^+
D. N O < C_2^{2-} < O_2^- < H e_2^+
```

Answer: B

Solution:

According to molecular orbital theory, the energy level of the given molecules are $C_2^{2^-} \rightarrow \sigma 1s^2 \sigma * 1s^2 \sigma 2s^2 \sigma * 2s^2 \pi 2p_x^2 = \pi 2p_y^2 \sigma 2p_x^2$ B. $O = \frac{1}{2}[10 - 4] = 3$ H $e_2^+ \rightarrow \sigma 1s^2 \sigma * 1s^1$ B. $O = \frac{1}{2}[2 - 1] = \frac{1}{2} = 0.5$ N $O \rightarrow \sigma 1s^2 \sigma * 1s^2 \sigma 2s^2 \sigma * 2s^2 \pi 2p_x^2 = \pi 2p_y^2 \sigma 2p_z^2 \pi * 2p_x^1$ B. $O \rightarrow \frac{1}{2}[10 - 5] = 2.5$ $O_2^- \rightarrow \sigma 1s^2 \sigma * 1s^2 \sigma 2s^2 \sigma * 2s^2 \sigma 2p_z^2 \pi 2p_x^2 = \pi 2p_y^2 \pi * 2p_x^2 \pi * 2p_y^2$ B. $O = \frac{1}{2}[10 - 7] = 1.5$ So, the correct order of their increasing bond order is $H e_2^+ < O_2^- < N O < C_2^{2^-}$

Question80

The correct order of C - O bond length among CO, CO_3^{2-} , CO_2 is (2007)

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Options:

A. $CO < CO_3^{2-} < CO_2$

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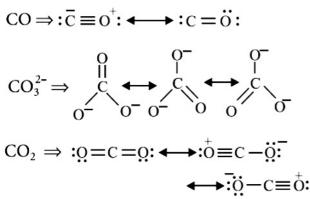
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B.
$$CO_3^{2-} < CO_2 < CO$$

C. CO < $CO_2 < CO_3^{2-}$
D. $CO_2 < CO_3^{2-} < CO$

Answer: C

Solution:



More single bond character in resonance hybrid, more is the bond length. Hence the increasing bond length is $CO < CO_2 < CO_3^{2-}$

Question81

In which of the following pairs, the two species are isostructural? (2007)

Options:

A. SO_3^{2-} and NO_3^{-}

B. BF $_{\rm 3}$ and N F $_{\rm 3}$

C. $\mathrm{BrO_3}^-$ and X $\mathrm{eO_3}$

D. SF $_4$ and X eF $_4$

Answer: C

Solution:

Hybridisation of Br in BrO_3^- : H = $\frac{1}{2}(7 + 0 - 0 + 1) = 4i.e, \text{sp}^3$ hybridisation Hybridisation of Xe in X O - 3: H = $\frac{1}{2}(8 + 0 - 0 + 0) = 4 \text{ i.e sp}^3$ hybridisation In both BrO_3^- and X eO₃, the central is sp³ hybridised and contains one lone pair of electrons, hence in both the cases, the structure is trigonal pyramidal.

Question82

Which of the following is not isostructural with SiCl $_4$ (2006)

Options:

A. N H $_4^+$

B. SCl $_4$

C. SO_4^{2-}

D. PO₄³⁻

Answer: B

Solution:

Solution:

 SCl_4 , N H₄⁺, SO_4^{2-} and PO_4^{3-} ions are the examples of molecules/ions which are of AB₄ and have tetrahedral structure SCl_4 is AB₄(lone pair) type species. Although the arrangements of five sp³d hybrid orbitals in space is trigonal bipyramidal, due to the presence of one lone pair of electron in the basal hybrid orbital, the shape of AB₄ (lone pair) species gets distorted and becomes distorted tetrahedral or see-saw.

Question83

Which of the following species has a linear shape? (2006)

Options:

A. 0₃

B. NO_2^-

C. SO_2

D. N O_2^+

Answer: D

Solution:

 NO_2^- :Due to sp² hybridisation of N-atom and the presence of one lone pair on it, NO_2^- has angular shape.

$$O_3: O_{116.8^\circ} O_{O}$$
 V-shaped

..

 SO_2 : Due to the presence of one lone pair of electrons in one of the three sp^2 – hybrid orbitals and sp^2 hybridisation of S or S⁺ atom, SO_2 molecule has aguilar (V-shaped) structure

 $\mathrm{N}~\mathrm{O_2}^+\text{:}\mathsf{Due}$ to sp hybridisation of N $^+\text{,}~\mathrm{N}~\mathrm{O_2}^+$ ion has linear shape

Question84

Which of the following is not a correct statement? (2006)

Options:

A. Multiple bonds are always shorter than corresponding single bonds



- B. The electron-deficient molecules can act as Lewis acids
- C. The canonical structures have no real existence
- D. Every AB_5 molecule does in fact have square pyramid structure.

Answer: D

Solution:

For AB_5 molecules, there are three possible geometries i.e. planar pentagonal, square pyramidal and trigonal bipyramidal. Out of these three geometries, it is only trigonal pyramidal shape in which bond pair-bond pair repulsions are minimum and hence this geometry is the most probable geometry of AB_5

Question85

Which of the following molecules has trigonal planar geometry? (2005)

Options:

A. BF ₃

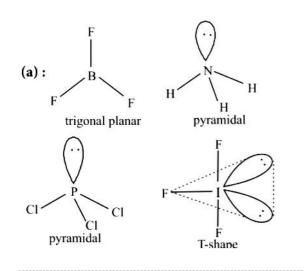
B. N H ₃

C. PCl₃

D. IF 3

Answer: A

Solution:



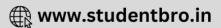
Question86

The correct order in which the $\rm O-O$ bond length increases in the following is (2005)

Options:

A. $O_2 < H_2O_2 < O_3$

- B. $O_3 < H_2O_2 < O_2$
- C. H $_{2}O_{2} < O_{2} < O_{3}$



D. $O_2 < O_3 < H_2O_2$

Answer: D

Solution:

Solution: Bond lengths of O – O in O₂ is 1.21Å, in H₂O₂ is 1.48Å and in O₃ is 1.28Å. Therefore, correct order of the O – O bond length is H₂O₂ > O₃ > O₂

Question87

The surface tension of which of the following liquid is maximum? (2005)

Options:

A. C_2H_5OH

B. CH ₃OH

C. H $_2$ O

D. C_6H_6

Answer: C

Solution:

Solution: Hydrogen bonding in H $_2$ O > C $_2$ H $_5$ OH > CH $_3$ OH Hence, H $_2$ O has maximum surface tension.

Question88

Among the following, the pair in which the two species are not isostructural is (2004)

Options:

A. SiF $_4$ and SF $_4$

B. I $\mathrm{O_3}^-$ and X $\mathrm{eO_3}$

C. BH $_4^-$ and N H $_4^+$

D. PF $_6^-$ and SF $_6^-$

```
Answer: A
```

Solution:

Solution:

SiF $_4$ has symmetrical tetrahedral shape which is due to sp³ hybridisation of the central silicon atom in its excited state configuration. SF $_4$ has distorted tetrahedral or see-saw geometry which arises due to sp³d hybridisation of central sulphur atom and due to the presence of one lone pair of electrons in one of the equatorial hybrid orbital.

Question89

In a regular octahedral molecule, M X $_6$ the number of X – M – X bonds at 180° is

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(2004)

Options:

- A. three
- B. two
- C. six
- D. four.
- Answer: A

Solution:

In octahedral molecule, six hybrid orbitals directed towards the corners of a regular octahedron with a bond angle of $90^{\rm circ}$.

Octahedral d'sp³ 90°

According to this geometry, the number of X - M - X bonds at 180° must be three.

Question90

H $_2$ O is dipolar, whereas BeF $_2$ is not. It is because (2004)

Options:

A. the electronegativity of $F\,$ is greater than that of O

B. H $_2$ O involves hydrogen bonding whereas BeF $_2$ is discrete bonding

C. H $_2\mathrm{O}$ is linear and BeF $_2$ is angular

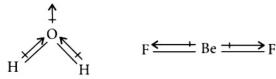
D. H $_2$ O is angular and BeF $_2$ is linear.

Answer: D

Solution:

Solution:

The overall value of the dipole moment of a polar molecule depends on its geometry and shape, i.e. vectorial addition of dipole moment of the constituent bonds. Water has angular structure with bond angle 105° as it has dipole moment. However BeF₂ is a linear molecule since dipole moment summation of all the bonds present in the molecule cancel each other.



Question91

In BrF $_3$ molecule, the lone pairs occupy equatorial positions to minimize (2004)

Options:

A. lone pair - bond pair repulsion only

B. bond pair - bond pair repulsion only

C. lone pair - lone pair repulsion and lone pair - bond pair repulsion





D. lone pair - lone pair repulsion only.

Answer: D

Solution:

Solution:

Bent T-shaped geometry in which both lone pairs occupy the equatorial positions of the trigonal bipyramid. Here (l p - l p) repulsion = 0, (l p - bp) repulsion = 4 and (bp - bp) repulsion = 2

Question92

Which one of the following statements is not correct for sigma- and pi- bonds formed between two carbon atoms? (2003)

Options:

A. Sigma-bond is stronger than a pi-bond.

- B. Bond energies of sigma- and pi-bonds are of the order of 264kJ /mol and 347kJ /mol , respectively.
- C. Free rotation of atoms about a sigma bond is allowed but not in case of a pi bond.
- D. Sigma-bond determines the direction between carbon atoms but a pi-bond has no primary effect in this regard.

Answer: B

Solution:

We know C - C = 347 kJ / molC = C = 619 kJ / mol

Question93

Which of the following has $p\pi - d\pi$ bonding? (2002)

Options:

- A. NO_3^{-}
- B. SO_3^{2-}

C. BO_{3}^{3-}

D. CO_3^{2-}

Answer: B

Solution:

In sulphite ion, the central atom sulphur is ${\rm sp}^3$ hybridised. Electronic structure of S atom in excited state



The three p electrons form σ bonds with three oxygen atoms - with one position (of the tetrahedron) being occupied by a lone pair. The d electron (excluded from hybridisation) forms π bond with one oxygen atom. i.e. $p\pi - d\pi$ bonding occurs.

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Question94

In N O_3^- ion number of bond pair and lone pair of electrons on nitrogen atom are (2002)

Options:

- A. 2,2
- B. 3,1
- C. 1,3
- D. 4,0

Answer: D

Solution:

$$o^- - \underset{O}{N} \rightarrow O$$

In N O_3^- ion, Due to the presence of one negative charge, number of valence electrons = 5 + 1 = 6. One O-atom forms two bonds (= bond) and two O-atom are shared with two electrons of N atom Thus, 30 -atoms are shared with 8 electrons of N -atom. N -atom. Number of bond pairs (or shared pairs) = 4 Number of lone pairs = 0

Question95

Which of the following is isoelectronic? (2002)

Options:

A. CO_2 , N O_2

B. N O_2^- , CO_2

C. CN ⁻, CO

D. SO₂, CO₂

Answer: C

Solution:

In CO, the number of electrons = 6 + 8 = 14[Z of C = 6 and O = 8] Electronic configuration of molecular orbital of CO : $(\sigma 1s)^2(\sigma^* 1s)^2(\sigma 2s)^2(\sigma^* 2s)^2(\pi 2p_x)^2(\sigma 2p_z)^2$ CN ⁻ have also get (6 + 7 + 1)14 electrons and the configuration is similar to that of CO. CN ⁻ and CO are isoelectronic.

Question96

Which of the following two are isostructural? (2001)

Options:

A. $X eF_{2}$, IF_{2}^{-} B. N H₃, BF₃ C. CO_{3}^{2-} , SO_{3}^{2-} D. PCl₅, I Cl₅ **Answer: A**

Solution:

Solution:

Compounds having same shape with same hybridisation are known as isostructural. X eF $_2$, IF $_2$ \rightarrow both are sp³d hybridised linear molecules.

Question97

In which of the following bond angle is maximum? (2001)

Options:

A. N H $_3$

B. N H $_4^+$

C. PCl₃

D. SCl₂

Answer: B

Solution:

Solution: Bond angle is maximum in ${\rm N\,H\,_4}^+$ tetrahedral molecule with bond angle 109°

Question98

Nitrogen forms N $_2$, but phosphorus does not form P $_2$, however, it converts P $_4$, reason is (2001)

Options:

A. triple bond present between phosphorus atom

B. $p\pi$ – $p\pi$ bonding is weak

C. $p\pi$ – $p\pi$ bonding is strong

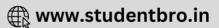
D. multiple bonds form easily.

Answer: B

Solution:

Solution: For strong π -bonding, $p\pi$ - $p\pi$ bonding should be strong. In case of P, due to larger size as compared to N -atom, $p\pi - p\pi$ bonding is not so strong.

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Question99

In X – H Y, X and Y both are electronegative elements. Then (2001)

Options:

A. electron density on \boldsymbol{X} will increase and on \boldsymbol{H} will decrease

B. in both electron density will increase

C. in both electron density will decrease

D. on X electron density will decrease and on H increases.

Answer: A

Question100

d π – pπ bond present in (2000)

Options:

A. CO₃²⁻

B. PO₄³⁻

 $C. NO_3^{-}$

D. NO_2^{-}

Answer: B

Solution:

In PO_4^{3-} , P atom has vacant d-orbitals, thus it can form $p\pi - d\pi$ bond. 'N' and 'C' have no vacant 'd' orbitals in their valence shell, so they cannot form such bond.

Question101

Right order of dissociation energy N $_2$ and N $_2^{\,+}$ is (2000)

Options:

A. N₂ > N₂⁺

B. N₂ = N₂⁺

C. N $_{2}^{+}$ > N $_{2}^{-}$

D. none.

Answer: A

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Solution:

$$\begin{split} &\mathrm{N}_{2}(14) \rightarrow (\sigma 1 \mathrm{s})^{2}, \, (\sigma^{*} 1 \mathrm{s})^{2}, \, (\sigma 2 \mathrm{s})^{2}, \, (\sigma^{*} 2 \mathrm{s})^{2} (\pi 2 \mathrm{p}_{\mathrm{x}})^{2}, \, (\pi 2 \mathrm{p}_{\mathrm{y}})^{2}, \, (\sigma 2 \mathrm{p}_{\mathrm{z}})^{2} \\ &\mathrm{In} \, \mathrm{N}_{2}, \, \mathrm{bond} \, \mathrm{order} \, = \, \frac{\mathrm{N}_{b} - \mathrm{N}_{a}}{2} = \, \frac{10 - 4}{2} = 3 \\ &\mathrm{In} \, \mathrm{N}_{2}^{+}, \, \mathrm{bond} \, \mathrm{order} \, = \, \frac{9 - 4}{2} = 2 \cdot 5 \\ &\mathrm{As} \, \mathrm{the} \, \mathrm{bond} \, \mathrm{order} \, \mathrm{in} \, \mathrm{N}_{2} \, \mathrm{is} \, \mathrm{more} \, \mathrm{than} \, \mathrm{N}_{2}^{+} \, \mathrm{so} \, \mathrm{the} \, \mathrm{dissociation} \, \mathrm{energy} \, \mathrm{of} \, \mathrm{N}_{2} \, \mathrm{is} \, \mathrm{higher} \, \mathrm{than} \, \mathrm{N}_{2}^{+} \end{split}$$

Question102

Which species does not exhibit paramagnetism? (2000)

Options:

A. N $_2^+$

B. O_2^{-}

C. CO

D. NO

Answer: C

Solution:

Solution: In 'CO' (14 electrons), there is no unpaired electron in its molecular orbital. Therefore, this does not exhibit paramagnetism.

Question103

The number of anti-bonding electron pairs in ${O_2}^{2-}$ molecular ion on the basis of molecular orbital theory is (Atomic number of O is 8) (1998)

Options:

A. 3

B. 2

C. 5

D. 4

Answer: D

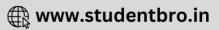
Solution:

 $O_2^{2-}(18) \rightarrow (\sigma 1s)^2$, $(\sigma^* 1s)^2 (\sigma 2s)^2$, $(\sigma^* 2s)^2 (\sigma 2p_z)^2$, $(\pi 2p_x)^2$, $(\pi 2p_y)^2$, $(\pi^* 2p_x)^2$, $(\pi^* 2p_y)^2$ Thus, the no. of antibonding electrons in O_2^{2-} ion is 8(4 pairs)

Question104

In PO_4^{3-} ion, the formal charge on each oxygen atom and P – O bond order respectively are

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(1998)

C

Options:

A. -0.75, 1.25 B. -0.75, 1.0

C. -0.75, 0.6

D. -3, 1.25

Answer: A

Solution:

The total charge = -3 So the average formal charge on each 'O' atom is -3/4 = -0.75Again total no. of electrons in the valence shell of PO_4^{3-} ion = 5 + 8 = 13No. of electrons involved in bond formation in PO_4^{3-} ion = 13 - 3 = 10 PO_4^{3-} ion = 13 - 3 = 10No. of bonds in $PO_4^{3-} = \frac{10}{2} = 5$ \Rightarrow Average P - O bond order = $\frac{5}{4} = 1.25$

Question105

N $_2$ and O $_2$ are converted into monocations, N $_2^{\ +}$ and O $_2^{\ +}$ respectively. Which is wrong? (1997)

Options:

A. In O_2^+ paramagnetism decreases.

B. N $_2^+$ becomes diamagnetic.

C. In N $_2^+$, the N - N bond weakens.

D. In O_2^{+} , the O – O bond order increases.

Answer: B

Solution:

Diamagnetism is caused due to the absence of unpaired electrons. But in N 2⁺, there is unpaired electron. So, it is paramagnetic.

Question106

N $_2$ and O $_2$ are converted into monocations, N $_2^{\ +}$ and O $_2^{\ +}$ respectively. Which is wrong? (1997)

Options:

A. In O_2^+ paramagnetism decreases.

B. N $_2^+$ becomes diamagnetic.

C. In N_2^+ , the N - N bond weakens.

D. In O_2^+ , the O – O bond order increases.

Answer: B

Question107

The bond length between hybridised carbon atom and other carbon atom is minimum in (1996)

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Options:

A. propene

B. propyne

C. propane

D. butane.

Answer: B

Solution:

Solution: The C-C bond length = 1.54Å A C = C bond length = 1.34Å and C = C bond length = 1.20Å since propyne has a triple bond, therefore it has minimum bond length.

Question108

Which of the following has sp^2 -hybridisation? (1996)

Options:

A. BeCl₂

B. C₂H₂

C. C_2H_6

D. C_2H_4

Answer: D

Solution:

Solution: BeCl $_2$ and C_2H $_2$ have sp -hybridisation and C_2H $_6$ has sp^3 -hybridisation.

Question109

Which of the following species is paramagnetic? (1995)

Options:

- A. CO
- B. CN⁻

 $C. O_2^{2-}$

D. NO

Answer: D

Solution:

Solution:

Paramagnetism is caused by the presence of atoms, ions or molecules with unpaired electrons, i.e. : $\dot{N} = \ddot{O}$:

Question110

The correct order of the O-O bond length in $\rm O_2$, H $_2\rm O_2$ and $\rm O_3$ is (1995)

Options:

A. $O_2 > H_2O_2 > O_3$

B. H $_{2}O_{2} > O_{3} > O_{2}$

C. $O_2 > O_3 > H_2O_2$

D. $O_3 > H_2O_2 > O_2$

Answer: B

Solution:

Bond length of O-O in O_2 is 1.21\AA (O=O); in H $_2O_2$ is $1.48 \text{\AA}(\text{H}\,O-\text{OH}$) and in O_3 is 1.28\AA



Question111

The ground state electronic configuration of valence shell electrons in nitrogen molecule (N $_2$) is written as K K, $\sigma 2s^2$, $\sigma^* 2s^2 \pi 2p_x^2 = \pi 2p_y^2 \sigma 2p_z^2$. Hence the bond order in nitrogen molecule is (1995)

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Options:

- A. 2
- В. З
- C. 0
- D. 1

Answer: B

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Solution:

Number of electrons in bonding orbitals N $_{\rm b}$ = 10 and number of electrons in antibonding orbitals N $_{\rm a}$ = 4 Therefore bond order = 1/2(N $_{\rm b}$ - N $_{\rm a}$) = 1/2(10 - 4) = 3

Question112

Which of the following molecules has the highest bond order? (1994)

Options:

A. 0₂⁻

B. O_2

C. O_2^{+}

D. O₂²⁻

Answer: C

Solution:

Solution: The bond order of $O_2^{+} = 2.5$, $O_2^{-2} = 1$, $O_2^{-} = 1.5$ and that of $O_2 = 2$

Question113

Which of the following molecules does not possess a permanent dipole moment? (1994)

Options:

A. CS_2

B. SO₃

C. H $_2$ S

D. SO_2

Answer: A

Solution:

Solution: The structure of CS_2 is linear and therefore it does not have permanent dipole moment. It is represented as S = C = S.

Question114

The table shown below gives the bond dissociation energies (E $_{\rm diss}$) for single covalent bonds of carbon (C) atoms with element A, B, C and D. Which element has the smallest atoms?

Bond	E _{diss} (kJ mol ⁻¹)
C-A	240
С-В	328
C-C	276
C - D	485

(1994)

Options:

A. C

- B. D
- C. A

D. B

Answer: B

Solution:

Solution:

Smaller the atom, stronger is the bond and greater the bond dissociation energy. Therefore the bond C - D has the greatest energy or smallest atoms.

Question115

Among the following which compound will show the highest lattice energy? (1993)

Options:

A. KF

B. NaF

C. CsF

D. RbF

Answer: B

Solution:

Solution:

For compounds containing ions of same charge, lattice energy increases as the size of ions decreases. Thus, NaF has highest lattice energy.

Question116

Which one of the following is the correct order of interactions? (1993)

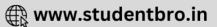
Options:

A. Covalent < hydrogen bonding < van der Waals' < dipole-dipole

B. van der Waals' < hydrogen bonding < dipole-dipole < covalent

C. van der Waals' < dipole-dipole < hydrogen bonding < covalent

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D. Dipole-dipole < van der Waals' < hydrogen bonding < covalent.

Answer: B

Solution:

Solution: (b) The van der Waals" forces are weakest forces and covalent bond is strongest, so the order of interactions is van der Waals' < H-bonding < dipole-dipole < covalent.

Question117

Which one of the following has the shortest carbon carbon bond length? (1992)

Options:

A. Benzene

B. Ethene

C. Ethyne

D. Ethane

Answer: C

Solution:

Solution:

There is a triple bond in ethyne molecule ($H - C \equiv C - H$) and due to this triple bond, carbon-carbon bond distance is shortest in ethyne.

Question118

Which structure is linear? (1992)

Options:

A. SO_2

B. CO_2

C. CO_3^{2-}

D. SO_4^{2-}

Answer: B

Solution:

Solution:

 CO_2 molecule is sp -hybridised and thus it is linear, while CO_3^{2-} is planar (sp² -hybridised), SO_2 is an angular molecule with sp² hybridisation and SO_4^{2-} is tetrahedral (sp³ – hybridised)

Question119

Strongest hydrogen bond is shown by (1992)



Options:

- A. water
- B. ammonia
- C. hydrogen fluoride
- D. hydrogen sulphide.

Answer: C

Solution:

Solution: H -F shows strongest H -bonds because fluorine is most electronegative.

Question120

In compound X , all the bond angles are exactly $109^{\circ}28^{\circ}$, X is (1991)

Options:

A. chloromethane

B. carbon tetrachloride

C. iodoform

D. chloroform.

Answer: B

Solution:

As all C - Cl bonds are directed towards the corner of a regular tetrahedron.

Question121

Among LiCl , BeCl $_{\rm 2}$, BCl $_{\rm 3}$ and CCl $_{\rm 4}$, the covalent bond character follows the order (1990)

Options:

```
A.

BeCl_2 > BCl_3 > CCl_4 < LiCl

B.

BeCl_2 < BCl_3 < CCl_4 < LiCl

C.

LiCl < BeCl_2 < BCl_3 < CCl_4

D.

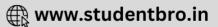
LiCl > BeCl_2 > BCl_3 > CCl_4

Answer: C
```

Solution:

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Along the period, as we move from Li \rightarrow Be \rightarrow B \rightarrow C, the electronegativity increases and hence the E N difference between the element and Cl decreases and accordingly the covalent character increases. Thus LiCl < BeCl $_2 <$ BCl $_3 <$ CCl $_4$ is correct.

Question122

The complex ion $[Co(NH_3)_6]^{3+}$ is formed by sp³d² hybridisation. Hence the ion should possess (1990)

Options:

A. octahedral geometry

B. tetrahedral geometry

C. square planar geometry

D. tetragonal geometry.

Answer: A

Solution:

Solution: According to VSEPR theory, a molecule with 6 bond pairs must be octahedral.

Question123

Which statement is not correct? (1990)

Options:

A. A sigma bond is weaker than a pi bond.

B. A sigma bond is stronger than a pi bond.

C. A double bond is stronger than a single bond.

D. A double bond is shorter than a single bond.

Answer: A

Solution:

Solution:

A sigma bond is formed by end-to-end overlap of the bonding orbitals.

Whereas, a pi - bond is formed by side-to-side overlap of (usually) two p-orbitals. This places the electron density above and below the plane of the nuclei with a node in the middle. The nodal region reduces the degree to which the orbitals overlap, and therefore, a pi bond is weaker than a sigma bond between the same two elements.

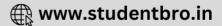
Hydrogen bonds are a strong type of dipole-dipole interaction. As a Rule of Thumb, they are weaker than covalent and ionic ("intramolecular") bonds", but stronger than most dipole-dipole interactions.

Question124

Which one shows maximum hydrogen bonding? (1990)

Options:

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A. H ₂O

- B. H₂Se
- C. H $_2$ S
- D. HF

Answer: A

Solution:

Solution:

 $\rm H_{2}O$ shows maximum H -bonding because each $\rm H_{2}O$ molecule is linked to four $\rm H_{2}O$ molecules through H -bonds.

Question125

Linear combination of two hybridized orbitals belonging to two atoms and each having one electron leads to the formation of (1990)

Options:

A. sigma bond

B. double bond

C. co-ordinate covalent bond

D. pi bond.

Answer: A

Question126

Which one of the following formulae does not correctly represent the bonding capacities of the two atoms involved? (1990)

Options:

A.

$$\begin{bmatrix} H \\ I \\ H - P - H \\ I \\ H \end{bmatrix}^{+}$$

B.
F \overbrace{O}^{F}
C.

$$H-C=C_{O-H}$$

Answer: D

Solution:

 $H - C = C = C^* - O - H$ The asterisk (*) marked carbon has a valency of 5 and hence this formula is not correct because carbon has a maximum valency of 4.

Question127

Which of the following molecules does not have a linear arrangement of atoms? (1989)

Options:

A. H₂S

B. C_2H_2

C. BeH $_{\rm 2}$

D. CO_2

Answer: A

Solution:

Solution:

For linear arrangement of atoms the hybridisation is sp (bond angle = 180°). Only H ₂S has sp³ -hybridisation and hence it has angular shape while C₂H ₂, BeH ₂ and CO₂ all involve sp -hybridisation and hence has linear arrangement of atoms.

Question128

Which of the following does not apply to metallic bond? (1989)

Options:

- A. Overlapping valence orbitals
- B. Mobile valence electrons
- C. Delocalized electrons
- D. Highly directed bonds

Answer: D

Solution:

Solution:

Metallic bonds have electrostatic attraction on all sides and hence do not have directional characteristics.

Question129





In which one of the following molecules the central atom can be said to adopt sp^2 hybridization? (1989)

Options:

- A. BeF₂
- B. BF 3
- C. C_2H_2

D. NH₃

Answer: B

Solution:

BF $_{\rm 3}$ involves sp^2 -hybridisation.

F B F F

Question130

H $_2{\rm O}$ has a net dipole moment while BeF $_2$ has zero dipole moment because (1989)

Options:

- A. H $_2\mathrm{O}$ molecule is linear while BeF $_2$ is bent
- B. BeF $_2$ molecule is linear while H $_2$ O is bent
- C. fluorine has more electronegativity than oxygen
- D. beryllium has more electronegativity than oxygen.

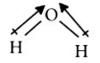
Answer: B

Solution:

 BeF_2 is linear and hence it has zero dipole moment.

$$F \longrightarrow Be \longrightarrow F$$

while H $_2\mathrm{O}$ is a bent molecule and hence it has a non-zero dipole moment



Question131

The angle between the overlapping of one ${\bf s}$ -orbital and one ${\bf p}$ -orbital is (1988)

Options:

A. 180°



B. 120°

C. 109°28′

D. 120°, 60°

Answer: A

Solution:

The type of overlapping between s- and p -orbitals occurs along internuclear axis and hence the angle is $180^\circ.$

Question132

Equilateral shape has (1988)

Options:

A. sp hybridisation

B. sp² hybridisation

C. sp³ hybridisation

D. d sp^3 hybridisation.

Answer: B

Solution:

Equilateral or triangular planar shape involves ${\rm sp}^2$ hybridisation. e . g., BCl $_3.$

