

## Hybridisation

41. (d)

**Explanation:**

Hydrogen peroxide ( $\text{H}_2\text{O}_2$ ) has a **non-planar** structure.

- Each oxygen is  $\text{sp}^3$  hybridized, forming a **bent geometry**.
- The molecule has a **dihedral angle of  $\sim 101^\circ$**  between the two O-H planes.
- $\angle \text{H}-\text{O}-\text{O} = \angle \text{O}-\text{O}-\text{H}' \approx 97^\circ$ .

Thus,  $\text{H}_2\text{O}_2$  has a **skew (non-planar)** shape, correctly described in option (d)

42. (b) 4

**Explanation:**

In **ethylene** ( $\text{C}_2\text{H}_4$ ), the two carbon atoms are connected by a **double bond** ( $\text{C}=\text{C}$ ).

- A double bond consists of **one sigma ( $\sigma$ )** and **one pi ( $\pi$ )** bond.
- Each bond involves **two shared electrons**, so total =  $2(\sigma) + 2(\pi) = 4$  **shared electrons** between the two carbons.

43. (c)  $\text{sp}^3$ ,  $\text{sp}^2$ ,  $\text{sp}$ ,  $\text{sp}^2$

**Explanation:**

Let's analyze each carbon:

1.  $\text{C}_1 (\text{CH}_3-)$  → single bonds only →  $\text{sp}^3$
2.  $\text{C}_2 (-\text{CH}=)$  → part of a double bond →  $\text{sp}^2$
3.  $\text{C}_3 (= \text{C}=)$  → part of two double bonds →  $\text{sp}$
4.  $\text{C}_4 (= \text{CH}_2)$  → part of a double bond →  $\text{sp}^2$

Hence, hybridization sequence =  $\text{sp}^3$ ,  $\text{sp}^2$ ,  $\text{sp}$ ,  $\text{sp}^2$



44. (a) Acetate ion is  $CH_3-C\begin{array}{c} \overset{O}{\parallel} \\ \backslash \\ O^- \end{array}$  i.e. one  $C-O$  single bond and one  $C=O$  double bond.

45. (c) **sp hybridized**

**Explanation:**

In **acetylene ( $C_2H_2$ )**, each carbon atom forms:

- One **sigma ( $\sigma$ )** bond with the other carbon atom
- One **sigma ( $\sigma$ )** bond with a hydrogen atom
- Two **pi ( $\pi$ )** bonds\*\* between the two carbons\*\*

Thus, each carbon is **sp hybridized**, forming a **linear molecule** with a bond angle of **180°**

46. (c) Benzene has all carbons  $sp^2$  hybridised and planar in shape.

47. (d) In methane C is  $sp^3$  hybridized and bond angle is  $109^\circ$ .

48. (b)  $120^\circ$

Explanation:  $sp^2$  hybrid orbitals are arranged trigonal-planar at  $120^\circ$

49. (c)  $H_3C^+$

Explanation: The carbocation ( $CH_3^+$ ) has three  $\sigma$ -bonds/positions and an empty p orbital  $\rightarrow sp^2$ .

50. (a) Tetrahedral

Explanation: Each C is  $sp^3$  hybridized and bonded to four other C atoms in a tetrahedral network.



51. (b)  $\text{BCl}_3$

Explanation: B in  $\text{BCl}_3$  has three  $\sigma$ -bonds and no lone pair  $\rightarrow \text{sp}^2$ , trigonal planar.

52. (a)  $\text{dsp}^2$

Explanation: The Cu(II) in the tetraammine complex is commonly square-planar ( $\text{dsp}^2$  hybridisation).

53. (c)  $\text{sp-sp}^2$

Explanation: The C–C single bond connects an sp-hybridized carbon (from the  $\equiv$ ) to an  $\text{sp}^2$ -hybridized carbon (from the =).

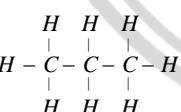
54. (d)  $\text{CH}_3-\text{C}-\text{H}$  O (i.e., the  $\text{CH}_3$  carbon is  $\text{sp}^3$ )

Explanation: A methyl carbon ( $\text{CH}_3$ ) is bonded by single bonds (four  $\sigma$ ) and is  $\text{sp}^3$  hybridized. (Interpreted the options so the  $\text{CH}_3$ -containing choice is the one with  $\text{sp}^3$  carbon.)

55. (b)  $\text{sp}^2$

Explanation: The  $120^\circ$  angle and trigonal arrangement around B indicate  $\text{sp}^2$  hybridisation for boron.

56. (d)



There are 10 shared pairs of electrons.

57. (b)  $1/2$

**Explanation:**

In  $\text{sp}$  hybridisation, one s orbital mixes with one p orbital, forming two equivalent hybrid orbitals.

Therefore,

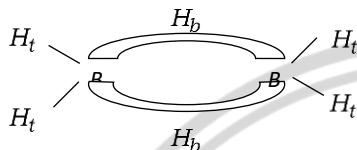


s-character = 1/2

p-character = 1/2

58. (a) The diborane molecule has two types of  $B - H$  bond :

- (i)  $B - H_t$  – It is a normal covalent bond.
- (ii)  $B - H_b$  – It is a three centred bond.



59. (a)  $sp^3$

**Explanation:** The circled carbon in  $\text{CH}_3-\text{OCl}$  is bonded to three hydrogen atoms and one oxygen atom through single ( $\sigma$ ) bonds. Hence, it uses  $sp^3$  hybrid orbitals in bond formation.

60. (c)  $\text{H}_2\text{O}_2 > \text{O}_3 > \text{O}_2$

**Explanation:**

- In  $\text{O}_2$ , the bond order = 2 → shortest bond.
- In  $\text{O}_3$ , the bond order = 1.5 → intermediate bond length.
- In  $\text{H}_2\text{O}_2$ , the bond order = 1 → longest bond.

Therefore, bond length order is  $\text{H}_2\text{O}_2 > \text{O}_3 > \text{O}_2$ .

