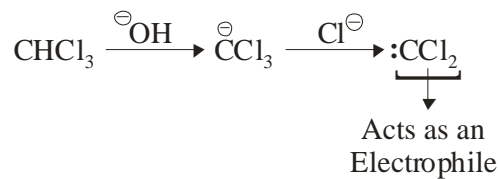


## PART-B : CHEMISTRY

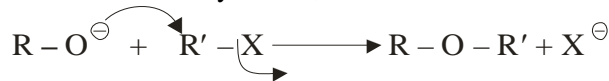
31. (B)  
Oxidation state of P is getting changed from  $-3$  to  $+5$  during this conversion. That is, P in  $\text{Cu}_3\text{P}$  is getting oxidized to  $\text{H}_3\text{PO}_4$ .
32. (C)  
 $\text{PH}_3$  is a Drago's compound in which bond angle is approximately  $90^\circ$ .
33. (B)  
Here Mn is in highest oxidation state ( $+7$ ), so no d-electrons are there. So the color arises from ligand to metal charge transfer transition.
34. (D)  
Because Bi do not show  $+5$  oxidation state, because of Inert Pair Effect.
35. (A)  
Due to the formation of tetraamminezinc(II) complex:  
 $\text{Zn}^{2+} + \text{NH}_4\text{OH} \longrightarrow [\text{Zn}(\text{NH}_3)_4]^{2+}$

36. (C)



37. (C)

In Williamson's synthesis,



Thus, it involves  $\text{S}_{\text{N}}2$  mechanism.

38. (C)

39. (D)

$$\text{Miliequivalents of } \text{H}^+ = 75 \times \frac{1}{5} = 15$$

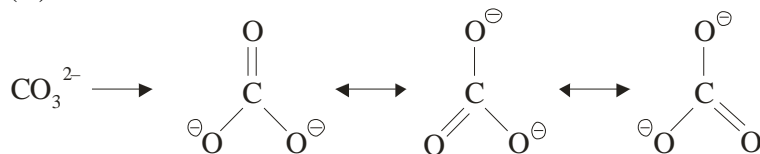
$$\text{Miliequivalents of } \text{OH}^\ominus = 25 \times \frac{1}{5} = 5$$

$$\text{Net Miliequivalents of } \text{H}^+ = 10$$

$$\text{Concentration of } \text{H}^+ = \frac{10}{100} = 0.1 = 10^{-1}$$

$$\begin{aligned} \text{pH} &= -\log [\text{H}^+] \\ &= -\log (0.1) \\ &= 1 \end{aligned}$$

40. (D)



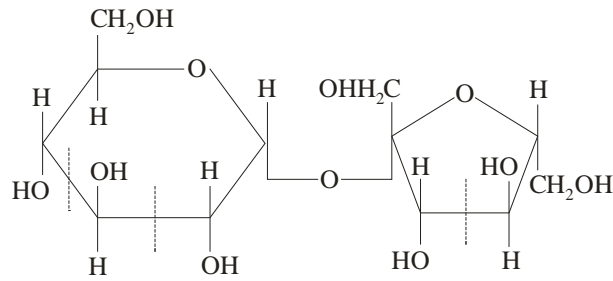
Due to backbonding in  $\text{N}(\text{SiH}_3)_3$ , it has planar structure.

In  $\text{BCl}_3$ ; B and in  $\text{CO}_3^{2-}$  C has  $\text{sp}^2$  hybridisation.

41. (B)

Due to inter electronic repulsions

42. (C)



43. (C)

$$\log \frac{x}{m} = \log k + \frac{1}{n} \log P$$

$$\log \frac{x}{m} = 0.699 + (\tan 45^\circ) (\log (0.5))$$

$$\log \frac{x}{m} = 0.3979 = 0.40$$

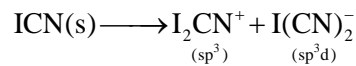
$$\frac{x}{m} = 2.5$$

$$x = 2.5 \text{ g/g adsorbent}$$

44. (D)

Cerussite ( $\text{PbCO}_3$ ) undergoes calcination.

45. (A)



46. (B)

47. (C)

$$550 = \left(\frac{1}{4}\right)(P_X^0) + \left(\frac{3}{4}\right)(P_Y^0)$$

$$2200 = P_X^0 + 3P_Y^0 \quad \dots\dots\dots \text{(A)}$$

$$560 = \left(\frac{1}{5}\right)(P_X^0) + \left(\frac{4}{5}\right)(P_Y^0)$$

$$2800 = P_X^0 + 4P_Y^0 \quad \dots\dots\dots \text{(B)}$$

On solving A and B, we get,

$$P_X^0 = 400 \text{ and } P_Y^0 = 600$$

48. (D)

49. (C)

$$\frac{Q}{F} = \text{moles} \cdot n_f$$

$$\frac{i \times t}{F} = \text{moles} \cdot n_f$$

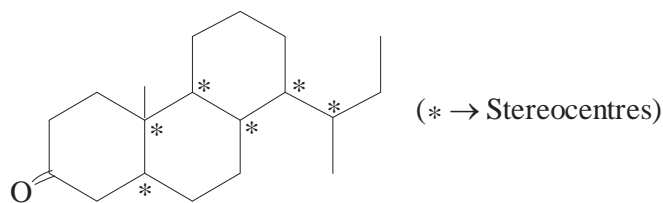
$$\frac{2 \times 5 \times 3600}{96500} = \frac{22.2}{177} \cdot n_f$$

$$\Rightarrow n_f = 3$$

Thus, oxidation state of metal in the metal salt is +3.

50. (B)  
CaF<sub>2</sub> has fluorite type structure in which coordination number of Ca<sup>2+</sup> is 8 and of F<sup>-</sup> it is 4.

51. (C)



52. (A)  
Effect of densities and temperature will nullify each other and pressure will be same for both.

$$d = \frac{PM}{RT}$$

$$\text{So } P = \frac{dRT}{M}$$

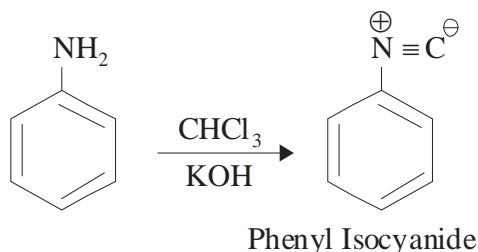
53. (B)

$$\text{pH} = \frac{\text{pK}_1 + \text{pK}_2}{2}$$
$$= \frac{2.15 + 7.2}{2} = 4.675$$

54. (C)

More the electronegative elements attached on P, more is the partial positive charge, more is the extent of back-bonding, more is the planarity. Since, F is the most electronegative among these, so P<sub>3</sub>N<sub>3</sub>F<sub>6</sub> is most planar.

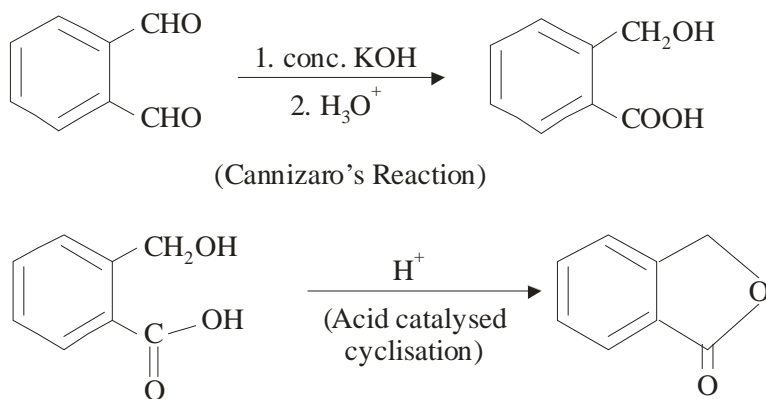
55. (A)



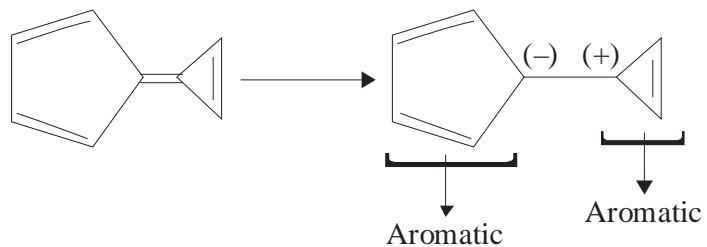
56. (C)

Steric inhibition of resonance is expected in case of substitution at ortho-position. So, (C) is correct.

57. (A)

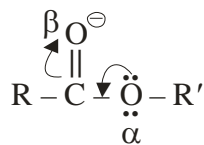


58. (B)



59. (B)

60. (B)



Hence, O denoted by  $\beta$  is more basic due to negative charge.