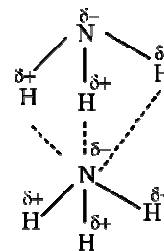


Daily Tutorial Sheet-4

Level – 1

- 46.(B)** Nitrogen is third most electronegative and hydrogen is highly electropositive and thus strong dipoles develop in NH_3 molecule causing strong dipole-dipole interactions. Note that carbon is less electronegative as compared to nitrogen and hence the dipoles (positive – negative charge centers developed in CH_4 molecule are not as strong as they are in NH_3 and in CO_2 the dipoles developed will be even weaker as both oxygen and carbon are electronegative atoms and hence the electronegativity differences will be less compared to NH_3 and CH_4 .



- 47.(B)** units of $a = \text{L}^2 \text{ atm mol}^{-2}$

$$\text{units of } b = \text{L mol}^{-1} \quad \therefore \quad \text{units of } \frac{a}{b} = \text{L atm mol}^{-1}$$

- 48.(C)** Real gases behave ideally at low pressure and high temperature

- 49.(D)** Low pressure region $Z = \left(1 - \frac{a}{RTV}\right)$

- 50.(C)** $Z = 1 + \frac{Pb}{RT} \Rightarrow Z = 1 + \frac{b}{(RT/P)} = \left(1 + \frac{b}{V}\right)$

- 51.(A)** van der Waal's equation $\left(P + \frac{an^2}{V^2}\right)(V - nb) = nRT$

$$\text{For one mole } \left(P + \frac{a}{V^2}\right)(V - b) = RT$$

$$\text{At low } P, V \text{ is very high } \Rightarrow V - b \approx V$$

$$\text{So equation becomes } \left(P + \frac{a}{V^2}\right)V = RT$$

- 52.(D)** $U_{\text{avg}} = \sqrt{\frac{8RT}{\pi m}} \Rightarrow U_{\text{avg}}^2 \times \frac{\pi M}{8} = RT;$

$$KE = \frac{3}{2}RT = \left(\frac{3\pi M}{16}\right)U_{\text{avg}}^2$$

- 53.(D)** All the three states co-exist at triple point.

- 54.(C)** $P_{\text{Real}} = P_i - \frac{n^2 a}{V^2}$

$$a \uparrow P_r \downarrow$$

$$P_2 < P_3 < P_1$$

- 55.(B)** At high P , $P \gg \frac{n^2 a}{V^2}$. So 'a' can be neglected.

- 56.(A)** At constant P & n , $V \propto T$

$$\therefore \quad \text{For a 10\% volume increase, there is 10\% increase in temperature.}$$

- 57.(CD)** at constant volume

$$P \propto T$$

$$P = KT$$

$$P = K(t^\circ\text{C} + 273.15)$$

at constant P

$$V \propto T$$

$$V = KT$$

$$V = K(t^\circ\text{C} + 273.15)$$

58.(C) Van der Waal equation : $\left(P + \frac{n^2 a}{V^2}\right)(v - nb) = nRT$

59.(D) $Z = \frac{PV}{RT}$

60.(C) At high pressure, $P \gg \frac{an^2}{V^2}$

\therefore Van der Waal equation becomes
 $P(v - nb) = nRT$