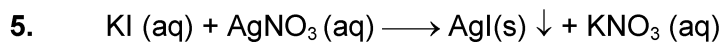


### Answer Key

- |  |                |               |                |         |
|--|----------------|---------------|----------------|---------|
| 1. (D)   | 2. (B)         | 3. (D)        | 4. (C)         | 5. (D)  |
| 6. (C)   | 7. (A)         | 8. (BD)       | 9. (ABC)       | 10. (B) |
| 11. (C)  | 12. (2)        | 13. (2)       | 14. (160)      |         |
| 15. (0.04 mol)   | 16. (23.25min) | 17. (5 times) | 18. (M = 53.8) |         |
| 19. (H <sub>20</sub> P, 45560 amu)   | 20. (18.27%)   | 21. (1669.9)  |                |         |
| 22. (m <sub>L</sub> = 112.7)   |                |               |                |         |
| 23. (i) Tetraamminedichloroplatinum(IV) chloride,<br>(ii) Volume of NH <sub>3</sub> (aq.) needed = 2 mL,<br>(iii) Diamminesilver(I) Chloride |                |               |                |         |

### Solution

1. (D)      2. (B)
3. 40 mL O<sub>3</sub> dissolve in 100 g water at 300 K and 1 atm  
 40 × 4 mL O<sub>3</sub> dissolve in 400 g water at 300 K and 1 atm  
 ∴ m.m. P so (40 × 4) × 4 mL O<sub>3</sub> dissolve in 400 g water at 300 K and 4 atm
- $$n_{\text{O}_3 \text{ dissolve}} = \frac{4 \times 640 \times 10^{-3}}{0.0821 \times 300} = 0.1$$
- or mass of O<sub>3</sub> = 4.8 g
4. Let n<sub>B</sub> mole of B present in 1 mole of mixture that has been vaporized. Thus,  $y_B = \frac{n_B}{1}$
- Mole fraction of B in the remaining liquid phase will be  $x_B = \frac{1-n_B}{1}$
- $$x_B = \frac{P - P_T^\circ}{P_B^\circ - P_T^\circ} \quad \dots\dots(1)$$
- $$[P = P_T^\circ + (P_B^\circ - P_T^\circ) x_B]$$
- and  $y_B = \frac{P_B}{P} \Rightarrow \frac{P_B^\circ x_B}{P} \quad \dots\dots(2)$
- After substitution of values of x<sub>B</sub> and y<sub>B</sub> in (1) and (2)
- we get  $1 - n_B = \frac{P - P_T^\circ}{P_B^\circ - P_T^\circ} \quad \dots\dots(3)$
- and  $n_B = \frac{(1 - n_B) P_B^\circ}{P} \quad \dots\dots(4)$
- or  $n_B = \frac{P_B^\circ}{P + P_B}$
- so  $1 - \frac{P_B^\circ}{P + P_B} = \frac{P - P_T^\circ}{P_B^\circ - P_T^\circ}$
- $$\Rightarrow P = \sqrt{P_B^\circ \cdot P_T^\circ} = \sqrt{100 \times 900}$$
- $$\Rightarrow 300 \text{ torr}$$



	KI	+	AgNO <sub>3</sub>	→	KNO <sub>3</sub>	+	AgI↓
Initial:	0.3 mole		0.2mole		–		–
Final:	0.1 mole		–		0.2		0.2

Total moles of solute ions =  $(0.1 + 0.2) \times 2$  ( $\because i = 2$ )

$$[\text{Solute}] = \frac{0.6}{4} \text{M}$$

$$\Delta T_f = K_f \cdot \frac{0.6}{4}$$

$$= 1.86 \times \frac{0.6}{4} = 0.279 \text{K}$$

6. Given that,

$$L_f = 1436.3 \text{ cal/mol} = \frac{1436.3}{18} \text{ cal/g}$$

$$T_f = 273 \text{K}$$

$$K_f = \frac{RT_f^2}{1000l_f} = \frac{2 \times 273 \times 273}{1000 \times \frac{1436.3}{18}} = 1.87$$

$$\therefore \Delta T_f = \frac{1000 \times K_f \times W}{m \times W}$$

$$= 1000 \times 1.87 \times \frac{0.2}{0.8 \times 18} = 25.97^\circ \text{C}$$

$$\therefore T_f = 0 - 25.97 = -25.97^\circ \text{C}$$

7. In vaporization of a pure solvent at a constant temperature,  $\Delta S$  will be maximum.

8. (BD)

9.  $P = MRT$

$$P = \frac{n}{V} \times RT$$

$$P = \frac{w}{V \times m} \times RT$$

$$\frac{P}{d} = \frac{RT}{m}$$

10.  $P = P_A^0 X_A + P_B^0 X_B$

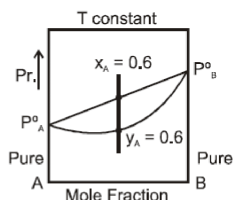
$$\Rightarrow 500 \times 0.6 + 800 \times 0.4$$

$$\Rightarrow 620 \text{ torr}$$

$$y_A = \frac{P_A}{P} \Rightarrow \frac{300}{620}$$

$$\Rightarrow 0.48 ; y_B = 0.52$$

11.  $P_A^0 = 500$  ;  $P_B^0 = 800$  torr



When most of the liquid has vaporized  $x_A = 0.6$  (given) would be  $y_A = 0.6$

$$y_A = \frac{P_A^0 x_A}{P_A^0 x_A + P_B^0 (1 - x_A)} \Rightarrow 0.6 = \frac{500 x_A}{500 x_A + 800 (1 - x_A)}$$

$$x_A = 0.70; x_B = 0.30$$

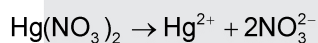
12. (2) 13. (2) 14. (160)

15. The expression for the molality of  $\text{Hg}(\text{NO}_3)_2$  solution is  $m = \frac{\Delta T_f}{K_f} = \left( \frac{0.0558}{1.86} = 0.03 \text{ m} \right)$ .

But 1000 g contains  $= \frac{3.24}{324} = 0.01$  moles of  $\text{Hg}(\text{NO}_3)_2$ .

The observed molality is thrice of the calculated value.

Thus, each molecule of  $\text{Hg}(\text{NO}_3)_2$  dissociates to give three ions.



The molality of  $\text{HgCl}_2$  solution is  $\frac{0.0744}{1.86} = 0.04 \text{ m}$ .

The number of moles of  $\text{HgCl}_2 = \frac{10.84}{271} = 0.04 \text{ mol}$ .

The observed value is equal to the calculated value. Hence, the molecules of  $\text{HgCl}_2$  do not undergo dissociation.

16. Density of methanol is 0.8 g/mL

2.5 L of methanol corresponds to a mass of  $0.8 \times 2500 = 2000 \text{ g}$

The molar mass of methanol is 32 g/mol

The number of moles of methanol  $= \frac{2000}{32} = 62.5 \text{ moles}$ .

Volume of water is 10 L

Density of water is 1 kg/L

Mass of water will be 10 kg

Molality of methanol will be  $\frac{62.5}{10} = 6.25 \text{ m}$

The depression in the freezing point  $\Delta T_f = K_f \times m$

$\Delta T_f = 1.86 \times 6.25 = 11.625^\circ\text{C}$

The rate of decrease of temperature  $= 0.5^\circ\text{C/min}$

The time for which there will be no danger to the radiator of the car  $= \frac{11.625}{0.5} = 23.25 \text{ min}$

17. We have the equation  $\pi = iCRT$

As urea is non-electrolyte,  $i = 1$

$$\pi_{\text{urea}} = iC_{\text{urea}} RT_1 \quad \dots\dots (1)$$

$$\pi_{\text{solution}} = iC_{\text{solution}} RT_2 \quad \dots\dots (2)$$

Dividing Equation 1 and 2,

$$\frac{\pi_{\text{urea}}}{\pi_{\text{solution}}} = \frac{C_{\text{urea}} T_1}{C_{\text{solution}} T_2}$$

$$\frac{500 \times 298}{105.3 \times 283} = \frac{C_{\text{urea}}}{C_{\text{solution}}}$$

We get,

$$5C_{\text{solution}} = C_{\text{urea}}$$

This means the solution was diluted 5 times with respect to the urea concentration.

18. Wt. of Solute (Urea) = 40 g =  $W_2$

$$\text{Wt. of Solvent (Water)} = 300 \text{ g} = W_1$$

$$\text{Mol. wt. of Solvent} = 18 \text{ g/mol} = M_1$$

Now, we know that, Relative lowering of Vapour Pressure =  $\frac{\text{Loss in weight of Solvent or Water bulbs}}{\text{Gain in weight of Sulphuric acid tube}}$

$$\frac{\Delta P}{P^0} = \frac{0.087}{2.036} = 0.0427$$

Now, Using Raoult's Law:

$$M_2 = \frac{W_2}{W_1} \times M_1 \times \frac{p^0}{\Delta p} = \frac{40}{300} \times 18 \times \frac{1}{0.0427} = 53.7 \text{ g/mol}$$

19.  $\Delta T_b = i \times K_f \times m$

$$5.93 \times 10^{-3} = \frac{(x+1) \times 0.52 \times 0.25 \times 1000}{M \times 10}$$

$$\frac{(x+1)}{M} = 4.56 \times 10^{-4} \quad \dots(i)$$

$$\frac{M}{100} = 23x \quad \dots(ii)$$

From equation (i) and (ii)

$$X = 20.49 \cong 20$$

Formula of protein =  $H_{20}P$

$$M = (2300 \times 20 - 20 \times 23) + 20 = 45560 \text{ amu}$$

20.  $\frac{0.5}{M} = 3.75 \times 10^{-3}$

$$M = 133.33$$

$$0.165 = (1 + \alpha) \times 1.86 \times 1.5 \times \frac{1000}{133.33 \times 150}$$

$$\alpha = 0.1827 = 18.27\%$$

21.  $\Delta H = 1659.9 \text{ Cal. at } 80 \text{ K}, \Delta H = R [313.7 \times 2.303 + 1.40655T]$

$$\frac{d \ln P}{dT} = \frac{\Delta H}{RT^2} \quad \dots (i)$$

$$\log P = 3.54595 - \frac{313.7}{T} + 1.40655 \log T$$

$$\ln P = 3.54595 \times 2.303 - \frac{313.7}{T} \times 2.303 + 1.40655 \ln T$$

$$\frac{d \ln P}{dT} = \frac{313.7 \times 2.303}{T^2} + \frac{1.40655}{T} \quad \dots (ii)$$

Comparing equation (i) & (ii)

$$\Delta H = R[313.7 \times 2.303 + 1.40655T]$$

at  $T = 80 \text{ K}$

$$\Delta H = 1669.9 \text{ cal.}$$

22. The relation is:

$$\frac{W_1}{W_2} = \frac{M_1 \times P_1^0}{M_2 \times P_2^0}$$

$$M_1 = \frac{2.5 \times 18 \times 526}{1 \times (736 - 526)} = 112.7 \text{ g/mol}$$

23.  $[\text{Pt}(\text{NH}_3)_4\text{Cl}_2]\text{Cl}_2 \rightarrow [\text{Pt}(\text{NH}_3)_4\text{Cl}_2]^{2+} + 2\text{Cl}^-$

$$\Delta T_f = i \times K_f \times m$$

$$0.0054 = i \times 1.80 \times 0.001$$

The van' Hoff factor,  $i = 3$

This suggests that 1 molecule of  $[\text{Pt}(\text{NH}_3)_4\text{Cl}_4]$  dissociates in solution to produce 3 ions.

Hence, the correct formula of the complex is  $[\text{Pt}(\text{NH}_3)_4\text{Cl}_3]\text{Cl}_2$



*Power of real gurus*