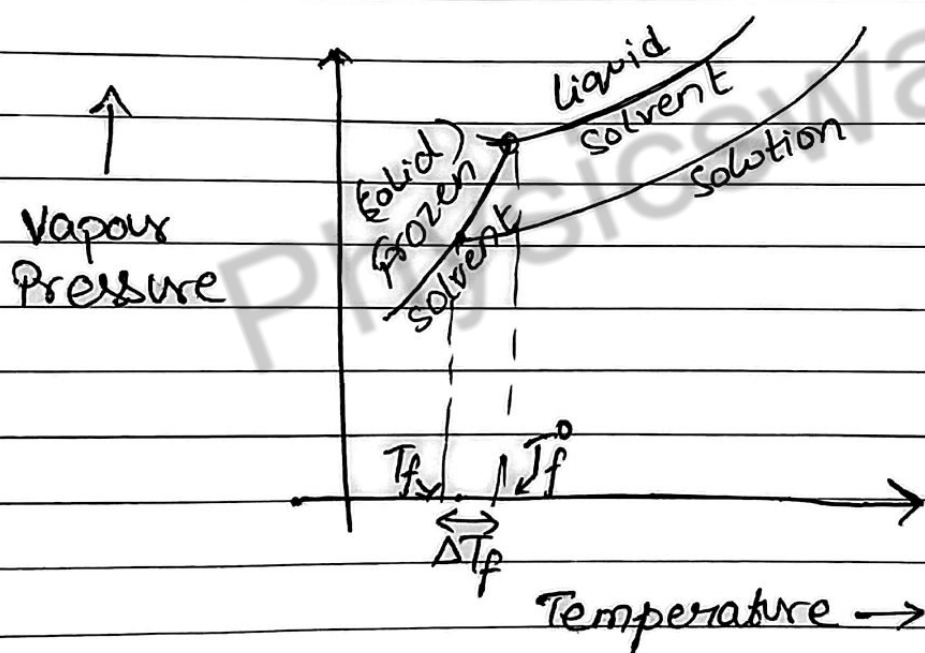


Depression in Freezing Point.

Freezing Point is the temperature at which the vapour pressure of a substance in its liquid phase becomes equal to the vapour pressure in the solid phase.

⇒ we also know, On addition of non volatile solute the vapour pressure of a solution decreases.



from the graph we can see,

- i) vapour pressure of non-volatile solute containing solution is lower than vapour pressure of pure liquid solvent.
- ii) Freezing Point of pure solvent is T_f^0
(vapour pressure of solvent = V.P. of frozen solvent)

ΔT_f is a Colligative property as it depends upon no of solute particles and not on nature of solute particles (2)

iii) Freezing Point of Solution is T_f

(Vapour Pressure of Solution = Vapour Pressure of Solid Solvent)
 $T_f < T_f^\circ$

Depression in Freezing point
 $\Delta T_f = T_f^\circ - T_f$

Experimentally, we find

$\Delta T_f \propto m$ (Colligative property)
Molality of solution

$$\Delta T_f = K_f m$$

Molal Depression Constant
Cryoscopic Constant

$K_f \rightarrow$ depends on solvent

$K_f \rightarrow$ unit $K \text{ kg/mol}$

ex: K_f for water is $1.86 K \text{ kg/mol}$

m (molality) = $\frac{\text{no of moles of non volatile solute}}{\text{Mass of Solvent (in kg)}}$

Q1) How much C_2H_5OH should be added to 1L H_2O so that it will not freeze at $-2^\circ C$?
($K_f = 1.86$) (density of water = 10^3 kg/m^3)

Solution: a) 20g b) 50g c) 80g d) 100g

(3)

1 L water = 1 kg water

 C_2H_5OH

Molar Mass = 46

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

$$2 = 1.86 \times \frac{\text{no of moles of solute}}{\text{Mass of solvent (in kg)}}$$

$$2 = 1.86 \times \frac{x}{46}$$

$$11 \text{ kg}$$

$$x = 49.4 \text{ g} \approx 50 \text{ g}$$

Q2) The freezing point of a solution containing 50 cm^3 of ethylene glycol in 50 g of water is found to be -34°C . Assuming ideal behaviour, calculate the density of ethylene glycol. (K_f for water = $1.86^\circ\text{C kg/mol}$)

a) 1.02 g/cm^3 b) 2.12 g/cm^3 c) 1.13 g/cm^3 Solution:

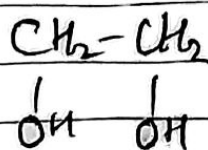
$$\Delta T_f = K_f \times m$$

$$34 = 1.86 \times \frac{\text{no of moles of solute}}{\text{Mass of solvent (in kg)}}$$

Let mass of ethylene glycol be x

$$34 = 1.86 \times \frac{x}{62}$$

$$\frac{50}{1000}$$



Molar Mass = 62

$$x = 56.66 \text{ g}$$

$$\text{density of ethylene glycol} = \frac{\text{mass}}{\text{volume}} = \frac{56.66 \text{ g}}{50 \text{ cm}^3} = 1.13 \text{ g/cm}^3$$

Q) A 5% solution (by mass) of cane sugar in water has freezing point of 271K . Calculate the freezing point of 5% glucose in water if freezing point of pure water is 273.15K .

Cane Sugar	$\text{C}_{12}\text{H}_{22}\text{O}_{11}$	($M=342$)
Glucose	$\text{C}_6\text{H}_{12}\text{O}_6$	($M=180$)

Solution:

$$\Delta T_f = K_f \times m \quad (\text{for cane sugar})$$

$$2.15 = K_f \times m \quad 5\% = 5\text{g solute in } 100\text{g solution}$$

$\Rightarrow 5\text{g cane sugar} + 95\text{g water}$

$$2.15 = K_f \times \frac{5}{\frac{342}{1000}} \quad \text{--- (i)}$$

$$\Delta T_f = K_f \times m \quad (\text{for glucose})$$

$$\Delta T_f = K_f \times \frac{5}{\frac{180}{1000}} \quad \text{--- (ii) (5g glucose + 95g water)}$$

$$(ii) \div (i)$$

$$\frac{\Delta T_f}{2.15} = \frac{342}{180}$$

$$\Delta T_f = 4.09$$

$$T_f = T_f^0 - 4.09 = 273.15 - 4.09 = 269.06\text{K}$$

~~R_f~~ K_f is not given (rare case)

$$K_f = \frac{R \times (\overset{\text{in Kelvin}}{T_f^\circ})^2 \times M}{1000 \times \Delta H_{\text{fusion}}}$$

$R \rightarrow$ Universal Gas Constant (8.314 J/mol K
(value as per unit of ΔH_{fusion}) ^{OR} 2 calorie/mol K)

$T_f^\circ \rightarrow$ Freezing Point of Solvent (in Kelvin)

$M \rightarrow$ Molar Mass of Solvent

$\Delta H_{\text{fusion}} \rightarrow$ Molar Enthalpy of Fusion.

$$(\Delta H_{\text{fusion}} = M \times L_{\text{fusion}})$$

\downarrow
Molar Mass
of Solvent

\rightarrow Latent Heat of
fusion

Q3) Calculate Molar depression Constant of a solvent which has Freezing Point at 16.6°C and Latent Heat of fusion 180.75 J/g

Solution

$$T_f^\circ = 16.6 + 273 = 289.6 \text{ K}$$

Let M
be Molar
Mass
of
Solvent

$$\Delta H_{\text{fusion}} = M \times L_f = M \times 180.75 \text{ J/g}$$

$$R = 8.314 \text{ J/mol K}$$

$$K_f = \frac{R \times (T_f^\circ)^2 \times M}{1000 \times \Delta H_{\text{fusion}}} = \frac{8.314 \times (289.6)^2 \times M}{1000 \times M \times 180.75}$$

$$\boxed{K_f = 3.857 \text{ K kg/mol}}$$