

1

$$\text{MASSA} = 2.56 \text{ gr}$$

$$V = 500 \text{ mL} \quad \text{H}_2\text{O}$$

$$\text{DENSITA}' = d = 1 \text{ gr/cm}^3$$

$$\boxed{\text{cm}^3 = \text{mL}}$$

$$\text{molalite}' = \frac{\text{moli SOLUTO}}{\text{MASSA SOLVENTE}}$$

$$\text{PM}_{\text{ACIDO}} = 118.10 \text{ gr/mol}$$

$$n_{\text{ACIDO}} = \frac{\text{MASSA}}{\text{PM}_{\text{ACIDO}}} = \frac{2.56}{118.10} =$$

$$= 2.17 \cdot 10^{-2} \text{ moli}$$

$$\text{MASSA SOLVENTE} = V \cdot d = 500 \cdot 1$$

$$= 500 \text{ gr} = 0.5 \text{ Kg}$$

$$\text{molalita}^- = \frac{2.17 \cdot 10^{-2}}{0.5} = \underline{\underline{4.34 \cdot 10^{-2} \text{ m}}}$$

FRAZIONE MOLARE

$$n_{\text{H}_2\text{O}} = \frac{500}{18} = 27.7 \text{ mol}$$

$\rightarrow PM_{\text{H}_2\text{O}}$

$$\begin{aligned} \chi_{\text{ACIDO}} &= \frac{n_{\text{ACIDO}}}{n_{\text{ACIDO}} + n_{\text{H}_2\text{O}}} = \frac{2.17 \cdot 10^{-2}}{2.17 \cdot 10^{-2} + 27.7} \\ &= 7.83 \cdot 10^{-4} \end{aligned}$$

PERCENTUALE IN PESO

$$\begin{aligned} \% \text{ PESO ACIDO} &= \frac{M_{\text{ACIDO}}}{M_{\text{ACIDO}} + M_{\text{H}_2\text{O}}} \cdot 100 \\ &= \frac{2.56}{2.56 + 500} \cdot 100 \\ &= 0.51 \% \end{aligned}$$

②

$$[\text{CO}_2] = 0.0506 \text{ m}$$

$$K_H = 4.48 \cdot 10^{-5} \text{ M/mmHg}$$

$$P_{CO_2} = K_H / \Delta$$

APPROSSIMO $m \rightarrow M$

1000 gr di SOLUTO $H_2O \approx 1 L$

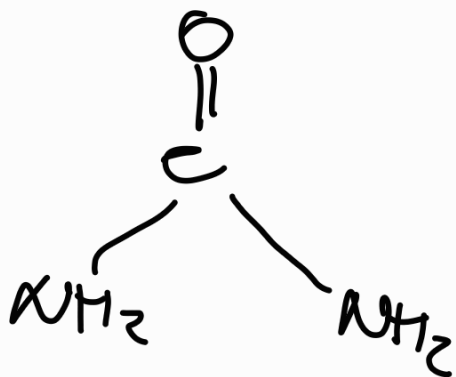
QUINDI $0.0506 m \approx 0.0506 M$

$$P_{CO_2} = K_H / 0.0506 = 1129.44 \text{ mmHg}$$

$$= \underline{\underline{1.49 \text{ atm}}}$$

(3)

UREA



$$PM_{UREA} = 60.06 \text{ gr/mol}$$

0.5 kg H_2O

$$n_{UREA} = \frac{MASS}{PM_{UREA}} = \frac{15}{60.06} = 0.25 \text{ mol}$$

$$\text{molalità UREA} = \frac{m_{\text{UREA}}}{\text{MASSA H}_2\text{O}}$$

$$= \frac{0.25}{0.5} = 0.5 \text{ m}$$

$$\Delta T_{\text{EB}} = K_b \cdot m_{\text{UREA}} = 0.5121 \cdot 0.5$$

$$= \underline{\underline{0.26^\circ \text{C}}}$$

④

$$\Delta T = K_f \cdot m$$

$$K_f = 1.86^\circ \text{C/m}$$

$$m = \frac{\Delta T}{K_f} = \frac{16.0}{1.86} = \underline{\underline{8.6 \text{ m}}}$$

b)

$$\% \text{ PESO} = \frac{\text{MASSA ETANOLLO}}{\text{MASSA ETANOLLO} + \text{MASSA H}_2\text{O}}$$

CONSIDERANDO 1 kg DI H₂O AURO'

8.6 moli ETANOLO DATA LA CONCENTRAZ.

$$\text{MASSA ETANOLO} = n_{\text{ETANOLO}} \cdot PM_{\text{ETANOLO}}$$

$$= 8.6 \cdot 46.07$$

$$= 396.2 \text{ gr}$$

$$\% \text{ PESO ETANOLO} = \frac{396.2}{396.2 + 1000} \cdot 100$$

↓
H₂O

$$= \underline{\underline{28.4 \%}}$$

⑤

0.255 g di X ($X = C_{10}H_8Fe$ FORM. MINIMA)

$$\text{MASSA BENZENE} = 11.0 \text{ gr}$$

$$\Delta T_{EB} = 80.26 - 80.10 = 0.16 ^\circ C$$

$$\Delta T_{EB} = K \cdot m \quad K = 2.53 ^\circ C/m$$

DA CUI POSSO DETERMINARE MOLALITA' X

$$m_X = \Delta T_{EB} / K = 6.32 \cdot 10^{-2} m$$

$$m_x = m_x / \text{MASSA BENZENE}$$

$$m_x = \text{MASSA BENZENE} \cdot m_x$$

$$\hat{=} 11.12 \cdot 10^{-3} \cdot 6.32 \cdot 10^{-2} = 7.03 \cdot 10^{-4} \text{ mol}$$

QUINDI $7.03 \cdot 10^{-4}$ mol di X PESANO 0.255 g

$$m_x = \frac{\text{MASSA } x}{PM_x}$$

$$PM_x = \frac{\text{MASSA } x}{m_x} = \frac{0.255}{7.03 \cdot 10^{-4}} = 362.73 \text{ g/mol}$$

$$PM_{\text{FORMULA MINIMA}} = 10 \cdot 12.01 + 8 \cdot 1.01 + 55.85 \\ = 184.03 \text{ g/mol.}$$

$$\frac{PM_x}{PM_{\text{FORMULA MINIMA}}} = \frac{362.73}{184.03} \approx 2$$

DEVO MOLTIPLICARE LA FORMULA MINIMA PER 2 $\Rightarrow C_{20}H_{16}Fe_2$

⑥

$$\text{MASSA (LiF)} = 52.5 \text{ gr}$$

$$\text{MASSA (H}_2\text{O)} = 306 \text{ gr}$$

$$\text{PM}_{\text{LiF}} = 25.94 \text{ gr/mol}$$

$$n_{\text{LiF}} = \frac{52.5}{25.94} = 2.02 \text{ mol}$$

$$m_{\text{LiF}} = \frac{n_{\text{LiF}}}{\text{MASSA (H}_2\text{O)}} = \frac{2.02}{306 \cdot 10^{-3}} = 6.61 \text{ m}$$

$$\Delta T = i \cdot K \cdot m_{\text{LiF}}$$

LiF ELETTROLITA QUINDI $i = 2$



$$\Delta T = 2 \cdot 1.86 \cdot 6.61 = \underline{\underline{24.59^\circ \text{C}}}$$

⑦

$$\hat{P} = 3.1 \text{ mm Hg}$$

$$T = 25^\circ \text{C}$$

$$= 298^\circ \text{K}$$

$$[\text{INS.}] = 1 \text{ gr/L}$$

$\hat{\Pi} = M \cdot R \cdot T$ NON ELETTROLITA E
QUINDI $\lambda = 1$

$$M = \hat{\Pi} / R \cdot T$$

$$\hat{\Pi} = 3.1 / 760 = 4.08 \cdot 10^{-3} \text{ atm}$$

QUINDI :

$$M = \frac{4.08 \cdot 10^{-3}}{R \cdot 298} = 1.67 \cdot 10^{-4} \text{ mol/L}$$

QUINDI IN 1 L DI SOLUZIONE
CONTIENE $1.67 \cdot 10^{-4}$ moli E

1.00 gr DI INSULINA :

$$PM_{\text{ins.}} = \frac{1.00}{1.67 \cdot 10^{-4}} = 5.99 \cdot 10^3 \frac{\text{gr}}{\text{mol}} \\ \approx 6000 \text{ gr/mol}$$

⑧ $P_{\text{SOLUZIONE}} = P_{\text{SOLVENTE}} \cdot X_{\text{SOLVENTE}}$

a) 0.35 mol in 1 kg H_2O

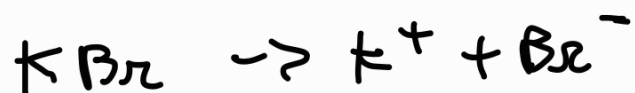
$$M_{H_2O} = \frac{1000}{18} = 55.5 \text{ mol}$$

$$X_{H_2O} = \frac{55.5}{55.5 + 0.35} = \underline{0.994}$$

b) NON ELETTROLITA QUINDI $i = 1$

$$X_{H_2O} = \frac{55.5}{55.5 + 0.5} = \underline{0.991}$$

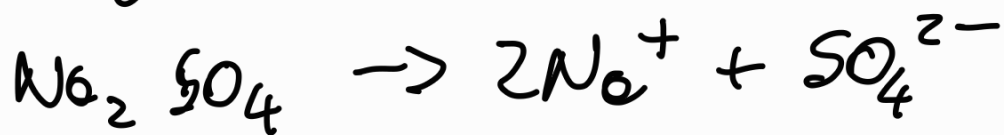
c) KBr ELETTROLITA QUINDI $i = 2$



QUINDI MOL SOLUTO $0.70 \cdot 2 = 0.40$

$$X_{H_2O} = \frac{55.5}{0.40 + 55.5} = \underline{0.993}$$

d) Na_2SO_4 ELETTROLITA



QUINDI AURO' $3 \cdot 0.20 = 0.60$ mol
DI SOLUTO

$$X_{H_2O} = \frac{55.5}{55.5 + 0.60} = \underline{0.989}$$

i) $\text{Na}_2\text{SO}_4 < \text{ZUCCHERO} < \text{KBr} < \text{GLICOLE ET.}$

$$d < b < c < a$$

ii)

$$\Delta T = (i \cdot m) \cdot K$$

e) $i \cdot m = 0.35 \text{ m}$

b) $i \cdot m = 0.50 \text{ m}$

c) $i \cdot m = 0.40 \text{ m}$

d) $i \cdot m = 0.60 \text{ m}$

IN PRATICA COME OVUNQUE ORDINAMENTO
CONTRARIO:

$$e < c < b < d$$

P. VAPORE PIÙ BASSA COMPORTA
UN PIÙ ALTO PUNTO DI EBOLLIZIONE.