

TEQB – Resolução Exame

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Conteúdo

Questão 1	2	Questão 4	12
Questão 2	6	Questão 5	15
Questão 3	8			

Questão 1

1 mol gas perfeito num processo reversível

1. 2.1 bar, 19 °C 2. 1.2 bar 3. 0.86 bar 4. .

- $\Delta H_{1 \rightarrow 2} = 844$

- $W_{3 \rightarrow 4} = -1671$

- 2→3: Adiabático $P V^\gamma = Cte$

- 3→4: T constante

Q1 a.

T em 2

$$T_2 = T_1 + \frac{\Delta H_{1 \rightarrow 2}}{n C_P} = T_1 + \frac{\Delta H_{1 \rightarrow 2}}{n (C_V + R)} = (19 + 273.15) + \frac{844}{1 * 3.5 * 8.31} \cong 321.15$$

Q1 b.

T em 3

$$\begin{aligned} P_3 V_3^\gamma &= P_3 \left(\frac{n R T_3}{P_3} \right)^{1.4} = \frac{n^{1.4} R^{1.4} T_3^{1.4}}{P_3^{0.4}} = P_2 V_2^\gamma = \frac{n^{1.4} R^{1.4} T_2^{1.4}}{P_2^{0.4}} \Rightarrow \\ \Rightarrow T_3 &= \left(\frac{n^{1.4} R^{1.4} T_2^{1.4} P_3^{0.4}}{P_2^{0.4} n^{1.4} R^{1.4}} \right)^{1/1.4} = T_2 \left(\frac{P_3}{P_2} \right)^{0.4/1.4} \cong 321.15 \left(\frac{0.86}{1.2} \right)^{0.4/1.4} \cong 291.99 \end{aligned}$$

Q1 c.

P em 4

$$\begin{aligned} - \int n R T \, dV/V &= -n R T_3 \ln(V_4/V_3) = -n R T_3 \ln \left(\frac{n R T_4}{P_4} - \frac{n R T_3}{P_3} \right) = W_{3 \rightarrow 4} \implies \\ \implies P_4 &= \left(\left(\exp \left(-\frac{W_{3 \rightarrow 4}}{n R T_3} \right) + \frac{n R T_3}{P_3} \right) \frac{1}{n R T_4} \right)^{-1} = \\ &= \left(\exp \left(-\frac{-1671}{1 * 8.31 * 291.99} \right) + \frac{1 * 8.31 * 291.99}{0.86 * 10^5} \right)^{-1} 1 * 8.31 * 291.99 \cong 1202.74 \end{aligned}$$

Questão 2

Sistema n-octano+entanol

Q2 a.

Calcole $\gamma_{etanol,I}$ quando $x_{n-octano} = 0.65$ a 75°C

$\gamma_{etanol,I}$

Questão 3

um pouco acima do ponto triplo ($175.5 \text{ K}, 1.86 * 10^{-6} \text{ bar}$)

$$P_{fus} = -1077.3 + 6.1309 T$$
$$\gtrsim (175.5 \text{ K}, 1.86 * 10^{-6} \text{ bar}) \wedge P < 500 \text{ bar}$$

- $V_{m,L} = 35.43 \text{ cm}^3 \text{ mol}^{-1}$
- $V_{m,S} = 32.47 \text{ cm}^3 \text{ mol}^{-1}$
- $P_{vap,L,215 \text{ K}} = 3.44 * 10^{-4} \text{ bar}$
- $C_{P,s} = 52.0 \text{ J K}^{-1} \text{ mol}^{-1}$
- $C_{P,l} = 70.8 \text{ J K}^{-1} \text{ mol}^{-1}$
- $C_{P,g} = 39.0 \text{ J K}^{-1} \text{ mol}^{-1}$

Q3 a.

Calcule P de sub a 173 K

$$P_{sub} = (-1077.3 + 6.1309 * T_t) \exp \left(\frac{-\Delta H_{sub}}{R} (T_1^{-1} - T_0^{-1}) \right) =$$

$$= (-1077.3 + 6.1309 * T_t) \exp \left(-\frac{T_1^{-1} - T_t^{-1}}{R} (\Delta H_{fus} + \Delta H_{vap}) \right)$$

$$\Delta H_{fus} = T_t \Delta V_{fus} \left(\frac{\partial P}{\partial T} \right)_{fus} = T_t (V_1 - V_0) \left(\frac{\partial P}{\partial T} \right)_{fus} =$$

$$= T_t (V_1 - V_0) \frac{\partial}{\partial T} (-1077.3 + 6.1309 T_t) * 10^6 =$$

$$= T_t (V_1 - V_0) * 6.1309 * 10^6 =$$

$$= 175.5 * (35.43 - 32.47) * 10^{-6} * 6.1309 * 10^6 \cong 3184.88$$

$$\ln P_{vap,L} = a + b/T \implies \ln 3.44 * 10 - b/215 = \ln 1.86 * 10^{-1} - b/175.5 \implies$$

$$\implies b = \frac{\ln (34.4/0.186)}{175.5^{-1} - 215^{-1}} \cong 4986.48 \implies$$

$$\implies a \cong 3.44 * 10 - 4986.48/215 \cong 11.21 \implies$$

$$\implies \ln P_{vap} \cong 11.21 + 4986.48/T$$

$$\Delta H_{vap} = R T^2 \frac{d \ln P}{dT} \cong R T^2 (4986.48/T_2) \cong 41\,459.93$$

$$P_{sub} \cong$$

$$\cong (-1077.3 + 6.1309 * 175.5) *$$

$$* \exp \left(-\frac{173^{-1} - 175.5^{-1}}{8.31} (3184.88 + 41\,459.93) \right) \cong$$

$$\cong 1383.83$$

Nota : Sobre a equação do P vap encontrada, esta não está correta, ao fazer a aproximação usando uma curva log em 1/T gerou valores em P muito superiores que o esperando, talvez seja melhor parametrizar em alguma outra curva como uma reta.

Q3 b.

$$\Delta S(180 \text{ K}, 200 \text{ bar}) \rightarrow (300 \text{ K}, 0.05 \text{ bar})$$

$$\left\{ \begin{array}{l} P_{fus,180 \text{ K}} = (-1077.3 + 6.1309 * 180) * 10^6 \cong 262.62 * 10^5 > 200 * 10^5 \\ P_{vap,180 \text{ K}} = (-11.21 + 4986.48/180) * 10^5 \cong 16.50 * 10^5 > 0.05 * 10^5 \\ T_{fus,200 \text{ bar}} = 200 + 1077.3/6.1309 \cong 375.72 > 300 \\ T_{vap,200 \text{ bar}} = \frac{4986.48}{\ln(200 * 10^5) - 11.21} \cong 890.25 \end{array} \right\}$$

$$\Delta S =$$

$$= \left(\begin{array}{l} \Delta S(l, 180 \text{ K}, (200 \rightarrow 16.50) \text{ bar}) \\ + \Delta S((l \rightarrow g), 180 \text{ K}, 16.50 \text{ bar}) \\ + \Delta S(g, (180 \rightarrow 300) \text{ K}, (16.50 \rightarrow 0.05) \text{ bar}) \end{array} \right) =$$

$$= \left(\begin{array}{l} 0 \\ + - n \Delta H_{vap}/180 \\ + \int n C_{P,g} dT/T - n R \ln(P_1/P_0) \end{array} \right) =$$

$$= \left(\begin{array}{l} -41459.93/180 \\ +39 * \ln 300/175.5 - 8.31 * \ln(0.05/16.5) \end{array} \right) \cong$$

$$\cong -209.45$$

Q3 c.

$$\Delta G(300 \text{ K}, 10 \text{ bar}) \rightarrow (300 \text{ K}, 0.05 \text{ bar})$$

$$\{ P_{fus,300 \text{ K}} = (1077.3 + 5.1309 * 300) * 10^6 \cong 2616.57 * 10^6 > 10 * 10^5 \}$$

$$\Delta G =$$

$$= \left(\begin{array}{l} \Delta G(s, (10 \rightarrow P_{fus}) \text{ bar}) \\ + \Delta G((s \rightarrow l), P_{fus} \text{ bar}) \\ + \Delta G(l, (P_{fus} \rightarrow P_{vap}) \text{ bar}) \end{array} \right) =$$

$$= \left(\begin{array}{l} \int_{P_1}^{P_2} V \, dP \\ + 0 \\ + \int_{P_0}^{P_1} V \, dP \end{array} \right) =$$

$$= \left(\begin{array}{l} V \int_{P_1}^{P_2} dp \quad (\text{vol liq constante em } \Delta P) \\ + \int_{P_0}^{P_1} \frac{n R T}{P} \, dP \end{array} \right) =$$

$$= \left(\begin{array}{l} V_l (P_2 - P_1) \\ + \frac{n R T}{\ln} (P_3/P_2) \end{array} \right)$$

Nota: Continuo a sofrer com a má parametrização da curva de P de vaporização

Questão 4

- $P_{vap,sol,30^{\circ}\text{C}} = 0.1026 \text{ bar}$
- $\ln P_{vap,eta} = 14.502 - 5084.7/T$
- $M_{eta} = 46.07 \text{ g mol}^{-1}$
- $M_{naf} = 128.17 \text{ g mol}^{-1}$
- $\rho_{sol} \approx \rho_{eta} = 0.789 \text{ g cm}^{-1}$

(i)

Pressão osmótica da sol a 20 C

$$\Pi = \frac{[i] P_{eta}}{n_{eta}}$$

(ii)

T de vap sol a 0.951 bar

$$\begin{aligned}
 t_{vap,sol} &= \left(t_{vap,eta}^{-1} - \ln x_{eta} \frac{R}{\Delta H_{vap,eta}} \right)^{-1} = \\
 &= \left(\left(\frac{5084.7}{14.502 - \ln P} \right)^{-1} - \ln(x_{eta}) \frac{R}{\Delta H_{vap,eta}} \right)^{-1} = \\
 &= \left(\frac{14.502 - \ln P}{5084.7} - \frac{\ln(x_{eta}) R}{\Delta H_{vap,eta}} \right)^{-1} ;
 \end{aligned}$$

$$\begin{aligned}
 \left. \frac{dP}{dT} \right|_{vap} &= \frac{\Delta H_{vap}}{T \Delta V_{vap}} = \frac{\Delta H_{vap}}{T (V_{vap,g} - V_{vap,l})} \cong \frac{\Delta H_{vap}}{T V_{vap,g}} \cong \frac{\Delta H_{vap}}{T (RT/P)} = \frac{P \Delta H_{vap}}{RT^2} \Rightarrow \\
 \Rightarrow \frac{dP/P}{dT} &= \frac{d \ln P}{dT} = \frac{d}{dT} (14.502 - 5084.7 T^{-1}) = 5084.7 T^{-2} = \frac{\Delta H_{vap}}{RT^2} \Rightarrow \\
 \Rightarrow \Delta H_{vap} &= 5084.7 R t_{vap,sol} = \left(\frac{14.502 - \ln P}{5084.7} - \frac{\ln(x_{eta}) R}{5084.7 R} \right)^{-1} = \\
 &= \frac{5084.7}{14.502 - \ln(0.951 * x_{eta})} \cong
 \end{aligned}$$

Questão 5

- $T = 30\text{ }^{\circ}\text{C}$
- $n_{oct,tot} = 3\text{ mol}$
- $n_{agua,tot} = 2\text{ mol}$
- $x_{oct,1} = 0.744$
- $x_{oct,2} = 0.002$

Q5 a.

Razão entre as quantidades (mol) da fase mais rica em n-octano e da fase mais rica em água

(i)

Fase mais rica em n-Octano

