

Worksheet #9 (Total number of points you can get is 3 pts)

1. For a certain chemical reaction one finds a enthalpy difference $\Delta H = -150 \text{ kJ/mol}$ and an entropy difference $\Delta S = -300 \text{ J/(K mol)}$. Assume that these two values do not depend on temperature. At what temperature will be the Gibb's energy difference $\Delta G = 0$?

$$\Delta G = \Delta H - T \Delta S = 0$$

$$\Rightarrow T \Delta S = \Delta H$$

$$\Rightarrow T = \frac{\Delta H}{\Delta S} = \frac{-150 \cdot 10^3 \text{ J/mol}}{-300 \text{ J/(K mol)}} = 500 \text{ K}$$

2. The equilibrium constant for the reaction



in benzene solution at 280 K is $K_c = 0.0022 \text{ mol/dm}^3$. Calculate ΔG° .

$$\Delta G^\circ = -RT \ln K_c^\circ$$

$$= -8.3145 \cdot 280 \cdot \ln(2.2 \cdot 10^{-3}) \text{ J/mol}$$

$$= -8.3145 \cdot 280 \cdot (-6.1193) \text{ J/mol}$$

$$= 14.25 \text{ kJ/mol}$$

The Gibbs energies of formation of A(g) and B(g) are 50 kJ/mol and

25.5 kJ/mol, respectively. The standard state is 1 bar and 300 K. Assume ideal behavior and a reaction $A \rightleftharpoons 2B$. Calculate the pressure where 50% of A is dissociated.

$$\Delta G^\circ = 51.1 \text{ kJ} - 50 \text{ kJ} = 1.1 \text{ kJ}$$

$$= -RT \ln K_p^\circ \Rightarrow \ln K_p^\circ = \frac{-\Delta G^\circ}{RT} = \frac{-1000 \text{ J/mol}}{8.3145 \text{ J/(K mol)} \cdot 300 \text{ K}} = 0.4009$$

$$\Rightarrow K_p^\circ = 0.67$$

$$\begin{array}{l} A \rightleftharpoons 2B \\ 1-x \quad 2x \quad \text{total } 1+x \\ P_A = \frac{1-x}{1+x} P \quad P_B = \frac{2x}{1+x} P \\ K_p^\circ = \frac{[2x/(1+x)]^2}{[1-x/(1+x)]} = \frac{4x^2}{1-x^2} \end{array}$$

$$P = \frac{1-x^2}{4x^2} K_p^\circ \text{ bar}$$

$$x = 0.5$$

$$\Rightarrow P = \frac{0.75}{1} \cdot 0.67 \text{ bar}$$

$$= 0.5025 \text{ bar}$$