

## CH 223 – Worksheet 6

1. The normal boiling point of ethanol ( $C_2H_5OH$ ) is  $78.3^\circ C$ , and its molar enthalpy of vaporization is  $38.56 \text{ kJ/mol}$ . What is the change in entropy in the system when  $68.3 \text{ g}$  of  $C_2H_5OH$  (g) at  $1 \text{ atm}$  condenses to a liquid at the normal boiling point?

$$68.3 \text{ g } C_2H_5OH \left( \frac{1 \text{ mol}}{46 \text{ g}} \right) = 1.485 \text{ mol } C_2H_5OH$$

$$\Delta S = -\frac{\Delta H}{T} = \frac{(1.485 \text{ mol } C_2H_5OH)(38.56 \frac{\text{kJ}}{\text{mol}})}{351 \text{ K}} \\ = 0.163 \frac{\text{kJ}}{\text{K}} = 163 \frac{\text{J}}{\text{K}}$$

2. The normal freezing point of 1-propanol ( $C_3H_8O$ ) is  $-127^\circ C$ . (a) Is the freezing an endothermic or exothermic process? (b) In what temperature range is the freezing of 1-propanol a spontaneous process? (c) In what temperature range is it a nonspontaneous process? (d) Is there any temperature at which liquid and solid 1-propanol are in equilibrium? Explain.

a) exothermic

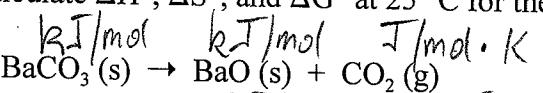
d) equilibrium occurs at

b)  $< -127^\circ C$

$-127^\circ C$  and 1 atm.

c)  $> -127^\circ C$

3. Using data from Appendix C, calculate  $\Delta H^\circ$ ,  $\Delta S^\circ$ , and  $\Delta G^\circ$  at  $25^\circ C$  for the following reaction:



$$-1216.3 \quad -553.5 \quad -393.5 \quad \Delta H$$

$$-1137.6 \quad -525.1 \quad -394.4 \quad \Delta G$$

$$112.1 \quad 70.42 \quad 213.6 \quad \Delta S$$

$$\Delta H = \sum n \Delta H^\circ(\text{products}) - \sum m \Delta H^\circ(\text{reactants})$$

$$\Delta H = \left[ -553.5 \frac{\text{kJ}}{\text{mol}} + -393.5 \frac{\text{kJ}}{\text{mol}} \right] - \left[ -1216.3 \frac{\text{kJ}}{\text{mol}} \right] = 269.3 \text{ kJ}$$

$$\Delta S = \sum n S^\circ(\text{products}) - \sum m S^\circ(\text{reactants})$$

$$\Delta S = \left[ 70.42 \frac{\text{J}}{\text{mol}\cdot\text{K}} + 213.6 \frac{\text{J}}{\text{mol}\cdot\text{K}} \right] - \left[ 112.1 \frac{\text{J}}{\text{mol}\cdot\text{K}} \right] = 171.32 \text{ J}$$

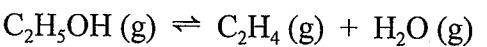
$$\Delta G_2 = \Delta H - T \Delta S = 269.3 \text{ kJ} - 298 \text{ K} (0.17132 \frac{\text{kJ}}{\text{mol}}) = 218.2 \text{ kJ}$$

$$\text{or } \Delta G_2 = \sum n \Delta G^\circ(\text{products}) - \sum m \Delta G^\circ(\text{reactants})$$

$$\Delta G^\circ = \left[ -525.1 \frac{\text{kJ}}{\text{mol}} + -394.4 \frac{\text{kJ}}{\text{mol}} \right] - \left[ -1137.6 \frac{\text{kJ}}{\text{mol}} \right] = 218.2 \text{ kJ}$$

They are the same.

4. Use data from Appendix C to calculate the equilibrium constant, K, at 298 K for the following reaction:



$$-168.5 \frac{\text{kJ}}{\text{mol}} \quad 68.11 \frac{\text{kJ}}{\text{mol}} \quad \downarrow -228.57 \frac{\text{kJ}}{\text{mol}}$$

$$\Delta G = \Delta G^\circ + RT \ln Q$$

$$\Delta G^\circ = \sum n \Delta G^\circ (\text{products}) - \sum m \Delta G^\circ (\text{reactants})$$

$$= [68.11 \frac{\text{kJ}}{\text{mol}} + -228.57 \frac{\text{kJ}}{\text{mol}}] - [-168.5 \frac{\text{kJ}}{\text{mol}}]$$

$$= 8.04 \frac{\text{kJ}}{\text{mol}}$$

at equilibrium

$$\Delta G = 0 = \Delta G^\circ + RT \ln K \text{ at eq. } K = Q$$

$$\therefore \Delta G^\circ = -RT \ln K \quad R = 8.314 \frac{\text{J}}{\text{mol} \cdot \text{K}}$$

$$-\frac{\Delta G^\circ}{RT} = \ln K$$

$$K = e^{-\Delta G^\circ/RT} = e^{-8.040 \frac{\text{J}}{\text{mol}} / 8.314 \frac{\text{J}}{\text{mol} \cdot \text{K}} \cdot 298 \text{ K}}$$

$$= 0.0390$$