

Chapter 18 -- Review Problem

Consider the gas-phase reaction: $\text{N}_2\text{O}_5 + \text{H}_2\text{O} \longrightarrow 2 \text{HNO}_3$
and the following thermodynamic data.

Compd	ΔH°_f (kJ/mole)	S° (J/mole-K)
$\text{N}_2\text{O}_5(\text{g})$	11.0	356
$\text{H}_2\text{O}(\text{g})$	- 242	189
$\text{HNO}_3(\text{g})$	- 174	156

- (a) Decide whether or not the above reaction is spontaneous at 25°C by calculating the value of the *appropriate* thermodynamic quantity.
- (b) Calculate the *temperature* (in $^\circ\text{C}$) at which the above reaction should have an equilibrium constant (K_p) equal to 1.00.

(a) $\Delta H^\circ = - 117 \text{ kJ}$

$$\Delta S^\circ = - 233 \text{ J/K} = - 0.233 \text{ kJ/K}$$

$$\Delta G^\circ = \Delta H^\circ - T\Delta S^\circ = - 117 \text{ kJ} - (298 \text{ K})(- 0.233 \text{ kJ/K}) = - 47.6 \text{ kJ}$$

The negative ΔG° value confirms that the reaction is spontaneous (at least under standard conditions).

- (b) The temp where $\ln K = 1.00$ is the temp at which $\Delta G = 0$.

$$\Delta G = - RT \ln K = - RT \ln(1.00) = 0$$

Because ΔH and ΔS are relatively independent of temperature, we can estimate this temp by setting $\Delta G = 0$ as follows.

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

$$\therefore T = \Delta H / \Delta S = (- 117 \text{ kJ}) / (- 0.233 \text{ kJ/K}) = 502 \text{ K}$$

$$T = 502 - 273 = 229 \text{ }^\circ\text{C}$$