You must show your work for full credit.

Exp 6 and 8 Questions

- 1. A 25.0 mL aliquot of a saturated CaSO₄ solution (no solid present) requires 32.1 mL of 0.0200 M EDTA to reach an end point.
- a) What is the Ca²⁺ concentration (in M) of the solution? (2 pts) $n(Ca^{2+}) = n(EDTA) = (0.0321 L)(0.0200 mol L^{-1}) = 6.42 \times 10^{-4} mol = [Ca^{2+}](0.025 L)$ $[Ca^{2+}] = 2.57 \times 10^{-2} M$
- b) Ignoring activities, what is the K_{sp} for CaSO₄ based on the titration data? (1 pt) $K_{sp} = [Ca^{2+}][SO_4^{2-}]$ and $[Ca^{2+}] = [SO_4^{2-}] = 2.57 \times 10^{-2}$ $K_{sp} = (2.57 \times 10^{-2})^2 = 6.59 \times 10^{-4}$
- **2.** HCl is added to a mixture containing Ca²⁺, Na⁺, Ag⁺, and K⁺.
- a) Which ion will precipitate out? (1 pt) Ag+
- b) What can be added to the precipitant to make it redissolve (once it is removed from the HCl and washed)? (1 pt). NH₃ (or NH₄OH)

- Thermodynamics problems
- <u>3.</u> Which of the following reactions leads to a decrease in entropy for the system? (1 pt)

a) $I_2(s) \leftrightarrow I_2(g)$ $d)2I(g) \leftrightarrow I_2(g)$

- b) $I_2(s) \leftrightarrow I_2(aq)$ c) $I_2(s) + H_2(g) \leftrightarrow 2HI(g)$
- **4.** Under what temperature regimes will the reaction: $N_2(g) + 3H_2(g) \rightleftharpoons 2NH_3(g)$ ($\Delta H^{\circ} = -91.8$
- k[/mol) be spontaneous? (1 pt) $\Delta S < 0$ so $\Delta G = \Delta H T \Delta S < at low T$
- b) high temperature c) low temperature d) always. a) never
- **<u>5.</u>** When any spontaneous reaction occurs, $\Delta S_{universe}$ is **<u>positive</u>** (give sign) (1 pt)

Chemistry 1B, Fall 2015 Quiz #7A KEY (one more on back)

6. Given that $\Delta G_f^{\circ}(HI) = 1.7 \text{ kJ/mol}$ (subscript f means formation), calculate K (the equilibrium constant) for $I_2(s) + H_2(g) \leftrightarrow 2HI(g)$ at 298 K. $R = 8.314 \text{ J mol}^{-1} \text{ K}^{-1}$ (2 pt)

 $\Delta G_{rxn}^{\circ} = 2\Delta G_f^{\circ}(HI) - \Delta G_f^{\circ}(H_2(g)) - \Delta G_f^{\circ}(I_2(s)) = 2(1.7kJ/mol)$ (others are elements in standard states so ΔG_f° are 0).

 $\Delta G_{rxn}^{\circ} = 3.4 \text{ kJ/mol} = -RTlnK \text{ or } K = exp(-\Delta G_{rxn}^{\circ}/RT) = e^{-3400J/(8.314 \text{ J mol}-1 \text{ K}-1)(298 \text{ K})} = e^{-1.37} = 0.25$