## CHEM 301 Assignment #4 (Optional)

Provide solutions to the following questions in a <u>neat and well organized</u> manner.

Clearly state assumptions and reference sources for any constants used.

(odd numbered questions only) December 7<sup>th</sup>

- 1. The Maximum Acceptable Concentration of mercury in water is 0.001 ppm. Is the MAC likely to be exceeded by the dissolution of a)  $\mathbf{HgS}(s)$  ( $K_{sp} = 1 \times 10^{-56}$ ) and b)  $\mathbf{HgO}(s)$  (solubility reported as 5.3 mg per 100 mL). Are either of your answers affected by the pH or pe of the water sample? Explain.
- **2.** An effluent waste stream contains 330 mg  $L^{-1}$  organic matter (suspended and dissolved) and 27 mg  $L^{-1}$  ammonium ion (as **N**). Write chemical equations for the complete oxidation of  $\{CH_2O\}$  and  $NH_4^+$  and calculate the total BOD load of this effluent in mg/L  $O_2$ .
- 3. A sewage treatment plant is designed to process  $3.0 \times 10^{12}$  L of sewage daily.
- a) What capacity (m<sup>3</sup>) is required for the primary settling lagoon if the residence time is to be 6 hours?
- b) If the influent water has a BOD of 850 ppm, what volume of oxygen at 15°C is required per day to reduce the BOD by 90%.
- **4.** Calculate the mass of lime ( $Ca(OH)_2(s)$ ), required to chemically treat 27,000 m<sup>3</sup> of waste water containing 6.1 mg L<sup>-1</sup> of phosphorous. Assume the following reaction goes to completion when a two fold molar excess of  $Ca(OH)_2$  is used.

$$5 \text{ Ca(OH)}_2 + 3 \text{ HPO}_4^{2-} \rightarrow \text{Ca}_5(\text{PO}_4)_3\text{OH} + 6 \text{ OH}^- + 3 \text{ H}_2\text{O}$$

**5.** A sewage sample contains 8.8 ppm of dissolved phosphorous in the form of ortho phosphate. It is brought to pH of 9.0 and  $[\mathbf{Ca}^{2^+}] = 4.7$  mM by the addition of  $\mathbf{Ca}(\mathbf{OH})_2$ . What is the concentration of dissolved phosphorous (as ppm **P**) when it's in equilibrium with precipitated calcium phosphate  $(\mathbf{K}_{sp} \ \mathbf{Ca_3(PO_4)_2} = 1 \times 10^{-24})$ ?

- **6.** Iron can occur as a carbonate mineral  $\mathbf{FeCO_3}$ , which has a  $K_{sp} = 3.1 \times 10^{-11}$ . Calculate the concentration of of iron (ppm) in groundwater in equilibrium  $\mathbf{FeCO_3}$  and  $\mathbf{CaCO_3}$ . The concentration of  $\mathbf{Ca^{2+}}(aq) = 120$  ppm. Hint: use the  $K_{sp}$  of  $\mathbf{CaCO_3}$  of 6.0 x 10<sup>-9</sup> to estimate the  $[\mathbf{CO_3}^{2-}(aq)]$ .
- 7. Using the solubility of  $\mathbf{FePO_4}(s)$  ( $K_{sp} = 1.3 \times 10^{-22}$ ) and the acidity constants for  $\mathbf{H_3PO_4}$ , determine if a concentration of  $\mathbf{Fe^{3+}}$  of 100. ppm would be sufficient to precipitate 90% of the phosphate from a solution initially containing  $1.0 \times 10^{-4}$  mol  $L^{-1}$  of total phosphate at a constant pH of 8.00.
- **8.** Gold ores are frequently leached with cyanide dissolving the gold according to;  $\mathbf{Au}(s) + 2 \mathbf{CN}^{-}(aq) + {}^{1}\!/_4 \mathbf{O}_2(g) + {}^{1}\!/_2 \mathbf{H}_2 \mathbf{O}(l) ===== \mathbf{Au}(\mathbf{CN})_2^{-}(aq) + \mathbf{OH}^{-}(aq)$  Which has an equilibrium constant,  $K = 1.2 \times 10^{17}$ . In order to prevent undue environmental contamination by cyanide, you wish to operate this process under conditions such that at least 98% of the  $\mathbf{CN}^{-}$  is converted to  $\mathbf{Au}(\mathbf{CN})_2^{-}$ . Your process operates at a pH 9.0 and the  $\mathbf{O}_2$  pressure inside the ore body is constant at 0.032 atm. Calculate the molar concentration of  $\mathbf{CN}^{-}$  required.
- 9. Filter alum  $Al_2(SO_4)_3$  is often used to remove phosphate from wastewater. A wastewater of pH 5.62 containing 25 ppm total phosphate is treated with alum until the equilibrium concentration of  $Al^{3+}$  is 4.0 x  $10^{-9}$  mol  $L^{-1}$ . What fraction of the phosphate is precipitated as  $AlPO_4(s)$ ? Consider only the equilibria below;

AlPO<sub>4</sub>(s) ==== Al<sup>3+</sup>(aq) + PO<sub>4</sub><sup>3-</sup>(aq) 
$$K_{sp} = 1.0 \times 10^{-21}$$
  
H<sub>2</sub>PO<sub>4</sub><sup>-</sup>(aq) ==== HPO<sub>4</sub><sup>2-</sup>(aq) + H<sup>+</sup>(aq)  $K_a = 6.2 \times 10^{-8}$   
HPO<sub>4</sub><sup>2-</sup>(aq) ==== PO<sub>4</sub><sup>3-</sup>(aq) + H<sup>+</sup>(aq)  $K_a = 4.8 \times 10^{-13}$ 

**10.** Lead solder on copper plumbing has been suggested as a possible source of low levels of lead (II) ions in drinking water. Using the standard reduction tables and the Nernst equation, comment on the possibility of  $\mathbf{Pb}^{2+}$  being present at or above the MAC of 10. ppb, if the water in contact with  $\mathbf{Pb}(s)$  contains  $\mathbf{Cu}^{2+}$  at 0.10 ppm (*ignore Cu*<sup>+</sup> *ions*).