ACTIVITY/WORKSHOP on IONIC EQUILIBRIUM Name:		
1. Predict the product(s) for the following reaction, and designate the conjugate acid and conjugate base. $HCHO_2(aq) \ + \ PO_4^{-3}(aq) \ \leftrightarrow ?$		
2. If $pH = 6.6$ , what are the molar concentrations of $H_3O^+(aq)$ and $OH^-(aq)$ in the solution?		
3. Lactic acid, $HC_3H_5O_3$ , has one acidic hydrogen. A 0.10 M solution of lactic acid has a pH of 2.44. Calculate its $K_a$ .		
4. A 0.200 M solution of bromoacetic acid, BrCH <sub>2</sub> COOH, is 13.2 percent ionized. Using this information, calculate [BrCH <sub>2</sub> COO <sup>-</sup> ], [H <sup>+</sup> ], [BrCH <sub>2</sub> COOH], AND K <sub>a</sub> for bromoacetic acid.		
5. Calculate the pH AND percent ionization of a $0.050$ M butanoic acid solution, $HC_4H_7O_2$ , $K_a = 1.5 \times 10^{-5}$		
6. Calculate the pH of a 5.0 M $H_3PO_4$ solution and the equilibrium concentrations of the species $H_3PO_4$ , $H_2PO_4$ , $HPO_4$ , and $PO_4$ . ( $K_{a1} = 7.5 \times 10^{-3}$ ; $K_{a2} = 6.2 \times 10^{-8}$ ; $K_{a3} = 4.8 \times 10^{-13}$ ).		
		easing acid strength, and explain your ordering:
В. НО	OCl, HOI, HOBr	
C. Se	H <sub>2</sub> , GeH <sub>4</sub> , AsH <sub>3</sub>	
D. CH	H <sub>3</sub> CO <sub>2</sub> H, FCH <sub>2</sub> CO <sub>2</sub> H, F <sub>2</sub> CHO	CO <sub>2</sub> H
8. A solution contains 45 mL of 0.100 M CH <sub>3</sub> COOH(aq), K <sub>a</sub> = 1.8 x 10 <sup>-5</sup> , and 55 mL of 0.100 M NaCH <sub>3</sub> CO <sub>2</sub> (aq).  A. Determine the pH of this solution.		
		ion when 10.0 mL of 0.350 M NaOH(aq) is added.
9. A 25.0 mL san of benzoic acid is A. Wh B. Wh C. Wh D. Wh E. Wh F. Wh Me	mple of 0.035 M benzoic acides 6.3 x 10 <sup>-5</sup> .  That is the pH, pOH, and [H <sub>3</sub> O <sup>+</sup> ] of the solution when it that is the pH of the solution at the line is the pH of the solution at the line is the pH of the solution when it that is the pH of the solution when is the pH of the solution at the pH of the solution at the pH of the solution at the pH of the solution when is the pH of the solution at the pH of the solution at the pH of the solution when is the pH of the solution at the pH of the solution when is the pH of the solution at the pH of the solution when is the pH of the solution	d, HC <sub>7</sub> H <sub>5</sub> O <sub>2</sub> , is titrated with 0.099 M NaOH solution. The $K_a$ the benzoic acid solution prior to the titration? 15.0 mL of 0.099 M NaOH solution is added to the original sample? half-way point of the titration? equivalence point of the titration? 35.0 mL of 0.099 M NaOH solution is added to the original sample? best indicator to use for this titration? Justify your answer. $K_a = 1 \times 10^{-5}$ $K_a = 1 \times 10^{-8}$
T NI	izarin yellow K	$X_a = 1 \times 10^{-11}$

- 10. At 25 °C, the value of  $K_{sp}$  for PbCl<sub>2</sub>(s) is 1.6 x 10<sup>-5</sup>, and the value of  $K_{sp}$  for AgCl(s) is 1.8 x 10<sup>-10</sup>.
  - A. If 60.0 mL of 0.0400 M NaCl(aq) is added to 60.0 mL of 0.0300 M Pb(NO<sub>3</sub>)<sub>2</sub>(aq), will a precipitate form? Assume that volumes are additive. Show calculations to support your answer.
  - B. Calculate the equilibrium value of [Pb<sup>+2</sup>(aq)] in a 1.00 L saturated PbCl<sub>2</sub> solution to which 0.250 mole of NaCl(s) has been added. Assume no volume change occurs.
  - C. If 0.100 M NaCl(aq) is slowly added to a beaker containing both 0.120 M AgNO<sub>3</sub>(aq) and 0.150 M Pb(NO<sub>3</sub>)<sub>2</sub>(aq) at 25 °C, which will precipitate first, AgCl(s) or PbCl<sub>2</sub>(s)? Show calculations to support your answer.
- 11. Calculate the molar solubility of lead(II) chloride ( $K_{sp} = 1.6 \times 10^{-5}$ ) in 0.050 M CaCl<sub>2</sub>(aq).
- 12. The solubility of  $Mg(OH)_2(s)$  in a particular buffer solution is 0.65 g/L. What must be the pH of the buffer solution?  $K_{sp}$  of  $Mg(OH)_2(s)$  is 1.8 x  $10^{-11}$ .
- 13. What minimum concentration of aqueous ammonia would be needed to dissolve 1.00 g of silver chloride and keep it in solution. The volume of the solution is to be 200 mL. The overall reaction is

$$AgCl(s) + 2NH_3 \rightarrow Ag(NH_3)_{2^+} + Cl$$

The solubility product of silver chloride is  $1.7 \times 10^{-10}$ , and the dissociation constant for  $Ag(NH_3)_2^+$  is  $6.0 \times 10^{-8}$ .

- 14. Sodium thiosulfate ("hypo") is used as a fixer for photographic films and papers. The thiosulfate ion complexes with silver salts that are unexposed to light, and thus dissolves them. The exposed silver salts were "developed" previously, and at this "fixing" stage are metallic silver, which does **not** complex. If it is desired that a given fixing bath be capable of dissolving 40 g of silver chloride per liter, what must be
- (a) the concentration of free thiosulfate ion when the solution has dissolved the 40 g of silver chloride per liter
- (b) the initial concentration of thiosulfate ion before reaction with the silver chloride? The overall reaction to be considered is

$$AgCl(s) + 2S_2O_3^{2-} \rightarrow [Ag(S_2O_3)2]^{3-} + Cl-$$

The solubility product for silver chloride is  $1.7 \times 10^{-10}$ , and the dissociation constant for the thiosulfatoargentate anion is  $1 \times 10^{-13}$ .