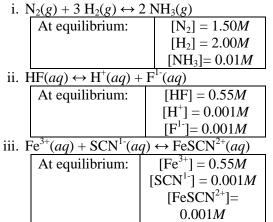
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# 8 – General Equilibrium/Solubility Equilibrium

# WS#1: INTRODUCTION TO EQUILIBRIUM

- 1. For the following three reactions,
  - a. Write the  $K_{eq}$  expression in terms of concentration,  $K_{c}$ .
  - b. Given the equilibrium concentrations, state whether each equilibrium is product-favored, reactantfavored, or fairly even ([products] ≈ [reactants]).
  - c. Calculate the value of  $K_c$ .



# Summarize:

Fill in the blanks with product-favored, reactant-favored, and approximately equal.

K <sub>c</sub>	State of Equilibrium
$K_{\rm c} >> 1$	
$K_{\rm c} << 1$	
$K_{\rm c} \approx 1$	

- 2. Knowing that pure water has a density of 1g/1mL calculate the mass of 1.00 Liter of water.
  - a. Calculate the number of moles in 1.00 L of  $H_2O$ .
  - b. What is the concentration (*M*) of water in water?
  - c. At this temperature, can you get more moles of water into this Liter of water?

The  $[H_2O]$  \_\_\_\_\_ (is / is not) constant.

# **Important Note:**

Since the concentrations of solids and liquids are constant, they are incorporated into the equilibrium constant,  $K_{eq}$ . That means, just leave them out of the  $K_c$  or  $K_p$  expression. Only include (g) and (aq)!

- 3. Write equilibrium expressions for each of the following reactions:
  - a.  $CaCO_3(s) \leftrightarrow CaO(s) + CO_2(g)$
  - b.  $Ni(s) + 4CO(g) \leftrightarrow Ni(CO)_4(g)$
  - c.  $5CO(g) + I_2O_5(s) \leftrightarrow I_2(g) + 5CO_2(g)$
  - d.  $Ca(HCO_3)_2(aq) \leftrightarrow CaCO_3(s) + H_2O(l) + CO_2(g)$
  - e.  $\operatorname{AgCl}(s) \leftrightarrow \operatorname{Ag}^+(aq) + \operatorname{Cl}^1(aq)$
- 4. Write the equilibrium expression in terms of partial pressures  $(K_p)$  for each of the following reactions.

Rate the reactions in order of their increasing tendency to proceed toward completion:

- a.  $4NH_3(g) + 3O_2(g) \leftrightarrow 2N_2(g) + 6H_2O(g)$   $K_p = 1x10^{228}$  atm b.  $N_2(g) + O_2(g) \leftrightarrow 2NO(g)$   $K_p = 5x10^{-31}$ c.  $2HF(g) \leftrightarrow H_2(g) + F_2(g)$   $K_p = 1x10^{-13}$ d.  $2NOCl(g) \leftrightarrow 2NO(g) + Cl_2(g)$   $K_p = 4.7x10^{-4}$  atm Question That You Should Packhla Takk
- A Question That You Should Be Able To Answer: Why don't the  $K_p$ 's in (b) and (c) have units?
- 7. a. Write the  $K_c$  expression for  $2 \operatorname{SO}_2(g) + \operatorname{O}_2(g) \leftrightarrow 2 \operatorname{SO}_3(g)$

Calculate the value of  $K_c$ :

c un	$c$ value of $M_c$ .	
	At equilibrium:	$[SO_2] = 1.50M$
		$[O_2] = 1.25M$
		$[SO_3] = 3.50M$

- b. If we **reverse** the equation, it is:  $2 \operatorname{SO}_3(g) \leftrightarrow 2 \operatorname{SO}_2(g) + \operatorname{O}_2(g)$ Write the  $K_c$  expression for this equation and calculate the new value of  $K_c$ : How does the expression and the value of  $K_c$  in 7(b) compare with those in 7(a)?
- c. If we now **multiply all of the coefficients by** <sup>1</sup>/<sub>2</sub>: SO<sub>3</sub>(g)  $\leftrightarrow$  SO<sub>2</sub>(g) + <sup>1</sup>/<sub>2</sub> O<sub>2</sub>(g) Write the  $K_c$  expression for this equation and calculate the new value of  $K_c$ : How do they compare with 7(b)?
- d. What would happen to the  $K_c$  expression and its value if we **doubled** the coefficients?

## Summarize:

Equation	K <sub>c</sub> expression & Value
doubled	
reversed	
halved	

8. Consider an equilibrium that occurs in two steps:  $H_2S(aa) \leftrightarrow H^+(aa) + HS^{1-}(aq)$ 

$$H_2S(aq) \leftrightarrow H(aq) + HS(aq)$$
$$\underline{HS^{1-}(aq)} \leftrightarrow \underline{H^+(aq)} + \underline{S^{2-}(aq)}$$

- a. Write the overall reaction.
- b. How do the  $K_c$ 's for the two steps ( $K_{c1}$  &  $K_{c2}$ ) relate to the  $K_c$  of the overall reaction ( $K_c$ )?

## WS#2: EQUILIBRIUM (K<sub>EQ</sub>) EXPRESSION

- 1. Write the expressions for the equilibrium constant  $K_c$  for the following reactions: a.  $4NH_3(g) + 7O_2(g) \leftrightarrow 4NO_2(g) + 6H_2O(l)$  b.  $3O_2(g) \leftrightarrow 2O_3(g)$
- 2. Write the equilibrium constant expressions for the following reactions. How are they related to one another? a.  $2N_2O(g) + 3O_2(g) \leftrightarrow 4NO_2(g)$ c.  $4NO_2(g) \leftrightarrow 2N_2O(g) + 3O_2(g)$

b. 
$$N_2O(g) + \frac{3}{2}O_2(g) \leftrightarrow 2NO_2(g)$$

3. Put the following *K* values in order of increasing product-favored ability.

- a.  $K = 4x10^{-5}$  b.  $K = 2x10^{-9}$  c.  $K = 7x10^{-5}$  d.  $K = 3x10^{-3}$ 4. Which of the following equations does  $K_c = K_p$ ? a.  $PCl_5(g) \leftrightarrow PCl_3(g) + Cl_2(g)$  d.  $H_2O(g) + CO \leftrightarrow H_2(g) + CO_2(g)$ b.  $2NOCl(g) \leftrightarrow 2NO(g) + Cl_2(g)$  e.  $2NO(g) \leftrightarrow N_2(g) + O_2(g)$ 
  - c.  $CaCO_3(s) \leftrightarrow CaO(s) + CO_3(g)$
- 5. If the mechanism of a chemical equilibrium consists of two reversible elementary steps, each with its own equilibrium constant  $K_{c1}$  and  $K_{c2}$ , what expression relates the equilibrium constant  $K_c$  for the overall equilibrium to the two constants  $K_{c1}$  and  $K_{c2}$ ?
- 6. Given the following equations:  $H_2O(g) + CO(g) \leftrightarrow H_2(g) + CO_2(g)$   $FeO(s) + CO(g) \leftrightarrow Fe(s) + CO_2(g)$   $Calculate the K_c value for: Fe(s) + H_2O(g) \leftrightarrow FeO(s) + H_2(g)$   $K_c = 4.8$   $K_c = 0.48$   $K_c = 0.48$   $K_c = 2.48$   $K_c = 2.48$  $K_c = 2.$

7. Consider the equilibrium:  $2 \operatorname{SO}_2(g) + \operatorname{O}_2(g) \leftrightarrow 2 \operatorname{SO}_3(g)$   $K_c = 4.36$ Calculate the value of "Q" for a situation in which  $[\operatorname{SO}_2] = 2.00M$ ,  $[\operatorname{O}_2] = 1.50M$ , and  $[\operatorname{SO}_3] = 1.25M$ . Does this mixture shift toward the reactants or products to reach equilibrium?

- 8. When converting  $K_c$  to  $K_p$  use the equation  $K_p = K_c(RT)^{\Delta n}$ . Where R = 0.0821, *T* is in Kelvin, and  $\Delta n$  is the difference in moles of gas between products and reactants ( $\Delta n = n_{\text{products}} n_{\text{reactants}}$ ). Write the  $K_p$  expression for the reaction in question (7) and calculate its value at 0°C. (Ans: 0.195)
- 9. The equation  $2NH_3(g) \leftrightarrow N_2(g) + 3H_2(g)$  has a value of  $K_c = 2.7 \times 10^{-4}$ . At STP, what is the value of  $K_p$ ? (Ans: 0.136)
- 10. For the equation,  $NH_4I(s) \leftrightarrow NH_3(g) + HI(g)$ , the total pressure is 4.2atm at equilibrium. What is  $K_p$ ? (Ans: 4.41)

#### WS#3: RICE TABLES

1. Consider the equilibrium:  $2N_2O(g) + O_2(g) \leftrightarrow 4NO(g)$ . 3.00 moles of NO(g) are introduced into a 1.00-Liter evacuated flask. When the system comes to equilibrium, 1.00 mole of  $N_2O(g)$  has formed. Determine the equilibrium concentrations of each substance. Calculate the  $K_c$  for the reaction based on these data.

	2 N <sub>2</sub> O	O <sub>2</sub>	4 NO
initial			
change			
equilibrium			

Remember: The "ice" box may be used with moles, molarity, or Liters (for gaseous equilibria)... never grams.

- 2. For the equation,  $H_2O(g) + CO(g) \leftrightarrow CO_2(g) + H_2(g) K_c = 0.235$ . If 2.00 moles of each  $H_2O$  and CO are put into a 10.0L container, what is the concentration of all species at equilibrium?(Ans:  $[CO_2] = [H_2] = 0.065$ ;  $[H_2O] = [CO] = 0.135$ )
- 3. For the equation,  $SO_2Cl_2(g) \leftrightarrow SO_2(g) + Cl_2(g) \tilde{K_p} = 4.8$ . If enough  $SO_2Cl_2$  is put into a container so its pressure is 8.00atm, what is the equilibrium pressure of all species? What is the total pressure? (Ans: Psoc = A-25atm, Psoc = 3.75, Pr = 12.25atm.)
- 4. For the reaction:  $PCl_5(g) \leftrightarrow PCl_3(g) + Cl_2(g)$ , if the initial pressure of  $PCl_5$  is 2.00atm and at equilibrium it is 15% dissociated, what is  $K_p$ ? (Ans: 0.0529)
- 5. Chlorine molecules will dissociate at high temperatures into chlorine atoms. At 3000°C, for example,  $K_c$  for the equilibrium shown is 0.55. If the partial pressure of chlorine molecules is 1.5atm, calculate the partial pressure of the chlorine atoms:  $Cl_2(g) \leftrightarrow 2Cl(g)$ . (Ans: 15atm)
- 6. Suppose that 0.50 moles of hydrogen gas, 0.50 moles of iodine gas, and 0.75 moles of hydrogen iodide gas are introduced into a 2.0 Liter vessel and the system is allowed to reach equilibrium.  $H_2(g) + I_2(g) \leftrightarrow 2 HI(g)$ . Calculate the concentrations of all three substances at equilibrium. At the temperature of the experiment,  $K_c$  equals  $2.0 \times 10^{-2}$ . (Ans:  $[H_2] = [I_2] = 0.41M$  and [HI] = 0.06M)

- 7. When 2.0mol of carbon disulfide and 4.0mol of chlorine are placed in a 1.0 Liter flask, the following equilibrium system results. At equilibrium, the flask is found to contain 0.30mol of carbon tetrachloride. What quantities of the other components are present in this equilibrium mixture?  $CS_2(g) + 3Cl_2(g) \leftrightarrow S_2Cl_2(g) + CCl_4(g)$  (Ans:  $[CS_2] = 1.7M$ ,  $[Cl_2] = 3.1M$ , and  $[S_2Cl_2] = 0.30M$ )
- 8. 3.0 moles each of carbon monoxide, hydrogen, and carbon are placed in a 2.0 Liter vessel and allowed to come to equilibrium according to the equation:  $CO(g) + H_2(g) \leftrightarrow C(s) + H_2O(g)$ . If the equilibrium constant at the temperature of the experiment is 4.0, what is the equilibrium concentration of water vapor? (Ans:  $[H_2O] = 1.0M$ )
- 9. Nitrosyl chloride NOCl decomposes to nitric oxide and chlorine when heated:  $2NOCl(g) \leftrightarrow 2NO(g) + Cl_2(g)$ . At 600K, the equilibrium constant  $K_p$  is 0.060. In a vessel at 600K, there is a mixture of all three gases. The partial pressure of NOCl is 675torr, the partial pressure of NO is 43torr and the partial pressure of chlorine is 23torr.
  - a. What is the value of the reaction quotient? (Ans: 0.093)
  - b. Is the mixture at equilibrium? (Ans: no)
  - c. In which direction will the system move to reach equilibrium? (Ans: shift to left, Q > K)
  - d. When the system reaches equilibrium, what will be the partial pressures of the components in the system? (Ans:  $P_{\text{NOCI}} = 681$ torr,  $P_{\text{NO}} = 37$ torr, and  $P_{\text{CI2}} = 20$ .torr)
- 10. Sulfuryl chloride decomposes at high temperatures to produce sulfur dioxide and chlorine gases:  $SO_2Cl_2(g) \leftrightarrow SO_2(g) + Cl_2(g)$ . At 375°C, the equilibrium constant  $K_c$  is 0.045. If there are 2.0 grams of sulfuryl chloride, 0.17 gram of sulfur dioxide, and 0.19 grams of chlorine present in a 1.0 Liter flask,
  - a. What is the value of the reaction quotient? (Ans:  $4.8 \times 10^{-4}$ )
  - b. Is the system at equilibrium? (Ans: no)
  - c. In which direction will the system move to reach equilibrium? (Ans: Q < K, so it will shift right)
- 11. Ammonium chloride is placed inside a closed vessel where it comes into equilibrium at 400°C according to the equation shown. Only these three substances are present inside the vessel. If  $K_p$  for the system at 400°C is 0.640, what is the pressure inside the vessel? NH<sub>4</sub>Cl(s)  $\leftrightarrow$  NH<sub>3</sub>(g) + HCl(g) (Ans: 1.60atm)
- 12. Bromine and chlorine react to produce bromine monochloride according to the equation,  $Br_2(g) + Cl_2(g) \leftrightarrow 2BrCl(g)$  and  $K_c = 36.0$  under the conditions of the experiment. If 0.180 moles of bromine gas and 0.180 moles of chlorine gas are introduced into a 3.0 Liter flask and allowed to come to equilibrium, what is the equilibrium concentration of the bromine monochloride? How much BrCl is produced? (Ans: [BrCl] = 0.090M and 0.27 mol)
- 13. When ammonia is dissolved in water, the following equilibrium is established. If the equilibrium constant is  $1.8 \times 10^{-5}$ , calculate the hydroxide ion concentration in the solution if 0.100mol of ammonia is dissolved in sufficient water to make 500 mL of solution. NH<sub>3</sub>(*aq*) + H<sub>2</sub>O(*l*)  $\leftrightarrow$  NH<sub>4</sub><sup>+</sup>(*aq*) + OH<sup>-</sup>(*aq*) (Ans: 0.0019*M*)

## WS#4: LE CHATELIER'S PRINCIPLE

WSł	74: LE CHATELIER'S PRINCIPLE						
1.	Consider the equilibrium $PCl_3(g) + Cl_2(g) \leftrightarrow PCl_3(g)$	$_{5}(g).$					
	How would the following changes affect the partial pressures of each gas at equilibrium?						
	$(\uparrow = increase; \downarrow = decrease; - = unchanged)$	·	<b>v</b>				
	a. addition of PCl <sub>3</sub>						
	b. removal of $Cl_2$						
	c. removal of PCl <sub>5</sub>						
	d. decrease in the volume of the container						
	e. addition of He without change in volume						
2.	How will each of the changes in question (3) affect	ct the $K_{eq}$ ? $(\uparrow =$	increase; $\downarrow = de$	ecrease; — =	unchanged)		
	a b c	d			U /		
3.	Indicate how each of the following changes affect	ts the amount of	each gas in the s	system below,	for which $\Delta H_{\text{reaction}}$		
3.	Indicate how each of the following changes affect $= +9.9$ kcal.		each gas in the s $\downarrow$ = decrease; -				
3.		$(\uparrow = increase;$		— = unchang	ged)		
3.	<ul><li>= +9.9 kcal.</li><li>a. addition of CO<sub>2</sub></li></ul>	$(\uparrow = increase;$	$\downarrow$ = decrease; -	— = unchang	ged)		
3.	= +9.9 kcal.	$(\uparrow = increase;$	$\downarrow$ = decrease; -	— = unchang	ged)		
3.	<ul><li>= +9.9 kcal.</li><li>a. addition of CO<sub>2</sub></li></ul>	$(\uparrow = increase;$	$\downarrow$ = decrease; -	— = unchang	ged)		
3.	<ul> <li>= +9.9 kcal.</li> <li>a. addition of CO<sub>2</sub></li> <li>b. addition of H<sub>2</sub>O</li> </ul>	$(\uparrow = increase;$	$\downarrow$ = decrease; -	— = unchang	ged)		
3.	<ul> <li>= +9.9 kcal.</li> <li>a. addition of CO<sub>2</sub></li> <li>b. addition of H<sub>2</sub>O</li> <li>c. addition of a catalyst</li> </ul>	$(\uparrow = increase;$	$\downarrow$ = decrease; -	— = unchang	ged)		
	<ul> <li>= +9.9 kcal.</li> <li>a. addition of CO<sub>2</sub></li> <li>b. addition of H<sub>2</sub>O</li> <li>c. addition of a catalyst</li> <li>d. increase in temperature</li> <li>e. decrease in the volume of the container</li> </ul>	$(\uparrow = \text{increase}; H_2(g) + $	$ \begin{array}{c} \downarrow = \text{decrease;} \\ CO_2(g) \leftrightarrow \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\$	— = unchang	ged)		
4.	<ul> <li>= +9.9 kcal.</li> <li>a. addition of CO<sub>2</sub></li> <li>b. addition of H<sub>2</sub>O</li> <li>c. addition of a catalyst</li> <li>d. increase in temperature</li> </ul>	$(\uparrow = \text{increase}; H_2(g) + $	$ \begin{array}{c} \downarrow = \text{decrease;} \\ CO_2(g) \leftrightarrow \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\$	— = unchang	ged)		

5.	Consider the equilibrium: How will the amount of chemicals at equilibrium be affected by	$2N_2O(g)$	+	$O_2(g)$	$\leftrightarrow$	4NO( <i>g</i> )
	a. adding N <sub>2</sub> O					
	b. removing $O_2$					
	c. increasing the volume of the container					
6.	d. adding a catalyst For the reaction,	$\overline{4\mathrm{NH}}_{2}(g)$	$+30_{2}$	$(g) \leftrightarrow 2$	N2(9) -	+ $6\overline{\text{H}_2\text{O}}(l)$
0.	How will the concentration of each chemical be affected by	11 (11)(8)	1 502	(8) 21	12(8)	
	a. adding $O_2$ to the system					
	b. adding $N_2$ to the system					
	c. removing $H_2O$ from the system					
	d. decreasing the volume of the container					

7. The following reaction is exothermic:  $Ti(s) + 2 Cl_2(g) \leftrightarrow TiCl_4(g)$ . List all the ways the yield of the product  $TiCl_4$ could be increased.

#### WS#5: READING A SOLUBILITY CURVE

- 1. Which substance is the most soluble at  $0^{\circ}$ C? At  $100^{\circ}C?$
- 2. How many grams of substance "B" will dissolve in 100g of water at 60°C? How about in 400g of water at the same temperature?
- Which substance shows the least change in solubility 3. form  $0^{\circ}C - 100^{\circ}C$  ?
- As you increase the temperature of the water, what 4. happens to the solubility of "A"?
- 5. As you increase the temperature of the water, what happens to the solubility of "B"?
- 6. As you increase the temperature of the water, what happens to the solubility of "C"?
- 7. Which substance(s) are probably a solid?
- 8. Which substance(s) are probably a gas?
- 9. If you put 40g of "A" into 100g of water at 50°C the solution would be?
- 10. If you put 40g of "B" into 100g of water at 50°C the solution would be?
- 11. If you put 40g of "A" into 100g of water at 20°C the solution would be?
- 12. Describe how to make a supersaturated solution of "B" at 50°C.

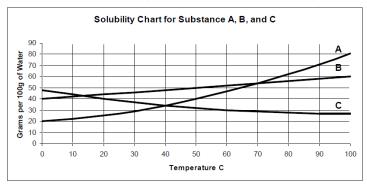
## WS#6: Solubility, *K*<sub>sp</sub>, and Common Ion Effect

Write a net ionic equation and the solubility product expression  $(K_{sp})$  for the following precipitation reaction.

- Solutions of sodium sulfate and calcium bromide are mixed. 1.
- 2. Solutions of aluminum (III) acetate and lithium hydroxide are mixed.
- 3. Solutions of iron (III) sulfate and sodium sulfide are mixed.
- 4. Solutions of aluminum (III) sulfate and calcium hydroxide are mixed.
- 5. Solutions of potassium chromate and lead (II) acetate are mixed.
- Solutions of silver (I) nitrate and ammonium sulfide are mixed. 6.
- Write the dissociation equation, solubility expression ( $K_{sp}$ ), and appropriate substitution scenario ( $x^2$ ,  $4x^3$ ,  $27x^4$ , or 7. 108x<sup>5</sup>) for the following ionic solids  $(x^2, 4x^3, 27x^4, \text{ or } 108x^5)$ . a.  $CaCO_3$ b.  $Al(OH)_3$ c.  $Mg_3(PO_4)_2$ d.  $Ag_2C_2O_4$

- 8. Determine the  $K_{sp}$  of silver (I) bromide, AgBr, given that its molar solubility is  $5.71 \times 10^{-7}$  moles per liter. (Ans:  $3.26 \times 10^{-13}$ ) 9. Determine the  $K_{sp}$  of zinc (II) bromide (ZnBr<sub>2</sub>), given that its molar solubility is  $2.52 \times 10^{-8}$  mole per liter. (Ans:  $6.40 \times 10^{-23}$ ) 10. Silver chloride, AgCl, has a  $K_{sp} = 1.77 \times 10^{-10}$ . Calculate its solubility in moles per liter. (Ans:  $1.33 \times 10^{-5} \text{ mol/L}$ ) 11. Aluminum phosphate, AlPO<sub>4</sub>, has a  $K_{sp}$  of  $9.83 \times 10^{-21}$ . What is its molar solubility in pure water? (Ans:  $9.91 \times 10^{-11} \text{ mol/L}$ )
- 12. A 100.0mL sample is removed from a sea water solution saturated in MgF<sub>2</sub> at 18°C. The water is completely
- evaporated from the sample and a 7.6mg deposit of  $MgF_2(s)$  is obtained. What is the  $K_{sp}$  for  $MgF_2$ ? (Ans: ) 13. Calculate the molar solubility of silver (I) chromate in water at 25°C. Find the  $K_{sp}$  for silver (I) chromate in the
- solubility table. (Ans: )
- 14. Calculate the solubility of solid Ca<sub>3</sub>(PO<sub>4</sub>)<sub>2</sub> ( $K_{sp} = 1.3 \times 10^{-32}$ ) in a 0.20*M* Na<sub>3</sub>PO<sub>4</sub> solution. (Ans: )

Look at the solubility curves on the graph below and then answer the following questions



- 15. The solubility of Ce(IO<sub>3</sub>)<sub>3</sub> in a 0.20*M* KIO<sub>3</sub> solution is  $4.4 \times 10^{-8}$  mol/L. Calculate  $K_{sp}$  for Ce(IO<sub>3</sub>)<sub>3</sub>. (Ans: )
- 16. A solution contains  $1.0 \times 10^{-5} M$  Na<sub>3</sub>PO<sub>4</sub>. What is the minimum concentration of AgNO<sub>3</sub> that would cause
- precipitation of solid Ag<sub>3</sub>PO<sub>4</sub> ( $K_{sp} = 1.8 \times 10^{-18}$ )? (Ans: ) 17. A solution is  $1 \times 10^{-4} M$  in NaF, Na<sub>2</sub>S, and Na<sub>3</sub>PO<sub>4</sub>. What would be the order of precipitation as a source of Pb<sup>2+</sup> is added gradually to the solution? The relevant  $K_{sp}$  values are  $K_{sp}$  (PbF<sub>2</sub>) =  $4 \times 10^{-8}$ ,  $K_{sp}$  (PbS) =  $7 \times 10^{-29}$ , and  $K_{sp}$  (Pb<sub>3</sub>(PO<sub>4</sub>)<sub>2</sub>) =  $1 \times 10^{-54}$ . (Ans: )

# **Answer Key**

WS#3

- 1. NA
- 2.  $4.0 \times 10^{-8}$
- 3.  $7.26 \times 10^{-9}$
- 4.  $6.54 \times 10^{-5} M$
- 5.  $1.1 \times 10^{-73}$
- 6.  $3.3 \times 10^{-8} \text{mol/L}$
- 7.  $6.4 \times 10^{-8} \text{mol/L}$
- 8. 2.3x10<sup>-11</sup>mol/L 9. 3.5x10<sup>-10</sup>

## WS#4

- 1.  $Q_{sp} = 5.32 \times 10^{-10}$ , yes a ppt will form 2.  $[Mg^{2+}] = 2.1 \times 10^{-6} M$ ,  $[F^{1-}] = 5.50 \times 10^{-2} M$ 3.  $[I^{1-}] = 2.6 \times 10^{-3} M$  for PbI<sub>2</sub> and  $[I^{1-}] = 5.3 \times 10^{-8} M$  for CuI; CuI will ppt first
- 4.  $[AgNO_3] > 5.6x10^{-5}M$
- 5. PbS(s) will ppt first, followed by  $Pb_3(PO_4)_2(s)$  and  $PbF_2$  will ppt last.