



Unit 3: Solubility Equilibrium

Chem 11 Review
Preparation for Solubility Equilibrium

Chem 11 Review Preparation for Solubility Equilibrium

CHEM 11 Review

It is expected that the student understands the concept of:

1. Strong electrolytes,
2. Weak electrolytes and
3. Nonelectrolytes.

CHEM 0011 – Unit 9 - Solubility of Ionic Compounds

<http://nobel.scas.bcit.ca/courses/wpmu/chem0011/unit-9/>

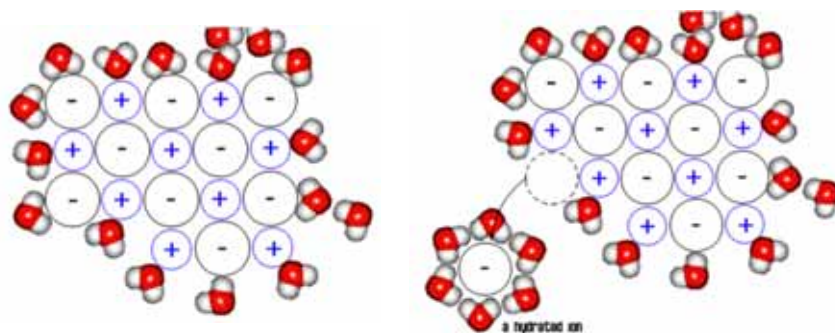
Chem 11 Review

Preparation for Solubility Equilibrium

CHEM 11 Review

The dissolution process.

<http://www.chem.iastate.edu/group/Greenbowe/sections/projectfolder/flashfiles/thermochem/solutionSalt.html>



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3

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1. Strong Electrolyte – Sodium chloride, NaCl

- Forms from metal and nonmetal
 eg: $\text{NaCl (s)} \rightarrow \text{Na}^+ \text{(aq)} + \text{Cl}^- \text{(aq)}$
 eg: $\text{CaCl}_2 \text{(s)} \rightarrow \text{Ca}^{2+} \text{(aq)} + 2 \text{Cl}^- \text{(aq)}$
 eg: $\text{NH}_4\text{NO}_3 \text{(s)} \rightarrow \text{NH}_4^+ \text{(aq)} + \text{NO}_3^- \text{(aq)}$
- Totally ionizes in water.
- Forms electrolytic solutions (conducts electricity) with charged ions.
- Has ionic bonds.
- Commonly called “salts” which are crystalline solids (high melting points and boiling points) and dissolves well.

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4

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2. Weak Electrolyte – Hydrofluoric acid, HF

Molecular Compounds – are weak or non-electrolytes when dissolved in water

- Forms from nonmetal with nonmetal
eg: $\text{HF (aq)} \rightleftharpoons \text{H}^+ \text{ (aq)} + \text{F}^- \text{ (aq)}$
eg: $\text{H}_2\text{O (aq)} \rightleftharpoons \text{H}^+ \text{ (aq)} + \text{OH}^- \text{ (aq)}$
- Partially ionizes in water.
- HF is a weak electrolyte (weakly conducts electricity).
- Has covalent bonds.
- Commonly called “molecules” and can be solid, liquids or gas.

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5

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3. Nonelectrolyte – Methanol, CH_3OH

- Does not ionize in water.
- CH_3OH does not conduct electricity.
- Has covalent bonds.
- Upon dissolving, the methanol molecules remain intact.

[http://highered.mcgraw-hill.com/olcweb/cgi/pluginpop.cgi?it=swf::525::530::/sites/dl/free/0072857684/322633/07_stg_wk_nonelelytes.swf::Strong Electrolyte, Weak Electrolyte, and Nonelectrolyte](http://highered.mcgraw-hill.com/olcweb/cgi/pluginpop.cgi?it=swf::525::530::/sites/dl/free/0072857684/322633/07_stg_wk_nonelelytes.swf::Strong%20Electrolyte,%20Weak%20Electrolyte,%20and%20Nonelectrolyte)

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6

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State whether the following substance is expected to form ionic or molecular solutions.

CuSO_4 (s) ionic solution

CH_4 (g) molecular solution

CrCl_3 (s) ionic solution

NaCH_3COO (s) ionic solution

In this course, we are focus on the IONIC compounds.

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7

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Preparation for Solubility Equilibrium

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Solubility of a substance:

Describes the amount of the substance that will dissolve in a solvent like water at a given temperature.

A solution which has dissolved the **MAXIMUM** amount of substance at a given temperature is said to be a **SATURATED** solution.

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8

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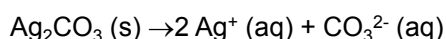
Preparation for Solubility Equilibrium

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It is expected that the student understands the concept of molarity and solution composition.

Eg: What is the concentration of all the ions present in a saturated solution of Ag_2CO_3 having a concentration of $1.2 \times 10^{-4} \text{ M}$? Calculate the concentration of all the ions in the solution.

When Ag_2CO_3 dissolves, the ions in solution are $\text{Ag}^+ (\text{aq})$ and $\text{CO}_3^{2-} (\text{aq})$.
The ratio of the ions is 2:1 $\text{Ag}^+ : \text{CO}_3^{2-}$.



$$[\text{Ag}^+] = 2.4 \times 10^{-4} \text{ M}$$

$$[\text{CO}_3^{2-}] = 1.2 \times 10^{-4} \text{ M}$$

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9

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It is expected that the student understands the concept of dilution and how to calculate the dilution occurring when solutions are mixed.

CHEM 0011 – Unit 9 - Preparation of Solutions by Dilution

http://nobel.scas.bcit.ca/wiki/index.php/Solution_dilution

$$M_1V_1 = M_2V_2$$

where M_1 = concentration of the original solution
 V_1 = volume of the original solution

where M_2 = concentration of the diluted solution
 V_2 = volume of the diluted solution

Note: that the units on M_1 and M_2 must be the same and the units on V_1 and V_2 must be the same

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10

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Dilution Problem

$$M_1V_1 = M_2V_2$$

If 5.0 mL of 0.020 M Cl^- is added to 15.0 mL of 0.012 M Br^- , what is the molarity of Cl^- and Br^- ions?

Consider the Cl^- ions:

$$M_1 = [\text{Cl}^-]_{\text{final}} = ?$$

$$M_2 = [\text{Cl}^-]_{\text{original}} = 0.020 \text{ M}$$

$$V_1 = \text{Total volume of final solution} = 20.0 \text{ mL}$$

$$V_2 = \text{Volume of original } \text{Cl}^- \text{ solution} = 5.0 \text{ mL}$$

$$[\text{Cl}^-]_{\text{final}} = M_2V_2/V_1 = (0.020 \text{ M})(5.0 \text{ mL}) / 20.0 \text{ mL} = 0.0050 \text{ M}$$

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11

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Preparation for Solubility Equilibrium

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Dilution Problem

$$M_1V_1 = M_2V_2$$

If 5.0 mL of 0.020 M Cl^- is added to 15.0 mL of 0.012 M Br^- , what is the molarity of Cl^- and Br^- ions?

Consider the Br^- ions:

$$M_1 = [\text{Br}^-]_{\text{final}} = ?$$

$$M_2 = [\text{Br}^-]_{\text{original}} = 0.012 \text{ M}$$

$$V_1 = \text{Total volume of final solution} = 20.0 \text{ mL}$$

$$V_2 = \text{Volume of original } \text{Br}^- \text{ solution} = 15.0 \text{ mL}$$

$$[\text{Br}^-]_{\text{final}} = M_2V_2/V_1 = (0.012 \text{ M})(15.0 \text{ mL}) / 20.0 \text{ mL} = 0.0090 \text{ M}$$

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12

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It is expected that the student understands the concept of writing chemically correct chemical formula when given cations and anions.

Chem 0011 – Unit 5 – Writing Chemical Formulae

http://nobel.scas.bcit.ca/wiki/index.php/Writing_chemical_formulae

Eg: Write the chemical formula for the compound formed between:

- | | |
|--|--|
| 1. Na and Br: | NaBr |
| 2. K and I: | KI |
| 3. Ba and F: | BaF ₂ |
| 4. K ⁺ and nitrate ion: | KNO ₃ |
| 5. Ca ²⁺ and acetate ion: | Ca(C ₂ H ₃ O ₂) ₂ |
| 6. ammonium ion and Cl ⁻ ion: | NH ₄ Cl |
| 7. Fe ³⁺ and carbonate ion: | Fe ₂ (CO ₃) ₃ |
| 8. Mg ²⁺ and phosphate ion: | Mg ₃ (PO ₄) ₂ |

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13

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It is expected that the student knows how to balance chemical equations.

Chem 0011 – Balancing Chemical Equations

http://nobel.scas.bcit.ca/wiki/index.php/Balancing_chemical_equations

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14

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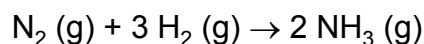
Preparation for Solubility Equilibrium

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It is expected that the student understands calculations involving limiting reagent and solve stoichiometry-related reactions.

If 55.0 grams of N_2 (g) is reacted with 55.0 g H_2 (g), how many grams of ammonia gas, NH_3 (g), is formed?

Write the balanced chemical equation:



Convince yourself that:

1. N_2 (g) is the limiting reagent
2. H_2 (g) is in excess
3. The answer is 66.9 g NH_3 (g).

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15

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Preparation for Solubility Equilibrium

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It is expected that the student understands the concept of predicting solubility using the solubility table.

On a test or exam, a solubility table will be provided. The student is expected to know how to use it.

Low solubility – does not dissolve in water.

Predict whether the following is soluble in water:

PbI_2	low solubility
MgSO_4	soluble
CuCO_3	low solubility
LiCl	soluble

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16

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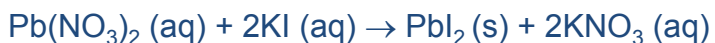
Preparation for Solubility Equilibrium

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It is expected that the student understands the concept of predicting solubility using the solubility table.

Eg: Use the solubility table to predict whether the following double replacement reaction will occur? In other words, what happens when solutions of $\text{Pb}(\text{NO}_3)_2$ (aq) and KI (aq) are mixed? If a reaction occurs, write the balanced chemical equation.

Answer:



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17

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Preparation for Solubility Equilibrium

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It is expected that the student understands the concept of predicting solubility using the solubility table.

Putting it all together!

Eg: What happens when 5.34 g of calcium chloride and 9.85 g of aluminum chloride are dissolved in 4.50 L of solution?

- (a) Write the reaction if it occurs.
- (b) Find the concentrations of all the ions in solution.

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18

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Preparation for Solubility Equilibrium

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Eg: What happens when 5.34 g of calcium chloride and 9.85 g of aluminum chloride are dissolved in 4.50 L of solution?

- (a) Write the reaction if it occurs.
- (b) Find the concentrations of all the ions in solution.

Answer:

(a) No reaction occurs. All ions stay in solution.

(b) FW (CaCl_2) = 111 g/mole; FW (AlCl_3) = 133.36 g/mole

CaCl_2 : 5.34 g / 111 g/mole = 0.0481 moles CaCl_2 produces 0.0481 moles Ca^{2+} ions
 = 2(0.0481) moles of Cl^- ions.

AlCl_3 : 9.85 g / 133.36 g/mole = 0.0739 moles AlCl_3 produces 0.0739 moles Al^{3+} ions.
 = 3(0.0739) moles Cl^- ions.

$$[\text{Ca}^{2+}] = 0.0481 \text{ moles} / 4.50 \text{ L} = 0.0107 \text{ M}$$

$$[\text{Al}^{3+}] = 0.0739 \text{ moles} / 4.50 \text{ L} = 0.0164 \text{ M}$$

$$[\text{Cl}^-] = [2(0.0481) + 3(0.0739)] / 4.50 \text{ L} = 0.0706 \text{ M}$$

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19

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Preparation for Solubility Equilibrium

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- (a) What happens when NaCl (aq) and AgNO_3 (aq) solutions are mixed?
- (b) Find the concentrations of all the ions in solution when 2.72 g of NaCl and 3.75 g AgNO_3 are dissolved in 5.20 L solution.
- (c) Write the balanced chemical equation.
- (d) Write the complete ionic equation.
- (e) Write the net ionic equation.
- (f) Identify the spectator ions.

Answers:

(a) and (c) NaCl (aq) + AgNO_3 (aq) \rightarrow NaNO_3 (aq) + AgCl (s)

(b) $[\text{Na}^+] = 0.00894 \text{ M}$, $[\text{Cl}^-] = 0.0047 \text{ M}$, $[\text{Ag}^+] \approx 0 \text{ M}$, $[\text{NO}_3^-] = 0.00424 \text{ M}$

(d) Na^+ (aq) + Cl^- (aq) + Ag^+ (aq) + NO_3^- (aq) \rightarrow Na^+ (aq) + NO_3^- (aq) + AgCl (s)

(e) Ag^+ (aq) + Cl^- (aq) \rightarrow AgCl (s)

(f) Na^+ (aq) and NO_3^- (aq) are spectator ions.

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20



Solubility Equilibrium

(Hebden Unit 3 – page 73-108)

Solubility Equilibrium

Hebden – Unit 3 (page 73-108)

Solubility Equilibrium

We will cover the following topics:

1. Define the term “solubility”.
2. What is a saturated solution?
3. What is solubility equilibrium?
4. What is the meaning of K_{sp} and what is the solubility product constant expression?
5. Calculate K_{sp} from solubility data.
6. Compute molar solubility from K_{sp} .

Solubility Equilibrium

Hebden – Unit 3 (page 73-108)

1. Define the term “solubility”.

Solubility is a measure of how much a solute can dissolve in a solvent at a given temperature.

In this course, we will focus on water as the solvent.

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23

Solubility Equilibrium

Hebden – Unit 3 (page 73-108)

2. What is a saturated solution?

A solution which has dissolved the **MAXIMUM** amount of substance at a given temperature is said to be a **SATURATED** solution.

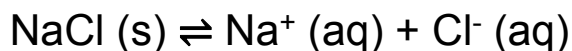
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24

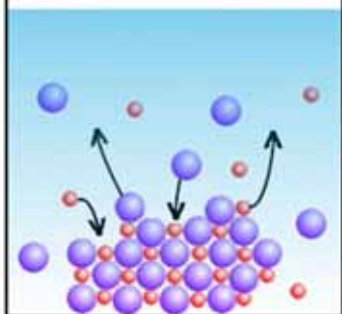
Dynamic Equilibrium

Hebden – Unit 2 (page 37-69)

3. What is solubility equilibrium?



A saturated sodium chloride solution



A saturated solution has an established equilibrium between dissolved and un-dissolved particles.

Rate of dissolution = Rate of crystallization

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25

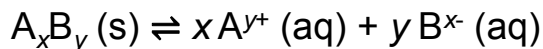
Solubility Equilibrium

Hebden – Unit 3 (page 73-108)

Solubility Equilibrium

Compounds of **low solubility** form **EQUILIBRIA** involving the solid and its aqueous ions.

For a general compound $\text{A}_x\text{B}_y \text{ (s)}$ which is slightly soluble in water, the equilibrium between the solid and its soluble ions is established.



$$*K_{\text{eq}} = [\text{A}^{y+}]^x [\text{B}^{x-}]^y$$

* $[\text{A}_x\text{B}_y]$ is not included in the equilibrium expression because A_xB_y is a solid.

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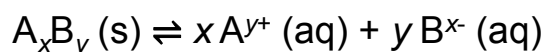
26

Solubility Equilibrium

Hebden – Unit 3 (page 73-108)

Solubility Equilibrium

For a general equilibrium



where $A_xB_y(s)$ is a compound with low solubility.

$$K_{eq} = [A^{y+}]^x [B^{x-}]^y$$

Since the K_{eq} is the product of the soluble ions, a special name is given to the K_{eq} , called

K_{sp} , Solubility Product Equilibrium Constant

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27

Solubility Equilibrium

Hebden – Unit 3 (page 73-108)

4. What is the meaning of K_{sp} and what is the solubility product constant expression?

Look at the Solubility Product Constant Reference Sheet

SOLUBILITY PRODUCT CONSTANTS AT 25°C

Name	Formula	K_{sp}
Silver bromide	AgBr	5.4×10^{-13}
Silver chloride	AgCl	1.8×10^{-10}

What do these numbers mean?

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28

Solubility Equilibrium

Hebden – Unit 3 (page 73-108)

Solubility Product Constant Reference Sheet

SOLUBILITY PRODUCT CONSTANTS AT 25°C

Name	Formula	K_{sp}
Silver bromide	AgBr	5.4×10^{-13}
Silver chloride	AgCl	1.8×10^{-10}

The solubility constant equilibrium is:



K_{sp} is the product of molar solubility of the ions in a saturated solution. This is the solubility product constant expression.

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29

Solubility Equilibrium

Hebden – Unit 3 (page 73-108)

Solubility Product Constant Reference Sheet

SOLUBILITY PRODUCT CONSTANTS AT 25°C

Name	Formula	K_{sp}
Silver bromide	AgBr	5.4×10^{-13}
Silver chloride	AgCl	1.8×10^{-10}

At 25°C, $K_{sp} (\text{AgCl}) > K_{sp} (\text{AgBr})$

$$[\text{Ag}^+][\text{Cl}^-] > [\text{Ag}^+][\text{Br}^-]$$

This means that AgCl is more soluble than AgBr.

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30

Solubility Equilibrium

Hebden – Unit 3 (page 73-108)

Solubility Product Constant Reference Sheet

Look up K_{sp} for strontium fluoride, SrF_2 . Write the solubility product constant expression for SrF_2 .

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31

Solubility Equilibrium

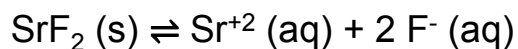
Hebden – Unit 3 (page 73-108)

Solubility Product Constant Reference Sheet

SOLUBILITY PRODUCT CONSTANTS AT 25°C

Name	Formula	K_{sp}
Strontium fluoride	SrF_2	4.3×10^{-9}

The solubility constant equilibrium is:



This is the solubility product constant expression.

$$K_{sp} = [\text{Sr}^{+2}][\text{F}^-]^2 = 4.3 \times 10^{-9}$$

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32

Solubility Equilibrium

Hebden – Unit 3 (page 73-108)

5. Calculate K_{sp} from solubility data.

One way to determine the value of K_{sp} for a salt is to measure its **solubility** (i.e. How much of the salt is required to produce a saturated solution.) Solubility is expressed in grams/L.

Molar solubility: is the number of moles of salt dissolved in one liter of its saturated solution.

That is, the concentration of a dissolved solid present in a saturated solution, expressed in molarity. (moles/L)

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33

Solubility Equilibrium

Hebden – Unit 3 (page 73-108)

5. Calculate K_{sp} from solubility data.

The solubility of AgBr in water was determined to be 1.38×10^{-4} g/L at 25°C. Calculate the K_{sp} for AgBr at that temperature.

Step 1: Convert 1.38×10^{-4} g/L to moles/L to get molar solubility of AgBr.

$$1.3 \times 10^{-4} \text{ g/L} \cdot \frac{1}{187.77 \text{ g/mole AgBr}} = 7.34 \times 10^{-7} \text{ moles/L}$$

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34

Solubility Equilibrium

Hebden – Unit 3 (page 73-108)

5. Calculate K_{sp} from solubility data.

The solubility of AgBr in water was determined to be 1.38×10^{-4} g/L at 25°C. Calculate the K_{sp} for AgBr at that temperature.

Step 2: Set up ICE table.

	AgBr (s)	\rightleftharpoons	Ag ⁺ (aq)	+	Br ⁻ (aq)
[I]	-		0		0
[C]			7.34×10^{-7}		7.34×10^{-7}
[E]			7.34×10^{-7}		7.34×10^{-7}

Initially no ions in the solution.

Stoichiometric ratio

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35

Solubility Equilibrium

Hebden – Unit 3 (page 73-108)

5. Calculate K_{sp} from solubility data.

Example 1: The solubility of AgBr in water was determined to be 1.38×10^{-4} g/L at 25°C. Calculate the K_{sp} for AgBr at that temperature.

Step 3: Write the K_{sp} expression.

$$K_{sp} = [\text{Ag}^+][\text{Br}^-] = (7.34 \times 10^{-7})(7.34 \times 10^{-7}) = 5.39 \times 10^{-13}$$

Look up the K_{sp} of AgBr in the Solubility Product Constant Reference Sheet

SOLUBILITY PRODUCT CONSTANTS AT 25°C

Name	Formula	K_{sp}
Silver bromide	AgBr	5.4×10^{-13}

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36

Solubility Equilibrium

Hebden – Unit 3 (page 73-108)

5. Calculate K_{sp} from solubility data.

Example 2: The molar solubility of silver chromate, Ag_2CrO_4 , in water is 6.7×10^{-5} moles/L at $25^\circ C$. What is the K_{sp} of Ag_2CrO_4 at $25^\circ C$?

	$Ag_2CrO_4 (s) \rightleftharpoons 2 Ag^+ (aq) + CrO_4^{2-} (aq)$		
[I]	-	0	0
[C]		$2(6.7 \times 10^{-5})^*$	6.7×10^{-5}
[E]		1.3×10^{-4}	6.7×10^{-5}

Stoichiometric ratio

* For each mole of Ag_2CrO_4 dissolved, there are 2 moles Ag^+ and 1 mole CrO_4^{2-} .

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37

Solubility Equilibrium

Hebden – Unit 3 (page 73-108)

5. Calculate K_{sp} from solubility data.

Example 2: The molar solubility of silver chromate, Ag_2CrO_4 , in water is 6.7×10^{-5} moles/L at $25^\circ C$. What is the K_{sp} of Ag_2CrO_4 at $25^\circ C$?

Write the K_{sp} expression.

$$K_{sp} = [Ag^+]^2[CrO_4^{2-}] = (1.3 \times 10^{-4})^2 (6.7 \times 10^{-5}) = 1.1 \times 10^{-12}$$

SOLUBILITY PRODUCT CONSTANTS AT $25^\circ C$

Name	Formula	K_{sp}
Silver chromate	Ag_2CrO_4	1.1×10^{-12}

Look up the K_{sp} of Ag_2CrO_4 in the Solubility Product Constant Reference Sheet

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38

Solubility Equilibrium

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6. Compute molar solubility from K_{sp} .

Example 1: What is the molar solubility of AgCl in pure water at 25°C?

	$\text{AgCl (s)} \rightleftharpoons \text{Ag}^+ \text{(aq)} + \text{Cl}^- \text{(aq)}$		
[I]	-	0	0
[C]		+x	+x
[E]		x	x

Stoichiometric ratio

$$K_{sp} = [\text{Ag}^+][\text{Cl}^-] = x^2 = 1.8 \times 10^{-10}$$

$$\text{Molar solubility} = x = 1.3 \times 10^{-5} \text{ M}$$

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39

Solubility Equilibrium

Hebden – Unit 3 (page 73-108)

6. Compute molar solubility from K_{sp} .

Example 2: Calculate the molar solubility of lead iodide in water from its K_{sp} at 25°C? $K_{sp} = 8.5 \times 10^{-9}$

	$\text{PbI}_2 \text{(s)} \rightleftharpoons \text{Pb}^{2+} \text{(aq)} + 2 \text{I}^- \text{(aq)}$		
[I]	-	0	0
[C]		+x	+2x
[E]		x	2x

Stoichiometric ratio

$$K_{sp} = [\text{Pb}^{2+}][\text{I}^-]^2 = (x)(2x)^2 = 4x^3 = 8.5 \times 10^{-9}$$

$$\text{Molar solubility} = x = 1.3 \times 10^{-3} \text{ M}$$

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40

Solubility Equilibrium

Hebden – Unit 3 (page 73-108)

Solubility Equilibrium

We have covered the following topics:

1. Define the term “solubility”.
2. What is a saturated solution?
3. What is solubility equilibrium?
4. What is the meaning of K_{sp} and what is the solubility product constant expression?
5. Calculate K_{sp} from solubility data.
6. Compute molar solubility from K_{sp} .

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41

Solubility Equilibrium

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Solubility Equilibrium

We will cover the following topics:

1. Common Ion Effect
 - Qualitatively using Le Chatelier's Principle
 - Quantitatively by K_{sp} calculation
2. Use K_{sp} to predict whether a precipitate will form in a solution (i.e. Compare Q with K_{sp})

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42

Solubility Equilibrium

Hebden – Unit 3 (page 73-108)

1. Common Ion Effect

Up to now, we have been studying the behaviour of low solubility salts dissolved in pure water.

What if other ions are present in the water?

Recall, we calculate the solubility of $\text{PbI}_2 (\text{s})$

Molar solubility PbI_2 in pure water $1.3 \times 10^{-3} \text{ M}$

Remember
this
number.

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43

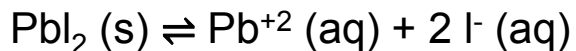
Solubility Equilibrium

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1. Common Ion Effect

Let's dissolve PbI_2 in a solution that already has sodium iodide, NaI , dissolved in it. How much of PbI_2 will dissolve?

To answer this question, think Le Chatelier's Principle. What does NaI do to the equilibrium?



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44

Solubility Equilibrium

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1. Common Ion Effect

Let's dissolve PbI_2 in a solution that already has sodium iodide, NaI , dissolved in it. How much of PbI_2 will dissolve?

1. Ask yourself, what is the solubility property of NaI . Look it up!!

SOLUBILITY OF COMMON COMPOUNDS IN WATER

The term soluble here means $> 0.1 \text{ mol/L}$ at 25°C

Negative Ions (Anions)	Positive Ions (Cations)	Solubility of Compounds
Chloride, Cl^- or Bromide, Br^- or Iodide, I^-	All others	Soluble
	Ag^+ , Pb^{2+} , Cu^+	Low Solubility

Is NaI soluble or insoluble in water?

CHEM 0012 Lecture Notes

45

Solubility Equilibrium

Hebden – Unit 3 (page 73-108)

1. Common Ion Effect

Let's dissolve PbI_2 in a solution that already has sodium iodide, NaI , dissolved in it. How much of PbI_2 will dissolve?

- Ask yourself, what is the solubility property of NaI . Look it up

Is NaI soluble or insoluble in water?

Answer: NaI is soluble!!

- What does it mean NaI is soluble?

It means that in solution,
 $\text{Na}^+ (\text{aq})$ and $\text{I}^- (\text{aq})$ ions are formed.

CHEM 0012 Lecture Notes

46

Solubility Equilibrium

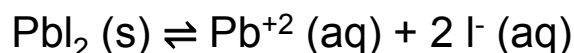
Hebden – Unit 3 (page 73-108)

1. Common Ion Effect

Let's dissolve PbI_2 in a solution that already has sodium iodide, NaI , dissolved in it. How much of PbI_2 will dissolve?

- Will dissolving PbI_2 in a solution full of Na^+ (aq) and I^- (aq) affect the solubility of PbI_2 ? **Yes!!**

Solubility of PbI_2 is controlled by the equilibrium that has a K_{sp} value equals to 8.9×10^{-9} . (from solubility constant table)



In the presence of NaI , the solution now has an extra source of I^- (aq). This is equivalent to stressing the equilibrium by increasing the concentration of I^- (aq). Le Chatelier's principle predicts that it will shift the equilibrium to the LEFT, causing LESS PbI_2 to dissolve.

Solubility Equilibrium

Hebden – Unit 3 (page 73-108)

1. Common Ion Effect

Calculate the molar solubility of PbI_2 in 0.10 M NaI solution.

Recall K_{sp} of PbI_2 at 25°C ? $K_{\text{sp}} = 8.5 \times 10^{-9}$

	$\text{PbI}_2 (\text{s}) \rightleftharpoons \text{Pb}^{+2} (\text{aq}) + 2 \text{I}^- (\text{aq})$		
[I]	-	0	0.10
[C]		+x	+2x
[E]		x	0.10 + 2x

I^- (aq) from
The 0.10 M NaI .

Stoichio-
metric
ratio

$$K_{\text{sp}} = [\text{Pb}^{+2}] [\text{I}^-]^2 = (x)(0.10 + 2x)^2 = 8.5 \times 10^{-9}$$

$$\text{Molar solubility} = x = 8.5 \times 10^{-7} \text{ M}$$

Much smaller!!
Recall that the molar solubility of PbI_2 in pure water is $1.3 \times 10^{-3} \text{ M}$!!

Solubility Equilibrium

Hebden – Unit 3 (page 73-108)

1. Common Ion Effect

Let's dissolve PbI_2 in a solution that already has sodium iodide, NaI , dissolved in it. How much of PbI_2 will dissolve?

In summary, dissolving PbI_2 in a solution of NaI will cause **LESS PbI_2 to dissolve** because of **common ion effect**.

1. Qualitatively, this is as predicted by Le Chatelier's Principle. The extra I^- (aq) from NaI will cause the solubility equilibrium of PbI_2 to shift to the left, forming more PbI_2 (s), or suppressing the dissolution of PbI_2 .
2. Quantitatively, this can be calculated using the solubility product constant of PbI_2 .
 - Molar solubility of PbI_2 in pure water is $1.3 \times 10^{-3} \text{ M}$.
 - Molar solubility of PbI_2 in a 0.10 M NaI solution is $8.5 \times 10^{-7} \text{ M}$.

CHEM 0012 Lecture Notes

49

Solubility Equilibrium

Hebden – Unit 3 (page 73-108)

2. We can use K_{sp} to determine whether a precipitate will form in solution.

This is done by computing Q , the ion product, and compare it to the K_{sp} .

- Precipitate will form if $Q > K_{\text{sp}}$
- No precipitate will form if:
 - $Q = K_{\text{sp}}$ (at saturation point)
 - $Q < K_{\text{sp}}$ (an unsaturated solution)

CHEM 0012 Lecture Notes

50

Solubility Equilibrium

Hebden – Unit 3 (page 73-108)

2. We can use K_{sp} to determine whether a precipitate will form in solution.

Suppose we wish to prepare 0.500 L of a solution containing 0.0075 mole of **NaCl** and 0.075 mole of **Pb(NO₃)₂**. Will a precipitate form?

This question draws on the following knowledge:

1. Solubility rules
2. Predict double-replacement reactions
3. Write chemical formulae
4. Balance chemical reaction
5. Write K_{sp} equilibrium constant expression

CHEM 0012 Lecture Notes

51

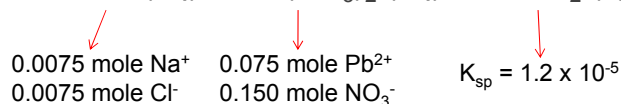
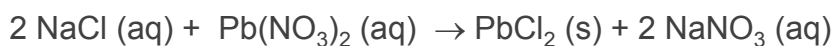
Solubility Equilibrium

Hebden – Unit 3 (page 73-108)

2. We can use K_{sp} to determine whether a precipitate will form in solution.

Suppose we wish to prepare 0.500 L of a solution containing 0.0075 mole of NaCl and 0.075 mole of Pb(NO₃)₂. Will a precipitate form?

Write the balanced equation:



Is there enough Pb²⁺ and Cl⁻ ions in solution to precipitate PbCl₂?

CHEM 0012 Lecture Notes

52

Solubility Equilibrium

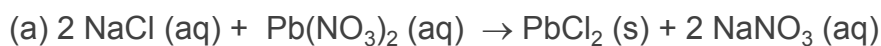
Hebden – Unit 3 (page 73-108)

2. We can use K_{sp} to determine whether a precipitate will form in solution.

Suppose we wish to prepare 0.500 L of a solution containing 0.0075 mole of NaCl and 0.075 mole of $Pb(NO_3)_2$.

(a) Will a precipitate form?

(b) Calculate the concentration of the ions after the reaction.

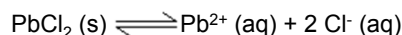


0.0075 mole Na^+
0.0075 mole Cl^-

0.075 mole Pb^{2+}
0.150 mole NO_3^-

$K_{sp} = 1.2 \times 10^{-5}$

Na^+ and NO_3^- are spectator ions.



$[Pb^{2+}] = 0.075 / 0.500 = 0.15 M$
 $[Cl^-] = 0.0075 / 0.500 = 0.015 M$

$Q = [Pb^{2+}] [Cl^-]^2 = (0.15)(0.015)^2 = 3.4 \times 10^{-5}$

$Q > K_{sp}$, precipitate will form!

CHEM 0012 Lecture Notes

53

Solubility Equilibrium

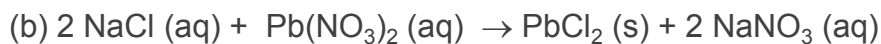
Hebden – Unit 3 (page 73-108)

2. We can use K_{sp} to determine whether a precipitate will form in solution.

Suppose we wish to prepare 0.500 L of a solution containing 0.0075 mole of NaCl and 0.075 mole of $Pb(NO_3)_2$.

(a) Will a precipitate form?

(b) Calculate the concentration of the ions after the reaction.



0.0075 mole NaCl
This is the limiting reagent.

0.075 mole $Pb(NO_3)_2$
Using up
0.00375 mole
 $Pb(NO_3)_2$

Producing
0.00375 mole
 $PbCl_2$

Producing
0.0075 mole
 $NaNO_3$

Left over is

$0.075 - 0.00375 = 0.07125$ mole $Pb(NO_3)_2 \rightarrow 0.1425 M Pb(NO_3)_2 \rightarrow 0.1425 M Pb^{2+}$

Solubility Equilibrium

Hebden – Unit 3 (page 73-108)

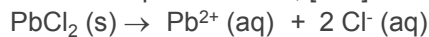
2. We can use K_{sp} to determine whether a precipitate will form in solution.

Suppose we wish to prepare 0.500 L of a solution containing 0.0075 mole of NaCl and 0.075 mole of $Pb(NO_3)_2$.

(a) Will a precipitate form?

(b) Calculate the concentration of the ions after the reaction.

(b) Na^+ and Cl^- are spectator ions, $[Na^+] = 0.015\text{ M}$ and $[NO_3^-] = 0.30\text{ M}$



I	0.1425	0
C	x	2x
E	0.1425 + x	2x

$$K_{sp} = 1.2 \times 10^{-5} = [Pb^{2+}][Cl^-]^2 = (0.1425 + x)(2x)^2 \approx (0.1425)(4x^2)$$

$$x = \sqrt{\frac{1.2 \times 10^{-5}}{(4)(0.1425)}} = 4.6 \times 10^{-3} \quad [Pb^{2+}] = 0.1425 + 0.0046 = 0.147\text{ M}$$

$$[Cl^-] = 2(0.0046) = 0.0092\text{ M}$$

Solubility Equilibrium

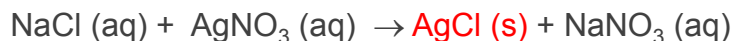
Hebden – Unit 3 (page 73-108)

2. We can use K_{sp} to determine whether a precipitate will form in solution.

What possible precipitate might form by mixing 50.0 mL of $1.0 \times 10^{-4}\text{ M}$ NaCl with 50.0 mL of $1.0 \times 10^{-6}\text{ M}$ $AgNO_3$?

Will a precipitate form?

Yes, there could be a possibility that a precipitate will form because AgCl has low solubility.



Whether AgCl will precipitate out will depend on the concentrations of the Ag^+ and Cl^- ions.

Think dilution when calculating the $[Ag^+]$ and $[Cl^-]$!!

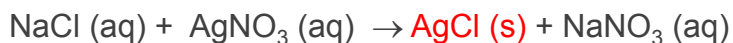
Solubility Equilibrium

Hebden – Unit 3 (page 73-108)

2. We can use K_{sp} to determine whether a precipitate will form in solution.

What possible precipitate might form by mixing 50.0 mL of 1.0×10^{-4} M NaCl with 50.0 mL of 1.0×10^{-6} M AgNO_3 ?

Will a precipitate form?



$$\text{moles of Ag}^+ = (0.0500 \text{ L}) (1.0 \times 10^{-6} \text{ M}) = 5.0 \times 10^{-8} \text{ moles}$$

$$\text{moles of Cl}^- = (0.0500 \text{ L}) (1.0 \times 10^{-4} \text{ M}) = 5.0 \times 10^{-6} \text{ moles}$$

$$[\text{Ag}^+] = [\text{NO}_3^-] = 5.0 \times 10^{-8} \text{ moles} / 0.1000 \text{ L} = 5.0 \times 10^{-7} \text{ M}$$

$$[\text{Na}^+] = [\text{Cl}^-] = 5.0 \times 10^{-6} \text{ moles} / 0.1000 \text{ L} = 5.0 \times 10^{-5} \text{ M}$$

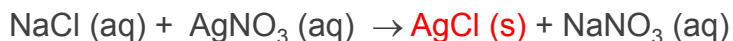
Solubility Equilibrium

Hebden – Unit 3 (page 73-108)

2. We can use K_{sp} to determine whether a precipitate will form in solution.

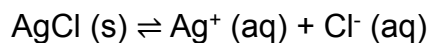
What possible precipitate might form by mixing 50.0 mL of 1.0×10^{-4} M NaCl with 50.0 mL of 1.0×10^{-6} M AgNO_3 ?

Will a precipitate form?



$$[\text{Ag}^+] = 5.0 \times 10^{-8} \text{ moles} / 0.1000 \text{ L} = 5.0 \times 10^{-7} \text{ M}$$

$$[\text{Cl}^-] = 5.0 \times 10^{-6} \text{ moles} / 0.1000 \text{ L} = 5.0 \times 10^{-5} \text{ M}$$



$$K_{sp} = 1.8 \times 10^{-10} = [\text{Ag}^+][\text{Cl}^-]$$

$$Q = (5.0 \times 10^{-7})(5.0 \times 10^{-5}) = 2.5 \times 10^{-11}$$

$Q < K_{sp}$, therefore, no precipitate forms.

Solubility Equilibrium

Hebden – Unit 3 (page 73-108)

Solubility Equilibrium

We have covered the following topics:

1. Common Ion Effect
 - Qualitatively using Le Chatelier's Principle
 - Quantitatively by K_{sp} calculation
2. Use K_{sp} to predict whether a precipitate will form in a solution (i.e. Compare Q with K_{sp})

CHEM 0012 Lecture Notes

59

Solubility Equilibrium

Hebden – Unit 3 (page 73-108)

Solubility Problems

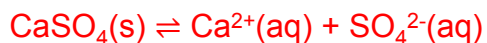
0.24 g CaSO_4 dissolve in 100.0 mL of water at 25°C to form a saturated solution. What is the K_{sp} for CaSO_4 at 25°C .

Solubility Equilibrium

Hebden – Unit 3 (page 73-108)

Solubility Problems

0.24 g CaSO_4 dissolve in 100.0 mL of water at 25°C to form a saturated solution. What is the K_{sp} for CaSO_4 at 25°C .



$$[\text{CaSO}_4] = \frac{(0.24 \text{ g CaSO}_4) \left(\frac{\text{mol CaSO}_4}{136.14 \text{ g CaSO}_4} \right)}{0.1000 \text{ L}} = 1.76 \times 10^{-2} \text{ M} = [\text{Ca}^{2+}] = [\text{SO}_4^{2-}]$$

$$K_{\text{sp}} = [\text{Ca}^{2+}][\text{SO}_4^{2-}] = (1.76 \times 10^{-2})^2 = 3.1 \times 10^{-4}$$

Solubility Equilibrium

Hebden – Unit 3 (page 73-108)

Solubility Problems:

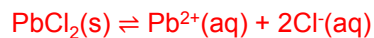
1. Calculate the molar solubility of PbCl_2 in pure water.
2. The molar solubility of Ag_2SO_4 in pure water is $1.2 \times 10^{-5} \text{ M}$. Calculate the K_{sp} .

Solubility Equilibrium

Hebden – Unit 3 (page 73-108)

Solubility Problems:

1. Calculate the molar solubility of PbCl_2 in pure water.



$$K_{sp} = 1.2 \times 10^{-5} = [\text{Pb}^{2+}][\text{Cl}^{-}]^2 = s(2s)^2 = 4s^3$$

$$s = \sqrt[3]{\frac{1.2 \times 10^{-5}}{4}} = 1.4 \times 10^{-2} \text{ M}$$

2. The molar solubility of Ag_2SO_4 in pure water is $1.2 \times 10^{-5} \text{ M}$. Calculate the K_{sp} .



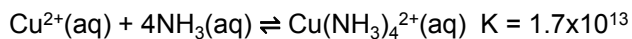
$$K_{sp} = [\text{Ag}^{+}]^2 [\text{SO}_4^{2-}] = [2(1.2 \times 10^{-5})]^2 (1.2 \times 10^{-5}) = 6.9 \times 10^{-15}$$

Solubility Equilibrium

Hebden – Unit 3 (page 73-108)

Solubility Problems:

A 200.0 mL sample of a solution that is $1.5 \times 10^{-3} \text{ M}$ in $\text{Cu}(\text{NO}_3)_2$ is mixed with a 250.0 mL sample of a solution that is 0.20 M in NH_3 . After the solution reaches equilibrium, what concentration of $\text{Cu}^{2+}(\text{aq})$ remains?

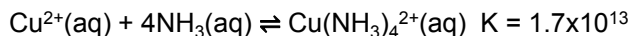


Solubility Equilibrium

Hebden – Unit 3 (page 73-108)

Solubility Problems:

A 200.0 mL sample of a solution that is 1.5×10^{-3} M in $\text{Cu}(\text{NO}_3)_2$ is mixed with a 250.0 mL sample of a solution that is 0.20 M in NH_3 . After the solution reaches equilibrium, what concentration of $\text{Cu}^{2+}(\text{aq})$ remains?



$$[\text{Cu}^{2+}]_0 = \frac{(200.0 \text{ mL})(1.5 \times 10^{-3} \text{ M})}{200.0 \text{ mL} + 250.0 \text{ mL}} = 6.7 \times 10^{-4} \text{ M}$$

$$[\text{NH}_3]_0 = \frac{(250.0 \text{ mL})(0.20 \text{ M})}{200.0 \text{ mL} + 250.0 \text{ mL}} = 0.11 \text{ M}$$

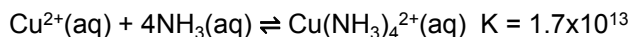
Since $K \gg 1$ let the reaction go to completion and then let it approach equilibrium

Solubility Equilibrium

Hebden – Unit 3 (page 73-108)

Solubility Problems:

A 200.0 mL sample of a solution that is 1.5×10^{-3} M in $\text{Cu}(\text{NO}_3)_2$ is mixed with a 250.0 mL sample of a solution that is 0.20 M in NH_3 . After the solution reaches equilibrium, what concentration of $\text{Cu}^{2+}(\text{aq})$ remains?



$$\text{I} \quad 6.7 \times 10^{-4} \quad 0.11 \quad 0$$

$$\text{R} \quad -6.7 \times 10^{-4} \quad -4(6.7 \times 10^{-4}) \quad 6.7 \times 10^{-4}$$

$$\text{C} \quad x \quad 4x \quad -x$$

$$\text{E} \quad x \quad 0.11 + 4x \quad 6.7 \times 10^{-4} - x$$

$$K = 1.7 \times 10^{13} = \frac{6.7 \times 10^{-4} - x}{(x)(0.11 + 4x)^4} \approx \frac{6.7 \times 10^{-4}}{(x)(0.11)^4} \rightarrow x = 2.7 \times 10^{-13} \text{ M assumption is OK}$$

$$[\text{Cu}^{2+}] = 2.7 \times 10^{-13} \text{ M} \quad [\text{NH}_3] = 0.11 \text{ M} + 4(2.7 \times 10^{-13} \text{ M}) = 0.11 \text{ M}$$

$$[\text{NH}_3] = 6.7 \times 10^{-4} \text{ M} - 2.7 \times 10^{-13} \text{ M} = 6.7 \times 10^{-4} \text{ M}$$

Solubility Equilibrium

Hebden – Unit 3 (page 73-108)

Solubility Problems:

1. What is the molar solubility of CaF_2 in 0.100 M NaF. $K_{\text{sp}} = 1.5 \times 10^{-10}$
2. A 30.0 mL sample of a solution that is 0.0500 M in $\text{Pb}(\text{NO}_3)_2$ is mixed with a 70.0 mL sample of a solution that is 0.00500 M in NaBr. Will a precipitate form?

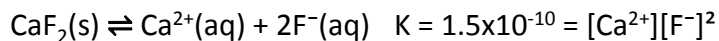
Solubility Equilibrium

Hebden – Unit 3 (page 73-108)

Solubility Problems:

1. What is the molar solubility of CaF_2 in 0.100 M NaF.

$$K_{\text{sp}} = 1.5 \times 10^{-10}$$



$$\begin{array}{lcl} \text{I} & 0 & 0.100 \end{array}$$

$$\begin{array}{lcl} \text{C} & s & 2s \end{array}$$

$$\begin{array}{lcl} \text{E} & s & 0.100+2s \end{array}$$

$$K = 1.5 \times 10^{-10} = s(0.100+2s)^2 \approx s(0.100)^2$$

$$s = 1.5 \times 10^{-8} \text{ M} \quad \text{assumption is OK}$$

Solubility Equilibrium

Hebden – Unit 3 (page 73-108)

Solubility Problems:

2. A 30.0 mL sample of a solution that is 0.0500 M in $\text{Pb}(\text{NO}_3)_2$ is mixed with a 70.0 mL sample of a solution that is 0.00500 M in NaBr. Will a precipitate form?

Must consider $\text{PbBr}_2(\text{s}) \rightleftharpoons \text{Pb}^{2+}(\text{aq}) + 2\text{Br}^{-}(\text{aq})$

$$[\text{Pb}^{2+}]_0 = \frac{(30.0 \text{ mL})(0.0500 \text{ M})}{100.0 \text{ mL}} = 0.0150 \text{ M}$$

$$[\text{Br}^{-}]_0 = \frac{(70.0 \text{ mL})(0.00500 \text{ M})}{100.0 \text{ mL}} = 0.00350 \text{ M}$$

$$Q = [\text{Pb}^{2+}][\text{Br}^{-}]^2 = (0.0150)(0.00350)^2 = 1.8 \times 10^{-7} < K_{\text{sp}} = 6.6 \times 10^{-6}$$

Since $Q < K_{\text{sp}}$ no precipitate forms

Solubility Equilibrium

Hebden – Unit 3 (page 73-108)

Solubility Problems

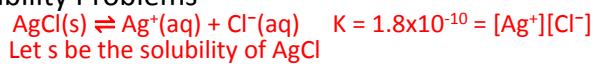
What is the solubility of AgCl in:

- a) Pure water
- b) $4.00 \times 10^{-3} \text{ M CaCl}_2$
- c) $4.00 \times 10^{-3} \text{ M Ca}(\text{NO}_3)_2$
- d) $4.00 \times 10^{-3} \text{ M CaBr}_2$

Solubility Equilibrium

Hebden – Unit 3 (page 73-108)

Solubility Problems



What is the solubility of AgCl in:

- a) Pure water

$$1.8 \times 10^{-10} = (s)(s) \rightarrow s = 1.3 \times 10^{-5} \text{ M}$$

- b) $4.00 \times 10^{-3} \text{ M CaCl}_2$

$$1.8 \times 10^{-10} = (s)(8.00 \times 10^{-3} + s) \approx (s)(8.00 \times 10^{-3}) \rightarrow s = 2.3 \times 10^{-8} \text{ M}$$

- c) $4.00 \times 10^{-3} \text{ M Ca(NO}_3)_2$ same as pure water

$$1.8 \times 10^{-10} = (s)(s) \rightarrow s = 1.3 \times 10^{-5} \text{ M}$$

- d) $4.00 \times 10^{-3} \text{ M CaBr}_2$

$$1.8 \times 10^{-10} = (s - 8.00 \times 10^{-3})(s) \text{ let } s = x + 8.00 \times 10^{-3}$$

$$1.8 \times 10^{-10} = (x)(x + 8.00 \times 10^{-3}) \approx (x)(8.00 \times 10^{-3}) \rightarrow x = 2.3 \times 10^{-8} \text{ M} \rightarrow s = 8.00 \times 10^{-3} \text{ M}$$