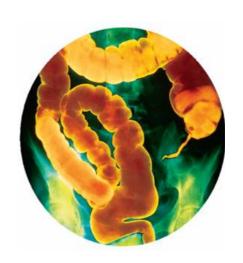
18.3 Solubility Equilibrium

18.3 Solubility Equilibrium

Connecting to Your World

Barium sulfate is ingested by a patient before X-ray images of the digestive tract are taken. Barium sulfate absorbs the X-rays, thereby producing light areas on the developed X-ray film. However, barium salts are usually toxic. You will learn why patients can ingest this poisonous substance without harm.



What is a solution?

- A solution is a mixture in which a solid has been dissolved into a liquid, usually water
- □ It is expressed as (aq) for its state of matter
- lonic compounds are solids at room temperature
- In order for the ions to interact with one another, they need to be dissolved in water, forming a solution

Solubility

- Dissolving and precipitating
 - Ex. Tooth decay from acids
 - Precipitation of salts in the kidneys which forms stones
 - Dissolved salts in rainwater
- A saturated solution is one where no more solute can dissolve in solution
- Solubility equations look like net ionic equations
 - \blacksquare BaSO_{4(s)} \longleftrightarrow Ba²⁺_(aq) + SO₄²⁻_(aq)

Equilibrium Expressions and Solubility

- Equilibrium expresses the degree to which the solid is soluble in water
- \square BaSO_{4(s)} \longleftrightarrow Ba²⁺_(aq) + SO₄²⁻_(aq)
 - This equation indicates that at equilibrium there is a presence of all three entities
 - However, we are only concerned with the concentration of ions in the solution, although we accept that there is some undissolved ionic compound present (BaSO_{4(s)})

□ The equilibrium reaction would be as follows:

Ionic Compound(s)
$$\leftrightarrow$$
 Cation⁺(aq) + Anion⁻(aq)

$$AgCl(s) \leftrightarrow Ag^{1+}_{(aq)} + Cl^{1-}_{(aq)}$$

$$AgCl(s) \Longrightarrow Ag^{+}(aq) + Cl^{-}(aq)$$

You can write an equilibrium expression for this process.

$$K_{\text{eq}} = \frac{[\text{Ag}^+] \times [\text{Cl}^-]}{[\text{AgCl}]}$$

$$K_{\rm eq} \times [{\rm AgCl}] = [{\rm Ag^+}] \times [{\rm Cl^-}] = K_{\rm sp}$$

- e.g. Write the Ksp expression for:
 - (i) AgCl
- (ii) BaSO₄
- (iii) PbCl₂
- (i) Ionization Equation: $AgCI_{(s)} \longrightarrow Ag^+_{(aq)} + CI^-_{(aq)}$

$$K_{sp} = [Ag^{+}_{(aq)}] [Cl^{-}_{(aq)}]$$

(ii) Ionization Equation: $BaSO_{4(s)} \longleftrightarrow Ba^{2+}_{(aq)} + SO_4^{2-}_{(aq)}$

$$K_{sp} = [Ba^{2+}_{(aq)}][SO_4^{2-}_{(aq)}]$$

(iii) Ionization Equation: $PbCl_{2(s)} \rightleftharpoons Pb^{2+}_{(aq)} + 2 Cl_{(aq)}$

$$K_{sp} = [Pb^{2+}_{(aq)}] [Cl_{(aq)}]^{2}$$

The Solubility Product Constant

- Solubility varies from one ionic compound to the next
- Some ionic compounds can dissolve quite significantly in water where others cannot
 - They are either soluble or insoluble
- □ Table 18.1, p. 561 highlights some of the <u>solubilities</u> of various ionic compounds
 - Water is always the solvent

Table 18.1

Solubilities of Ionic Compounds in Water					
Compounds	Solubility	Exceptions			
Salts of Group 1A metals and ammonia	Soluble	Some lithium compounds			
Ethanoates, nitrates, chlorates, and perchlorates	Soluble	Few exceptions			
Sulfates	Soluble	Compounds of Pb, Ag, Hg, Ba, Sr, and Ca			
Chlorides, bromides, and iodides	Soluble	Compounds of Ag and some compounds of Hg and Pb			
Sulfides and hydroxides	Most are insoluble	Alkali metal sulfides and hydroxides are soluble. Compounds of Ba, Sr, and Ca are slightly soluble.			
Carbonates, phosphates, and sulfites	Insoluble	Compounds of the alkali metals and of ammonium ions			

- Although an ionic compound may be considered insoluble, a small amount does actually dissolve
- □ The equilibrium reaction would be as follows:

Ionic Compound(s)
$$\leftrightarrow$$
 Cation⁺(aq) + Anion⁻(aq)

$$AgCI(s) \leftrightarrow Ag^{1+}_{(aq)} + CI^{1-}_{(aq)}$$

The **solubility product constant** (K_{sp}) , equals the product of the concentrations of the ions, each raised to a power equal to the coefficient of the ion in the dissociation equation.

$$K_{\rm sp} = [{\rm Ag}^+] \times [{\rm Cl}^-]$$

 $K_{\rm sp} = 1.8 \times 10^{-10}$



The smaller the numerical value of the solubility product constant, the lower the solubility of the compound.

Scale, formed by the precipitation of slightly soluble salts, builds up around faucets.



Why are insoluble substances hard to remove?

They do NOT dissolve in a solvent such as water.

Review

Any ionic solid placed in water establishes an equilibrium between its dissociated ions in solution and the solid that is undissolved

The solubility product constant is an equilibrium constant that describes the equilibrium between a solid and its ions in solution

Table 18.2, p. 562 – examine for high/low solubilities

Table 18.2							
Solubility Product Constants (K _{sp}) at 25°C							
Salt	K _{sp}	Salt	K _{sp}	Salt	K ₅p		
Halides		Sulfates		Hydroxides			
AgCl	1.8×10^{-10}	PbSO₄	6.3×10^{-7}	AI(OH) ₃	3.0×10^{-34}		
AgBr	5.0×10^{-13}	BaSO ₄	1.1×10^{-10}	Zn(OH) ₂	3.0×10^{-16}		
Agl	8.3×10^{-17}	CaSO ₄	$2.4 imes 10^{-5}$	Ca(OH) ₂	$6.5 imes 10^{-6}$		
PbCl₂	1.7×10^{-5}	Sulfides		Mg(OH)₂	7.1×10^{-12}		
PbBr₂	2.1×10^{-6}	NiS	4.0×10^{-20}	Fe(OH) ₂	7.9×10^{-16}		
Pbl₂	7.9×10^{-9}	CuS	8.0×10^{-37}	Carbonates			
PbF ₂	$3.6 imes 10^{-8}$	Ag₂S	8.0×10^{-51}	CaCO ₃	4.5×10^{-9}		
CaF ₂	3.9×10^{-11}	ZnS	3.0×10^{-23}	SrCO₃	9.3×10^{-10}		
Chromates		FeS	8.0×10^{-19}	ZnCO ₃	1.0×10^{-10}		
PbCrO₄	1.8×10^{-14}	CdS	1.0×10^{-27}	Ag ₂ CO ₃	8.1×10^{-12}		
Ag₂CrO₄	1.2×10^{-12}	PbS	3.0×10^{-28}	BaCO ₃	$5.0 imes 10^{-9}$		

SAMPLE PROBLEM 18.3

Finding the Ion Concentrations in a Saturated Solution

What is the concentration of lead ions and chromate ions in a saturated lead chromate solution at 25°C? ($K_{\rm sp}=1.8\times10^{-14}$)

Analyze List the knowns and the unknowns.

Knowns

$$\bullet K_{\rm sp} = 1.8 \times 10^{-14}$$

$$\bullet K_{\rm sp} = [{\rm Pb^{2+}}] \times [{\rm CrO_4}^{2-}]$$

• PbCrO₄(s)
$$\rightleftharpoons$$
 Pb²⁺(aq) + CrO₄²⁻(aq)

Unknowns

•
$$[Pb^{2+}] = ? M$$

•
$$[CrO_4^{2-}] = ? M$$

For each Pb²⁺ ion formed, one CrO₄²⁻ ion is formed.

Calculate Solve for the unknowns.

$$K_{\rm sp} = [{\rm Pb^{2+}}] \times [{\rm CrO_4^{2-}}] = 1.8 \times 10^{-14}$$

At equilibrium $[Pb^{2+}] = [CrO_4^{2-}]$. Substitute $[Pb^{2+}]$ for $[CrO_4^{2-}]$ in the expression for $K_{\rm sp}$ to get an equation with one unknown.

$$K_{\rm sp} = [{\rm Pb^{2+}}] \times [{\rm Pb^{2+}}] = [{\rm Pb^{2+}}]^2 = 1.8 \times 10^{-14}$$

Solve for [Pb²⁺].

$$[Pb^{2+}] = [CrO_4^{2-}] = \sqrt{1.8 \times 10^{-14}} = 1.3 \times 10^{-7} M$$

Finding the Ion Concentrations in a Saturated Solution

What is the concentration of lead ions and chromate ions in a saturated lead chromate solution at 25°C? ($K_{\rm sp}=1.8\times10^{-14}$)

1)
$$PbCrO_{4(s)} = Pb_{(aq)}^{2+} + CrO_{4(aq)}^{2-}$$
2) $K_{sp} = 1.8 \times 10^{-14}$
3) $1.8 \times 10^{-14} = [Pb_{(aq)}^{2+}][CrO_{4(aq)}^{2-}]$

$$= X \cdot X$$

$$[Pb_{(aq)}^{2+}] = [CrO_{4(aq)}^{2-}] = 1.3 \times 10^{-7} M$$

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Extra question

What is the concentation of barium and sulfate ions in a saturated barium sulfate solution at 25°C?

$$K_{sp} = 1.1 \times 10^{-10}$$

Example

(1)
$$PbCrO_{4(s)} = Pb_{(aq)}^{2+} + CrO_{4(aq)}^{2-}$$

(2) $K_{sp} = 1.8 \times 10^{-14}$

(3) $1.8 \times 10^{-14} = [Pb_{(aq)}^{2+}][CrO_{4(aq)}^{2-}]$

$$= X \cdot X$$

$$1.8 \times 10^{-14} = X^{2}$$

$$1.3 \times 10^{-7} = X$$

17. Lead(II) sulfide (PbS) has a $K_{\rm sp}$ of 3.0×10^{-28} . What is the concentration of lead(II) ions in a saturated solution of PbS?

$$PbS_{(s)} = Pba_{(s)} + Sa_{(s)}$$

$$Ksp = 3.0 \times 10^{-28}$$

$$Ksp = [3.0 \times 10^{-28}]$$

$$X = X$$

18. What is the concentration of calcium ions in a saturated calcium carbonate solution at 25°C? ($K_{\text{sp}} = 4.5 \times 10^{-9}$)

$$(a(0_3/3)) = (a(0_3)) + (\overline{0})$$
 $(a(0_3/3)) = (a(0_3)) + (\overline{0})$
 $(a(0)) = (a(0_3)) + (\overline{0})$
 $(a(0)) = (a(0)) + (a(0))$
 $(a(0)) = (a(0)) + (a(0))$
 $(a(0)) = (a(0)) + (a(0))$
 $(a$

What is the concentation of barium and sulfate ions in a saturated barium sulfate solution at 25° C?

$$K_{sp} = 1.1 \times 10^{-10}$$

$$1.10 \times 10^{-10} = x^2$$

 $Ba^{2+} SO_4^{2-} X = 1.0 \times 10^{-5}$

What is the concentation of barium and sulfate ions in a saturated barium sulfate solution at 25° C?

$$K_{sp} = 1.1 \times 10^{-10}$$

$$B_{0}S_{0}H_{15}$$
 $B_{0}(0_{1})$ $S_{0}H_{19}$
 $K_{S}p = [B_{0}^{2+}][S_{0}H_{19}]$
 $V_{1}/X_{10} = X_{0}^{2+}[S_{0}H_{19}]$
 $V_{2}/X_{10} = X_{0}^{2+}[S_{0}H_{19}]$

The Common Ion Effect

- When an ionic compound has reached saturation, another ionic compound can be added to increase the concentration of one of the ions
- Le Chatelier's principle says that when a stress is applied to an equilibrium, it re-establishes itself by adjusting its equilibrium position
- The reaction will shift to the left, resulting in more reactant (ionic compound(s)) forming

Common Ion Effect - continued...

As long as another compound, containing either a cation or anion from the original compound, is being added, the equilibrium position will adjust itself so that Ksp does not change

Original –
$$K_{sp} = [Pb^{2+}][CrO_4^{2-}] = 1.8 \times 10^{-14}$$

Adding Pb(NO₃)₂

Revised
$$- K_{5p} = [Pb^{2+}][CrO_4^{2-}] = 1.8 \times 10^{-14}$$

In this example, the lead ion is called a common ion. A **common ion** is an ion that is found in both salts in a solution. Adding lead nitrate to a saturated solution of PbCrO₄ causes the solubility of PbCrO₄ to decrease. The lowering of the solubility of an ionic compound as a result of the addition of a common ion is called the **common ion effect.** Adding sodium chromate to the solution of PbCrO₄ would also produce the common ion effect. The additional chromate ion, a different common ion, would similarly cause the solubility equilibrium to shift to the left and produce more PbCrO₄.

Common ion and common ion effect

Example: adding lead (II) nitrate to an existing saturated solution of lead (II) chromate



A saturated solution of lead(II) chromate is pale yellow.



When a few drops of lead nitrate are added to the solution, more lead(II) chromate precipitates.

FARSON

Definitions

- A common ion is the cation or anion that is found in both ionic compounds (salts) in a solution
- Common ion effect is the lowering of the solubility of an ionic compound as a result of the addition of a common ion
- Spectator ions are those that do not participate in the reaction. They stay in solution and do not precipitate

Precipitates

- Ksp can be used to determine whether a precipitate will form when solutions are mixed
- in the mixture is greater than the Ksp of the compound formed from the ions, a precipitate will form

Qsp - Solubility product quotient

Precipitation

- If we compare the reaction quotient, Q, for the current solution concentrations to the value of K_{sp}, we can determine whether precipitation will occur.
 - $Q = K_{sp}$: the solution is saturated; no precipitation
 - $Q < K_{sp}$: the solution is unsaturated; no precipitation
 - Q > K_{sp}: the solution is above saturation; the salt above saturation will precipitate

SAMPLE PROBLEM 18.4

Finding Equilibrium Ion Concentrations in the Presence of a Common Ion

Photographic film is covered with a light-sensitive emulsion containing silver bromide. The $K_{\rm sp}$ of silver bromide is 5.0×10^{-13} . What is the bromide-ion concentration of a 1.00-L saturated solution of AgBr to which 0.020 mol of AgNO₃ is added?

Analyze List the knowns and the unknown.

Knowns

• $K_{\rm sp} = 5.0 \times 10^{-13}$

Unknown

 \bullet [Br⁻] = ? M

- moles of AgNO₃ added = 0.020 mol
- volume of solution = 1 L
- $AgBr(s) \Longrightarrow Ag^{+}(aq) + Br^{-}(aq)$
- $K_{\rm sp} = [{\rm Ag^+}] \times [{\rm Br^-}]$

Express the concentrations of the two ions in one unknown. Let the equilibrium concentration of bromide ion from the dissociation be x. Then the equilibrium concentration of silver ion is x + 0.020.

Calculate Solve for the unknown.

Because of the small value of $K_{\rm sp}$, you can make a simplifying assumption: x will be negligibly small compared to 0.020. Thus the [Ag⁺] at equilibrium is approximately equal to 0.020M. Solve for x and substitute these values into the $K_{\rm sp}$ expression.

Ho the
$$K_{sp}$$
 expression.

$$K_{sp} = [Ag^{+}] \times [Br^{-}] = [Ag^{+}] \times x$$

$$x = \frac{K_{sp}}{[Ag^{+}]} = \frac{(5.0 \times 10^{-13})}{[Ag^{+}]}$$

$$= \frac{(5.0 \times 10^{-13})}{0.020}$$

$$= [Br^{-}] = 2.5 \times 10^{-11} M$$

The equilibrium concentration of bromide ion is $2.5 \times 10^{-11} M$.

19. What is the concentration of sulfide ion in a 1.0-L solution of iron(II) sulfide to which 0.04 mol of iron(II) nitrate is added? The $K_{\rm sp}$ of FeS is 8×10^{-19} .

Questions 19, 20, p. 564

Questions P 565 21-27

Ksp FeS=
$$8 \times 10^{-19}$$

mols of FeNO₃ added = 0.04 mol
volume = $1L$

$$FeS \longrightarrow Fe^{2+}_{(aq)} + S^{2-}_{(aq)}$$

$$Ksp = [Fe^{2+}][S^{2-}]$$

$$\frac{||S_{1}||^{2}}{||S_{1}||^{2}} = \frac{||S_{1}||^{2}}{||S_{1}||^{2}} = \frac{||S_{1}||^{2}}{||S_{1}||^{2}} = \frac{||S_{1}||^{2}}{||S_{1}||^{2}}$$

$$2 \times 10^{-17} = 5^{2-1}$$

Q 20-25

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Questions P 565 21-25

20. The $K_{\rm sp}$ of SrSO₄ is 3.2×10^{-7} . What is the equilibrium concentration of sulfate ion in a 1.0-L solution of strontium sulfate to which 0.10 mol of Sr(CH₃CO₂)₂ has been added?

0.10mol Sr (CH & a)2

$$SrS0_{4} = Sr_{10} = So_{10} = 3.2 \times 10^{-6}$$

$$SrS0_{10} = 3.2 \times 10^{-6}$$

23. Write the solubility product expression for

Ag₂CO₃.

$$Ag_2CO_3 = 2 + 1$$

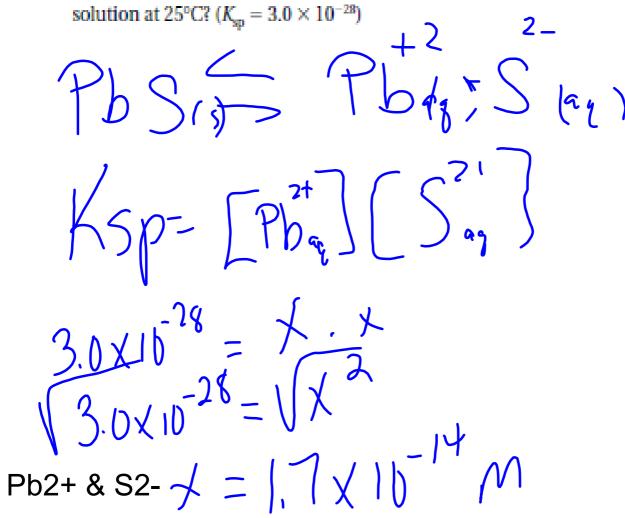
$$Ag_2CO_3 = 2 + 1$$

$$Ag_2CO_3 = 2 + 1$$

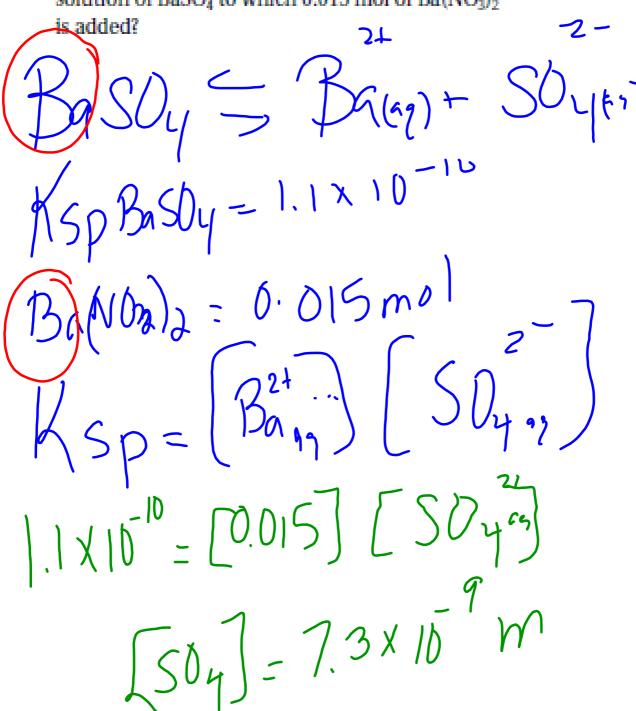
$$CO_3 = 2 + 1$$

$$Ksp = [Ag+]^2 = CO_3^2$$

24. What is the concentration of lead ions and sulfide ions in a saturated solution of lead sulfide (PbS) solution at 25°C? (K = 3.0 × 10⁻²⁸)



25. The K_{sp} of barium sulfate is 1.1 × 10⁻¹⁰. What is the sulfate-ion concentration of a 1.00 L saturated solution of BaSO₄ to which 0.015 mol of Ba(NO₃)₂



Suppose you have $0.50 \, \mathrm{L}$ of $0.002 M \, \mathrm{Ba(NO_3)_2}$ and mix it with $0.50 \, \mathrm{L}$ of $0.008 \, \mathrm{M} \, \mathrm{Na_2SO_4}$. You then have one liter of solution. The insoluble compound that could precipitate is barium sulfate. To predict whether a precipitate of $\mathrm{BaSO_4}$ will form, you need to determine the concentration of the ions after the dilution that occurs on mixing. Precipitation will occur if the product of the concentrations of the two ions ($\mathrm{Ba^{2+}}$ and $\mathrm{SO_4^{2-}}$) exceeds the K_{sp} of $\mathrm{BaSO_4}$, which is 1.1×10^{-10} . Because each solution was diluted with an equal volume of the other solution, the concentrations of both $\mathrm{Ba^{2+}}$ and $\mathrm{SO_4^{2-}}$ after mixing will be half of their original concentrations. Thus in the combined solution, [$\mathrm{Ba^{2+}}$] = 0.001 M and [$\mathrm{SO_4^{2-}}$] = 0.004 M. These concentrations can now be multiplied together as a trial product and the result compared with the K_{max} .

Step 1 - write double displacement equation

$$Ba(NO_3)_2 + Na_2SO_4$$
.

Step 2 - determine M of dilutions

Step 3 - Write out the Qsp expression for the salt that formed as the two ions it formed from.

Step 4 - Compare the Qsp of the salt to the Ksp of the salt to see if a precipitate will form.

$$Qsp = K_{sp} \text{ of BaSO}_4, \text{ which is } 1.1 \times 10^{-10}.$$

Suppose you have 0.50 L of $0.002M \text{ Ba}(\text{NO}_3)_2$ and mix it with 0.50 L of $0.008 \text{ M} \text{ Na}_2\text{SO}_4$. You then have one liter of solution. The insoluble compound that could precipitate is barium sulfate. To predict whether a precipitate of BaSO_4 will form, you need to determine the concentration of the ions after the dilution that occurs on mixing. Precipitation will occur if the product of the concentrations of the two ions $(\text{Ba}^{2+} \text{ and SO}_4^{2-})$ exceeds the K_{sp} of BaSO_4 , which is 1.1×10^{-10} . Because each solution was diluted with an equal volume of the other solution, the concentrations of both Ba^{2+} and SO_4^{2-} after mixing will be half of their original concentrations. Thus in the combined solution, $[\text{Ba}^{2+}] = 0.001M$ and $[\text{SO}_4^{2-}] = 0.004M$. These concentrations can now be multiplied together as a trial product and the result compared with the K

Step 1 - write double displacement equation

$$Ba(NO_3)_2 + Na_2SO_4$$
 BaSO₄ + 2 NaNO₃

Step 2 - determine M of dilutions

$$Ba(NO_3)_2$$
 0.002 M x 0.50 L =0.001 M
Add both solutions together $(0.50 + 0.50)L$
 Na_2SO_4 0.008 M x 0.50 L =0.004 M
 $(0.50 + 0.50)L$

Step 3 - Write out the Qsp expression for the salt that formed as the two ions it formed from.

Qsp =
$$[Ba^{2+}] \times [SO_4^{2-}]$$
 = [0.001] [0.004]
= 4.0 x 10⁻⁶

Step 4 - Compare the Qsp of the salt to the Ksp of the salt to see if a precipitate will form.

Qsp =
$$4.0 \times 10^{-6}$$
 K_{sp} of BaSO₄, which is 1.1×10^{-10} . Yes

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Questions 26-27

Question 26 look up Ksp of AgCl on table 18.2 page 562

Practice Problems 18.3

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26. Would precipitation occur when 500 mL of a 0.02M solution of AgNO₃ is mixed with 500 mL of a 0.001M solution of NaCl? Explain.

27. Which compound, FeS ($K_{\rm sp}=8.0\times10^{-19}$) or CuS ($K_{\rm sp}=8.0\times10^{-37}$), has the higher solubility?

FeS

Section 18.3 Assessment

- **21.** the smaller the $K_{\rm Sp}$, the lower the solubility of the compound
- **22.** If the product of the concentrations of two ions in a mixture is greater than the $K_{\rm sp}$ of the compound formed from the ions, a precipitate will form.
- **23.** $[Ag^+]^2 \times [CO_3^{2-}] = K_{sp}$

- **24.** $2 \times 10^{-14} M$ **25.** $7.3 \times 10^{-9} M$
- **26.** yes; the product $[Ag^+] \times [CI^-] = 0.01M \times 0.0005M = 5 \times 10^{-6} >> K_{sp} AgCI = 1.8 \times 10^{-10}$
- 27. FeS
- **28.** 4×10^{-20}

Will a precipitate form when 0.500L of 0.002M Ba(NO₃)₂ reacts with 0.500L of 0.008M Na₂SO₄? [K_{sp} (BaSO₄) = 1.1 x 10⁻¹⁰]

knowns

$$Ba(NO_3)_2 K_{sp} = 1.1 \times 10^{-10}$$

initial moles of $Ba(NO_3)_2 = 0.002 M$

initial moles of $Na_2SO_4 = 0.008 M$

volume of solution = 0.50L + 0.50L = 1.0 L

To predict if a precipitate will form the concentration after dilution needs to be calculated, if it exceeds the Ksp of $Ba(NO_3)_2$, a precipitate will form

Each solution was diluted with an equal volume of the other solution, so the concentration of each with be half after mixing:

moles of Ba(NO_3)₂ = 0.001 M

moles of $Na_2SO_4 = 0.004 M$

$$[Ba^{2+}] \times [SO_4^{2-}] = (0.001M) \times (0.004M) = 4 \times 10^{-6}$$

$$Ba(NO_3)_2 K_{sp} = 1.1 \times 10^{-10}$$

The trial product is larger therefore a precipitate will form , this will continue until the product of the concentration of the ions remaining in solution equals 1.1×10^{-10}

Finding Equilibrium Ion Concentration Using a Common Ion

Work through Sample Problem 18.4, p. 564.

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Section 18.3 Review

18.3 Practice problems

SECTION 18.3 SOLUBILITY EQUILIBRIUM

- Write the solubility product expression for a. Ca(OH)₂ and b. Ag₂CO₃.
- 2. What is the concentration of silver ions in a saturated solution of silver carbonate? The $K_{\rm sp}$ of Ag₂CO₃ is 8.1×10^{-12} .
- 3. The equilibrium concentration of hydroxide ions in a saturated solution of iron(II) hydroxide is $1.2 \times 10^{-5} M$ at a certain temperature. Calculate the $K_{\rm sp}$ of Fe(OH) $_2$ at this temperature.
- 4. Strontium carbonate has a $K_{\rm sp}=9.3\times10^{-10}$ at 25°C. What is the concentration of strontium ions in a saturated solution of SrCO₃?
- 5. What is the equilibrium concentration of silver ions at 25°C in a 1.0-L saturated solution of silver carbonate to which 0.20 mol of Na₂CO₃ has been added? The $K_{\rm sp}$ of Ag₂CO₃ is 8.1 \times 10⁻¹² at 25°C.
- 6. Will a precipitate of PbSO₄ form when 400.0 mL of 0.0050*M* MgSO₄ is mixed with 600.0 mL of 0.0020*M* Pb(NO₃)₂? The K_{SD} of PbSO₄ = 6.3×10^{-7} .
- 7. Will precipitation of CaCO $_3$ occur when 500.0 mL of 4.2 \times 10 ^{-3}M CaCl $_2$ is mixed with 500.0 mL of 2.6 \times 10 ^{-3}M Na $_2$ CO $_3$? The $K_{\rm sp}$ of CaCO $_3$ is 4.5 \times 10 $^{-9}$.
- 8. Which of these compounds would not decrease the solubility of $Mg(OH)_2$ when added to a saturated solution of the compound?

NaOH, MgCl₂, NaCl, KOH

Section 18.3

1. **a.**
$$Ca(OH)_2(s) \rightleftharpoons Ca^{2+}(aq) + 2OH^-(aq)$$

 $K_{sp} = [Ca^{2+}] \times [OH^-]^2$
b. $Ag_2CO_3(s) \rightleftharpoons 2Ag^+(aq) + CO_3^{2-}(aq)$
 $K_{sp} = [Ag^+]^2 \times [CO_3^{2-}]$
2. $Ag_2CO_3(s) \rightleftharpoons 2Ag^+(aq) + CO_3^{2-}(aq)$
 $K_{sp} = [Ag^+]^2 \times [CO_3^{2-}]$
 $Let x = [CO_3^{2-}]; 2x = Ag^+$
 $K_{sp} = (2x)^2(x) = 8.1 \times 10^{-12}$
 $4x^3 = 8.1 \times 10^{-12}$
 $x^3 = 2.0 \times 10^{-12}$
 $x = 1.3 \times 10^{-4}M = [CO_3^{2-}]$
 $[Ag^+] = 2x = 2.6 \times 10^{-4}M$

3.
$$Fe(OH)_2(s) \rightleftharpoons Fe^{2+}(aq) + 2OH^-(aq)$$

 $K_{sp} = [Fe^{2+}] \times [OH^-]^2$
 $[Fe^{2+}] = 0.5[OH^-] = 6.0 \times 10^{-6}$
 $K_{sp} = (6.0 \times 10^{-6})(1.2 \times 10^{-5})^2$
 $K_{sp} = 8.6 \times 10^{-16}$
4. $SrCO_3 \rightleftharpoons Sr^{2+}(aq) + CO_3^{2-}(aq)$
 $K_{sp} = [Sr^{2+}] \times [CO_3^{2-}]$
 $Let x = [Sr^{2+}] = [CO_3^{2-}]$
 $K_{sp} = x^2 = 9.3 \times 10^{-10}$
 $x = [Sr^{2+}] = 3.0 \times 10^{-5}M$

5.
$$K_{sp} = [Ag^+]^2 \times [CO_3^{2-}]$$

Let $[Ag^+] = 2x$; $[CO_3^{2-}] = x + 0.20$
assume $x << 0.20$ mol; $[CO_3^{2-}] = 0.20$
 $K_{sp} = (2x)^2(0.20) = 8.1 \times 10^{-12}$
 $x^2 = 1.0 \times 10^{-11}$
 $x = 3.2 \times 10^{-6}M = [CO_3^{2-}]$
 $[Ag^+] = 6.4 \times 10^{-6}M$

6.
$$K_{\rm sp}({\rm PbSO_4}) = 6.3 \times 10^{-7} = [{\rm Pb^{2+}}] \times [{\rm SO_4}^{2-}]$$

 $[{\rm SO_4}^{2-}] = 0.0050M = \frac{400.0 \text{ mL}}{1000.0 \text{ mL}} = 0.0020M$
 $[{\rm Pb^{2+}}] = 0.0020M = \frac{600.0 \text{ mL}}{1000.0 \text{ mL}} = 0.0012M$
 $[{\rm Pb^{2+}}] \times [{\rm SO_4}^{2-}] = (0.0012)(0.0020)$
 $= 2.4 \times 10^{-6}$

Because this product exceeds the $K_{\rm sp}$ value, precipitation will occur.

7.
$$K_{\rm sp}({\rm CaCO_3}) = 4.5 \times 10^{-9} = [{\rm Ca^{2+}}] \times [{\rm CO_3}^{2-}]$$

The total volume is 1000 mL, so $[{\rm Ca^{2+}}] = 0.0021 \, {\rm mol/L} = 0.0021 M$ $[{\rm CO_3}^{2-}] = 0.0013 \, {\rm mol/L} = 0.0013 M$ $[{\rm Ca^{2+}}] \times [{\rm CO_3}^{2-}] = (0.0021)(0.0013)$ $= 2.7 \times 10^{-6}$

Because this product exceeds the $K_{\rm sp}$ value, precipitation will occur.

8. NaCl has no ion in common with Mg(OH)2.

Example

Ex. Ksp of BaSO₄ = 1.1 x 10^{-10}

 $Ba(NO_3)_2 = 0.50L, 0.002M$

 $Na_2SO_4 = 0.50L, 0.008M$

Possible precipitate = $Ba(SO_4)_2$

The product of the ions barium and sulfate has to exceed the Ksp

Because they are mixed, the concentrations are cut in half, so...

Precipitate Example...

 $Ba(NO_3)_2 = 0.001M \text{ (mol/L)}$ $Na_2SO_4 = 0.004M \text{ (mol/L)}$

 $[Ba^{2+}] = 0.001M$ $[SO_4^{2-}] = 0.004M$

 $Q_{sp} = [Ba^{2+}] \times [SO_4^{2-}] = (0.001M) \times (0.004M) = 4 \times 10^{-6}$

 $K_{SD} = 1.1 \times 10^{-10}$ Does a precipitate form?

Finding Equilibrium Ion Concentration Using a Common Ion

Work through Sample Problem 18.4, p. 564.

Questions 19, 20, p. 564

Questions 21 – 28, p. 565

Section 18.3 Review