

KEEP IT SIMPLE SCIENCE

Chemistry Module 5

Chemical Equilibrium WORKSHEETS

	Worksheet 1	The Basics	Student Name
equi	hen any chemical reaction reac librium in a closed system, wha at the concentration of all the cl ent?	at becomes true	4. Why can a dynamic equilibrium NOT occur in an open system?
b) E	xplain why this is so, in the cas i) static equilibrium.	e of a:	5.Write an ionic equation to represent the dynamic equilibrium reached between the solid and the dissolved ions in each of these species, when a solution reaches saturation.a) potassium bromide
	ii) dynamic equilibrium.		
	, , , ,		b) magnesium chloride
	t are the thermodynamic reaso tions (eg combustion) are neve		c) iron (III) nitrate
			d) ammonium sulfate
	t is meant by "open" or "closed ems?	d" chemical	e) calcium carbonate



Worksheet 2 L Practice Questions

Le Chatelier's Principle

Student Name.....

1. $NO_{2(g)}$ reacts with itself as follows:

$$2NO_{2(g)} \longrightarrow N_2O_{4(g)}$$

The reaction to the right is exothermic, so heat can be considered as a "product".

$$2NO_{2(g)} \longrightarrow N_2O_{4(g)} + heat$$

Note that as the reaction proceeds to the right, 2 moles of gas form 1 mole of gas, so in a fixed volume container the pressure would drop as the reaction proceeds to the right.

Imagine a sealed container in which a mixture of these gases has reached equilibrium.

State which way the equilibrium would shift if each of the following disturbances were made to the mixture. Explain each answer.

- i) Increase in temperature.
- ii) Compress the mixture, thereby increasing pressure.
- iii) Injecting extra N₂O₄, without changing pressure.
- iv) Decrease the temperature.
- v) Spray in a little water. (NO₂ dissolves, N₂O₄ does not.)
- vi) Decreasing the total gas pressure by expanding the volume of the container.

2. If hydrogen iodide (covalent molecular) is dissolved in water, some of the molecules ionise, and an equilibrium is reached:

$$HI_{(aq)} \longrightarrow H^+_{(aq)} + I^-_{(aq)}$$

What is the effect on this equilibrium of: (Explain each)

- i) adding NaI_(aq) solution, which increases the concentration of iodide ions.
- ii) Adding NaOH, which reacts with H⁺ ions, and reduces their concentration.
- iii) Dissolving extra HI in the solution.
- iv) It is found that raising the temperature of an equilibrium mixture has the effect of increasing the concentration of ions. Deduce whether the reaction as written is exo- or endothermic.
- 3. Ammonia is manufactured from its elements by the reaction

$$N_{2(g)} + 3H_{2(g)} \longrightarrow NH_{3(g)} + heat$$

- i) To maximize the yield of ammonia, the reaction is carried out under very high pressure. Explain how this helps.
- ii) The temperature of the reaction is kept fairly high to speed up the <u>rate</u> of the reaction. What effect does higher temperature have on the equilibrium?
- iii) During the reaction, ammonia is constantly removed from the reaction vessel, and more reactant gases constantly pumped in. Explain the effect this has.



Worksheet 3

Chemical Equilibrium

Fill in the blanks

Many chemical reactions do not go to completion, but reach a a)......
in which both forward and reverse reactions are running at b).....

According to c)......'s Principle, if an equilibrium system is disturbed, the equilibrium will shift in the direction which d).....

Student Name.....

The value of K for the "reverse equation" is the j)..... of the original. The value of Keq is constant only at a given k)......

The values of K for an exothermic reaction

I)...... as temperature increases.

The values for an m)..... reaction
will n)..... as temp. increases.

Worksheet 4

Equilibrium Constant K_{eq}

Student Name.....

Each chemical species is in the gas state, unless indicated otherwise.

1. Write the expression for the equilibrium constant for:

a) $PCI_5 \rightleftharpoons PCI_3 + CI_2$

2. The reaction $H_{2(g)} + I_{2(g)} + \text{heat} \implies 2HI_{(g)}$

reached equilibrium at 400° C. Predict the effect on the equilibrium concentration of ${\rm HI}_{(\alpha)}$ of: (explain each answer)

i) increasing the temperature

b) $2H_2O \rightleftharpoons 2H_2 + O_2$

ii) compressing the mixture to a higher pressure

c) 2NO + CI₂ = 2NOCI

- iii) increasing pressure by pumping in extra H_{2(a)}
- d) $AI(OH)_{3(s)} \rightleftharpoons AI^{3+}_{(aq)} + 3OH^{-}_{(aq)}$
- iv) adding a catalyst

e) $2H_2S \rightleftharpoons 2H_2 + S_2$

v) increasing pressure by pumping in argon gas



Worksheet 5

Calculating K_{eq}

Practice Problems

Student Name.....

1.

The reaction

$$H_{2(g)} + I_{2(g)} + heat \implies 2HI_{(g)}$$

has reached equilibrium at 400°C. The concentrations at equilibrium are: $[H_2]$ = 0.0195molL⁻¹, $[I_2]$ = 0.0211molL⁻¹ and [HI] = 0.153molL⁻¹.

a) Calculate the value of $K_{\rm eq}$.

b) The temperature of this same gas mixture was changed, and the new value for K = 22.3. Was the mixture heated or cooled? Explain.

2. (cont)

c) At equilibrium, the concentration of hydrogen iodide gas [HI]=0.0552molL $^{-1}$. Find the concentrations of H $_2$ and I $_2$ gases. (Hint: they will be equal to each other)

3. Carbon monoxide reacts with steam as follows: $CO_{(g)} + H_2O_{(g)} \rightleftharpoons CO_{2(g)} + H_{2(g)}$

A mixture containing 10.0 moles each of CO gas and steam was introduced at high pressure and temperature into a 4.00 litre container. After equilibrium was reached it was found that there were 6.50 moles of CO_2 in the container.

a) Calculate the equilibrium concentrations of each reactant and product.

- A sample of pure hydrogen iodide gas (HI) was placed into a container and heated to 400°C. It decomposed in the reverse reaction to that described in Q1, and the mixture reached an equilibrium.
- a) Write an equation for the reaction.

b) Calculate the value of the equilibrium constant for this temperature.

- b) Write an expression for the equilibrium constant, and predict its value for equilibrium at 400°C. (Use answer Q1)
- c) What would have been different if the reaction had been carried out in the presence of a catalyst?



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(8)	Worksheet 6	Ionic Precipitations	
keep it simple science	Practice Problen	ns Student Name	

Feeb pair of ionic compounds below will produce a	4 coloium hudrovido 9 notocoium combonete		
Each <u>pair</u> of ionic compounds below will produce a precipitate if solutions are mixed together.	4. calcium hydroxide & potassium carbonate		
For each pair:	a)		
 a) name the precipitating compound. b) write a <u>full ionic equation</u> for the reaction, including all states. 	b)		
c) write the <u>net ionic equation</u> for the <u>equilibrium</u> . d) write an expression for the "Solubility Product"			
(K _{sp}) for the equilibrium.	c)		
1. calcium chloride & silver nitrate			
a)	d)		
b)	5. silver nitrate & magnesium sulfate		
,	a)		
c)	,		
,	b)		
d)			
	c)		
2. lead(II) nitrate & sodium carbonate			
a)	d)		
b)	6. potassium bromide & lead(II) nitrate		
	a)		
c)			
	b)		
d)	c)		
3. sodium sulfate & barium chloride	C)		
a)	d)		
u,	7. iron(III) chloride & calcium hydroxide		
b)	a)		
	" ,		
c)	b)		
d)	c)		
	d)		



Worksheet 7

K_{sp} and Solubility

Practice Problems

Student Name.....

Refer to the reactions in Worksheet 6.

Q4.

The $\rm K_{sp}$ value (at 25°C) for each of the insoluble products will be found in your Chem Data Sheet.

For each substance, calculate its max. solubility in water at 25°C. Show working.

Q1.

Q5.

Q2.

Q6.

Q3.

Q7.



Worksheet 8 Practice Problems

Predicting a Precipitate with $K_{\rm sp}$

Student Name.....

In each case, equal volumes of the two solutions were mixed together. Would a precipitate form at 25°C? Show all working & reasoning.

1. copper (II) sulfate, 0.005 molL⁻¹ and sodium carbonate, 0.001 molL⁻¹.

3. lead (II) nitrate, 0.05 molL⁻¹ and sodium chloride, 0.008 molL⁻¹.

4. potassium hydroxide, 0.002 molL⁻¹. and magnesium chloride, 0.006 molL⁻¹.

2. sodium phosphate, 0.0004 molL⁻¹. and iron(III) chloride, 0.0005 molL⁻¹.

5. sodium phosphate, 0.004 molL⁻¹. and silver nitrate, 0.05 molL⁻¹.



Answer Section

Worksheet 1

1.

a) The concentrations remain constant & do not change over time.

b)

- i) In a static equilibrium this is because all reaction has ceased.
- ii) In a dynamic equilibrium, the concentrations do not change because the forward and reverse reactions are running at the same rate.

2.

Reactions such as combustions are strongly exothermic and may also result in a large increase in entropy. These energy considerations make it almost impossible that they could run in reverse.

3.

An open system is one in which substances and/or energy can readily flow in or out from/to the surroundings.

A closed system is one in which no substances or energy can exchange with the environment. Although there can be no "perfect" closed system, a sealed, insulated container can be a very good approximation to one.

4.

In an open system, the products and heat energy of the reaction may be lost from the system and disperse into the environment. In such case it becomes impossible to establish a dynamic equilibrium because the temperature keeps changing and/or substances are no longer present to take part.

5

a)
$$KBr_{(s)} \rightleftharpoons K^+_{(aq)} + Br^-_{(aq)}$$

b)
$$\mathrm{MgCl}_{2(s)}$$
 \Longrightarrow $\mathrm{Mg^{2+}}_{(aq)}$ + $\mathrm{2Cl^{-}}_{(aq)}$

c)
$$Fe(NO_3)_{3(s)} = Fe^{3+}_{(aq)} + 3NO_3^{-}_{(aq)}$$

d)
$$(NH_4)_2SO_{4(s)} \implies 2NH_4^+_{(aq)} + SO_4^{2-}_{(aq)}$$

e)
$$CaCO_{3(s)}$$
 \Longrightarrow $Ca^{2+}_{(aq)}$ + $CO_3^{2-}_{(aq)}$

Worksheet 2

1.

- i) Left, to try to use heat & reduce temp.
- ii) Right, to reduce the pressure again.
- iii) Left, to decrease concentration of product.
- iv) Right, to release heat & increase temp.
- v) Left, to make more reactant & increase pressure.
- vi) Left, to increase pressure again.

2.

- i) Equilib. shifts left, to decrease conc. of iodide ion.
- ii) Equilib. shifts right, to increase conc. of hydrogen ion.
- iii) Equilib. shifts right, to decrease conc. of HI.
- iv) Endothermic (heat is a reactant), since higher temp causes shift to right to consume heat.

3

- i) High pressure causes equilibrium to shift right to reduce the total moles of gas and pressure.
- ii) Shifts equilibrium to left.
- iii) Removing product & adding reactant keeps shifting equilib. to right, so yield is maximised.

Worksheet 3

- a) dynamic equilibrium b) the same rate
- c) Le Chatelier's
- d) counteracts the disturbance
- e) temperature & concentration
- f) volume or pressure g) Equilibrium Constant
- h) products
- i) reactants
- j) reciprocal
- k) temperature
- I) decreases
- m) endothermic
- n) increase

Worksheet 4

1.

a)
$$K_{eq} = [PCI_3] \times [CI_2] / [PCI_5]$$

b)
$$K_{eq} = [H_2]^2 \times [O_2] / [H_2O]^2$$

c)
$$K_{eq} = [NOCI]^2 / [CI_2] \times [NO]^2$$

d)
$$K_{eq} = [OH^{-}]^{3} \times [AI^{3+}] / [AI(OH)_{3}]$$

e)
$$K_{eq} = [H_2]^2 \times [S_2] / [H_2S]^2$$

2.

- i) [HI] would increase as equilibrium shifts to the right in an attempt to use heat to counteract temp. increase.
- ii) no effect, since there are equal gas volumes on each side of the equation... compression increases all concentrations equally.
- iii) [HI] would increase as equil. shifts right to counteract greater $[H_2]$.
- iv) Catalysts have no effect on equil... only shorten the time to reach equil.
- v) No effect. Increasing pressure with another gas has no effect on concentrations of reactants or product.



Answer Section

Worksheet 5

a)
$$K_{eq} = [HI]^2 / [H_2] \times [I_2] = (0.153)^2 / (0.0195 \times 0.0211)$$

b) K_{eq} has decreased. Endothermic reactions increase K_{eq} with temp. increase. Therefore, temp. must have decreased.

2.
a)
$$2HI_{(g)} \Longrightarrow H_{2(g)} + I_{2(g)}$$
b)
$$K = [H_2] \times [I_2] / [HI]^2$$
Its value will be the reciprocal of 56.9 - 0.0176

Its value will be the reciprocal of 56.9 = 0.0176

c) 0.0176 =
$$[H_2] \times [I_2] / (0.0552)^2$$

 $\therefore [H_2] \& [I_2] = 5.78 \text{ molL}^{-1}$

3.

- a) Since all species react in equal molar quantities, then the molar quantities at equilibrium are: $n(CO_2) = 6.50 \text{ mol (given)}$ and $n(H_2) = 6.50 \text{ mol}$ n(CO) = 3.50 mol (since 6.50mol has been used up) $n(H_2O) = 3.50$ mol (as above) Since in a 4.00L container, and c = n/V
- \therefore c(CO₂) and c(H₂) = 6.50/4.00 = 1.625molL⁻¹ and c(CO) and c(H₂O) = 3.50/4.00 = 0.875molL⁻¹
- b) $K_{eq} = [CO_2]x[H_2]/[CO]x[H_2O]$ $= 1.625 \times 1.625 / (0.875 \times 0.875)$ = 3.45
- c) Equilibrium would have been achieved faster. The equil. itself would be exactly the same, with same value for Ked.

Worksheet 6

- a) silver chloride
- b) $Ca^{2+}_{(aq)} + 2Cl_{(aq)} + 2Ag_{(aq)}^{+} + 2NO_{3(aq)}^{-}$ \rightarrow 2AgCl_(s) + Ca²⁺_(aq) + 2NO₃-_(aq)
- $Cl_{(aq)}^- + Ag_{(aq)}^+$ \Longrightarrow AgCl_(s) equation may be written in reverse and that makes more sense for part (d)
- d) $K_{sp} = [Ag^{+}]x[CI^{-}]$

2

- a) lead(II) carbonate
- b) $Pb^{2+}_{(aq)} + 2NO_{3(aq)} + 2Na^{+}_{(aq)} + CO_{3(aq)}^{2-}$ \longrightarrow PbCO_{3(s)} + 2Na⁺_(aq) + 2NO_{3 (aq)}
- $Pb^{2+}_{(aq)} + CO_3^{2-}_{(aq)} \longrightarrow PbCO_{3(s)}$ c) equation may be written in reverse and that makes more sense for part (d)

d)
$$K_{sp} = [Pb^{2+}]x[CO_3^{2-}]$$

3.

a) barium sulfate

b)
$$Ba^{2+}_{(aq)} + 2Cl^{-}_{(aq)} + 2Na^{+}_{(aq)} + SO_{4}^{2-}_{(aq)}$$

BaSO_{4(s)} + $2Na^{+}_{(aq)} + 2Cl^{-}_{(aq)}$

 $Ba^{2+}_{(aq)} + SO_4^{2-}_{(aq)} \rightleftharpoons$ equation may be written in reverse and that makes more sense for part (d)

d)
$$K_{sp} = [Ba^{2+}]x[SO_4^{2-}]$$

4.

a) calcium carbonate

b)
$$Ca^{2+}_{(aq)} + 2OH^{-}_{(aq)} + 2K^{+}_{(aq)} + CO_{3}^{2-}_{(aq)}$$

$$\longrightarrow CaCO_{3(s)} + 2K^{+}_{(aq)} + 2OH^{-}_{(aq)}$$

- $Ca^{2+}_{(aq)} + CO_3^{2-}_{(aq)} \longrightarrow CaCO_{3(s)}$ equation may be written in reverse and that makes more sense for part (d)
- $K_{sp} = [Ca^{2+}]x[CO_3^{2-}]$

- a) silver sulfate
- b) $2Ag^{+}_{(aq)} + 2NO_{3(aq)}^{-} + Mg^{2+}_{(aq)} + SO_{4(aq)}^{-2-}$

$$\longrightarrow$$
 Ag₂SO_{4(s)} + Mg²⁺(aq) + 2NO₃(aq)

 $2\mathsf{Ag^+}_{(\mathsf{aq})} + \mathsf{SO_4}^{2\text{-}}_{(\mathsf{aq})} \quad \Longrightarrow \quad \mathsf{Ag_2SO_{4(\mathsf{S})}}$

equation may be written in reverse and that makes more sense for part (d)

d)
$$K_{sp} = [Ag^+]^2 \times [SO_4^{2-}]$$

- a) lead(II) bromide
- b) $Pb^{2+}_{(aq)} + 2NO_{3(aq)} + 2K^{+}_{(aq)} + 2Br_{(aq)}$

$$\longrightarrow$$
 PbBr_{2(s)} + 2K⁺_(aq) + 2NO₃ (aq)

 $Pb^{2+}_{(aq)} + 2Br_{(aq)}^{-}$ ⇒ PbBr_{2(s)}

equation may be written in reverse and that makes more sense for part (d)

d)
$$K_{sp} = [Pb^{2+}]x[Br^{2}]^{2}$$

7.

- a) iron(III) hydroxide
- b) $3Ca^{2+}_{(a\alpha)} + 6OH_{(a\alpha)}^{-} + 2Fe^{3+}_{(a\alpha)} + 6CI_{(a\alpha)}^{-}$

$$\longrightarrow$$
 2Fe(OH)_{3(s)} + 3Ca²⁺_(aq) + 6Cl⁻_(aq)

 $Fe^{3+}_{(aq)} + 3OH^{-}_{(aq)} \longrightarrow Fe(OH)_{3(s)}$ c)

equation may be written in reverse and that makes more sense for part (d)

d)
$$K_{sp} = [Fe^{3+}]x[OH^{-}]^{3}$$



Answer Section

Worksheet 7

1.

$$AgCl_{(s)} \longrightarrow Cl_{(aq)}^{-} + Ag_{(aq)}^{+}$$
 $K_{sp} = [Ag^{+}]x[Cl^{-}] = 1.77 \times 10^{-10}.$

Since $[Ag^+] = [CI^-] = "c"$, then $c^2 = 1.77x10^{-10}$

$$\therefore$$
 c = 1.77x10⁻¹⁰ = 1.33x10⁻⁵ molL⁻¹.

2.

$$PbCO_{3(s)} \rightleftharpoons Pb^{2+}_{(aq)} + CO_3^{2-}_{(aq)}$$

$$K_{sp} = [Pb^{2+}]x[CO_3^{2-}] = 7.40 \times 10^{-14}.$$

Since $[Pb^{2+}] = [CO_3^{2-}] = "c"$, then $c^2 = 7.40x10^{-14}$

$$\therefore$$
 c = $\sqrt{7.40 \times 10^{-14}}$ = 2.72 x 10⁻⁷ molL⁻¹.

3.

$$BaSO_{4(s)} \stackrel{}{\longrightarrow} Ba^{2+}_{(aq)} + SO_4^{2-}_{(aq)}$$

$$K_{sp} = [Ba^{2+}]x[SO_4^{2-}] = 1.08 \times 10^{-10}.$$

Since $[Ba^{2+}] = [SO_4^{2-}] = \text{``c''}$, then $c^2 = 1.08x10^{-10}$

$$\therefore$$
 c = $\sqrt{1.08 \times 10^{-10}}$ = 1.04 x 10⁻⁵ molL⁻¹.

4.

$$CaCO_{3(s)} \stackrel{}{\Longrightarrow} Ca^{2+}_{(aq)} + CO_3^{2-}_{(aq)}$$

$$K_{sp} = [Ca^{2+}]x[CO_3^{2-}] = 3.36 \times 10^{-9}.$$

Since $[Ca^{2+}] = [CO_3^{2-}] = \text{``c''}$, then $c^2 = 3.36 \times 10^{-9}$

$$\therefore$$
 c = $\sqrt{3.36 \times 10^{-9}}$ = 5.80 x 10⁻⁵ molL⁻¹.

5.

$$Ag_2SO_{4(s)} \Longrightarrow 2Ag^+_{(aq)} + SO_4^{2-}_{(aq)}$$

$$K_{sp} = [Ag^+]^2 \times [SO_4^{2-}] = 1.20 \times 10^{-5}.$$

If $[SO_4^{2-}] = "c"$, then $[Ag^+] = 2c$, so $K_{sp} = (2c)^2 \times c$

So
$$4c^3 = 1.20 \times 10^{-5}$$

 $\therefore c = \sqrt[3]{\frac{1.20 \times 10^{-5}}{4}} = 1.44 \times 10^{-2} \text{molL}^{-1}.$

6.

$$Pb^{2+}_{(aq)} + 2Br_{(aq)} \Longrightarrow PbBr_{2(s)}$$

$$K_{sp} = [Pb^{2+}]x[Br^{-}]^2 = 6.60 \times 10^{-6}.$$

If $[Pb^{2+}] = "c"$, then [Br] = 2c, so $K_{sp} = c \times (2c)^2$

So
$$4c^3 = 6.60 \times 10^{-6}$$

 $\therefore c = \sqrt[3]{\frac{6.60 \times 10^{-6}}{4}} = 1.18 \times 10^{-2} \text{molL}^{-1}.$

7.
$$Fe^{3+}_{(aq)} + 3OH_{(aq)} \longrightarrow Fe(OH)_{3(s)}$$

$$K_{sp} = [Fe^{3+}]x[OH^{-}]^{3} = 2.79 \times 10^{-39}.$$

If [Fe³⁺] = "c", then [OH-] = 3c, so $K_{sp} = c \times (3c)^3$

So
$$9c^4 = 2.79 \times 10^{-39}$$

 $\therefore c = \sqrt[4]{\frac{2.79 \times 10^{-39}}{9}} = 1.33 \times 10^{-10} \text{molL}^{-1}.$

Worksheet 8

1

copper(II) carbonate is the possible precipitate. $CuCO_{3(s)} \stackrel{}{\Longrightarrow} Cu^{2+}_{(aq)} + CO_3^{2-}_{(aq)}$

$$\begin{split} & \text{K}_{sp} = [\text{Cu}^{2+}]\text{x}[\text{CO}_3^{2-}] = 1.4 \text{ x } 10^{-10}. \\ & \text{Concentrations of each ion, after dilution by mixing equal} \\ & \text{volumes: } [\text{Cu}^{2+}] = 0.0025, [\text{CO}_3^{2-}] = 0.0005_{\text{molL}^{-1}}. \\ & \therefore \text{ actual Sol.Prod.} = 0.0025 \text{ x } 0.0005 = 1.25 \text{ x } 10^{-6}. \\ & \text{This is far higher than K}_{sp}, \text{ so pptte WILL form.} \end{split}$$

2. iron(III) phosphate is the possible precipitate. FePO_{4(s)} \rightleftharpoons Fe³⁺_(a0) + PO₄³⁻_(a0)

 $K_{\rm sp} = [{\rm Fe^{3+}}] \times [{\rm PO_4}^{3\text{-}}] = 9.91 \times 10^{-16}.$ Concentrations of each ion, after dilution by mixing equal volumes: $[{\rm Fe^{3+}}] = 0.0002$, $[{\rm PO_4}^{3\text{-}}] = 0.00025_{\rm molL^{-1}}.$ \therefore actual Sol.Prod. = 0.0002 x 0.00025 = 5.0 x 10⁻⁸. This is far higher than $K_{\rm sp}$, so pptte WILL form.

3. lead(II) chloride is the possible precipitate. PbCl_{2(s)} \rightleftharpoons Pb²⁺_(aq) + 2Cl⁻_(aq)

$$\begin{split} & \text{K}_{\text{sp}} = [\text{Pb}^{2+}]\text{x}[\text{Cl}^{-}]^2 = 1.7 \text{ x } 10^{-5}. \\ \text{Concentrations of each ion, after dilution by mixing equal volumes: } [\text{Pb}^{2+}] = 0.025, [\text{Cl}^{-}] = 0.004_{\text{molL}^{-1}}. \\ & \therefore \text{ actual Sol.Prod.} = 0.025 \text{ x } 0.004^2 = 4.0 \text{ x } 10^{-7}. \\ \text{This is less than } \text{K}_{\text{sp}}, \text{ so pptte WILL NOT form.} \end{split}$$

4. magnesium hydroxide is the possible precipitate. ${\rm Mg(OH)_{2(s)}}~{\rm Mg^{2+}_{(aq)}} + {\rm 2OH^{-}_{(aq)}}$

 $K_{sp} = [Mg^{2+}]x[OH^-]^2 = 5.61x \ 10^{-12}.$ Concentrations of each ion, after dilution by mixing equal volumes: $[Mg^{2+}] = 0.003$, $[OH^-] = 0.001_{molL^{-1}}.$ \therefore actual Sol.Prod. = $0.003 \ x \ 0.001^2 = 3.0 \ x \ 10^{-9}.$ This is higher than K_{sp} , so pptte WILL form.

silver phosphate is the possible precipitate. $Ag_3PO_{4(s)} \rightleftharpoons 3Ag^+_{(aq)} + PO_4^{3-}_{(aq)}$

 $K_{\rm sp} = [Ag^+]^3x[PO_4^{\ 3^-}] = 8.89 \ x \ 10^{-17}.$ Concentrations of each ion, after dilution by mixing equal volumes: $[Ag^+] = 0.025, [PO_4^{\ 3^-}] = 0.002_{\rm molL^{-1}}.$

∴ actual Sol.Prod. = 0.025^3 x 0.002 = 3.13 x 10^{-8} . This is far higher than K_{sp} , so pptte WILL form.