ph and solubility

This salt's solubility is pH dependent. How?

- * In a BASIC solution, the concentration of hydroxide ion in solution is high. , so solubility is LOWER than in pure water.
- * In an ACIDIC solution, we have a significant amount of hydronium, which can react with hydroxide. This lowers the hydroxide concentration and makes magnesium hydroxide MORE SOLUBLE

Generalizing

If a compound is BASIC, then it will be LESS SOLUBLE in basic solutions, and MORE SOLUBLE in acidic solutions!

If a compound is ACIDIC, then it will be MORE SOLUBLE in basic solutions, and LESS SOLUBLE in acidic solutions!

If a compound is NEUTRAL (neither acidic nor basic), then its solubility will be UNAFFECTED by pH

- are ions that result from the reaction of a Lewis base (like water, ammonia, hydroxide ion, etc.) with a metal ion
- The Lewis base attaches to the metal ion by forming a COORDINATE COVALENT BOND with the metal ion.

$$A_{g}^{\dagger}(aq) + 2 : NH_{3}(aq) \longrightarrow H_{3}N - A_{g}^{\dagger} - NH_{3}(aq)$$

$$\left(A_{g}^{\dagger}(aq) + 2NH_{3}(aq) \rightarrow A_{g}(NH_{3})_{2}^{\dagger}(aq)\right)$$

- The product of the reaction is called a "COMPLEX", and the attached Lewis bases are called "LIGANDS"

- Described by the FORMATION CONSTANT, Kf

$$A^{+} + B \rightleftharpoons AB^{+}$$

$$K_{p} = \frac{[AB^{+}]}{[A^{+}][B]}$$

ex:
$$Ag^{+} + 2NH_{3} = A_{g}(NH_{3})_{2}^{+}$$

$$K_{F} = \frac{[A_{g}(NH_{3})_{2}^{+}]}{[A_{g}^{+}][NH_{3}]^{2}} = 1.7 \times 10^{7}$$

What does this value for the equilibrium constant say about the favorability of the formation of the complex ion?

Since the formation of these complex ions is so favorable, we often assume that these reactions go to completion, and instead look at the small amount of complex ion that DISSOCIATES!

$$AB^{+} = A^{+} + B$$

$$Kd \text{ is called the DISSOCIATION CONSTANT, and it is equal to 1/Kf}$$

Kd is called the

$$(2 = [Ag^{\dagger}][NH_3]^2 = \frac{1}{[Ag(NH_3)_2^{\dagger}]} = \frac{1}{1.7 \times 10^7} = 5.9 \times 10^{-8}$$

AMPHOTERIC COMPOUNDS

- All metal hydroxides react with ACIDS, but SOME metal hydroxides can react with BASES by forming a complex ion.

$$A1(0H)_3(4) + 3H_30^+ = A1^{3+}(uq) + 3H_20(l)$$

... Aluminum hydroxide is soluble in acidic solutions.

... And it is also soluble in bases due to the formatkion of this complex ion!

- So aluminum hydroxide is relatively insoluble in pure water, but its solubility increases greatly if the pH goes either up or down.

COMPLEX IONS AND SOLUBILITY

- What is the effect of complex formation on solubility?

$$Ag(l(s) = Ag^{\dagger}(ag) + Cl^{\dagger}(ag)$$

$$Ag^{\dagger}(ug) + 2NH_3(ag) = Ag(NH_3)_2^{\dagger}$$

What will the presence of ammonia do to the solubility of silver chloride?

- Since the formation of the silver-ammonia complex is favorable, we expect that any dissolved silver ion would react with ammonia to make the complex.
- (2) This will REDUCE the concentration of free silver ion.
- The reduction of free silver ion will cause more silver chloride to dissolve (Le Chateleir's principle the equilibrium will try to produce more free silver ion to replace what the ammonia has removed)

So, the presence of a ligand which can form a complex with an ion from a salt will greatly INCREASE the solubility of that salt!

Calculate the solubility (in ppm) of silver chloride in 0.10 M ammonia solution.

$$Ag(l(s)) = Ag^{t}(aq) + Cl^{t}(aq); Ksp=1.8 \times 10^{-10}$$

 $Ag^{t}(aq) + 2NH_{3}(aq) = Ag^{t}(aq); Kf = 1.7 \times 10^{-7}$

How do we find Kc for the added reactions?

We must solve this equation to find out how much AgCl dissolves.

$$K_{SP} \times K_{f} = \frac{[A_{g}^{+}][C]^{-}[A_{g}(WH_{3})_{2}^{+}]}{[A_{g}^{+}][NH_{3}]^{2}}$$

$$K_{SP} \times K_{f} = \frac{[A_{g}(WH_{3})_{2}^{+}][C]^{-}}{[NH_{3}]^{2}} = K_{C}$$

$$NH_{3}^{-}$$

When you add two reactions to get a new reaction, the equilibrium constants of the original two reactions multiply to give you the new Kc

See section 14.2 for more details!

[Ag(NH3)2+][C1-]=0.00306	
[NH3]2	

$$\frac{\chi^{2}}{(0.10-2\chi)^{2}}=0.00306$$

Species	[Initial]	7	[Equilibrium]
Ag (N43)2	0	tχ	X
CIT	0	4%	X
NH ₃	0.10	-2x	0.10 - 2x

Ag: 1 x 107.9

C1: 1 x 35.45

143.35 g Ag(1/m)

1.1106345335 x = 0.00553172667

So "x" is equal to the MOLAR solubility of sinver chloride. Let's convert it to ppm! $= [Ag(I)]\lambda$

Compare the solubility of AgCl in distilled water with the solubility in 0.10 M ammonia AND 0.10 M NaCl.

(We see a significant drop in the solubility of AgCl in the presence of the common ion chloride.

So in general,

- * The presence of a ligand (which forms a complex with one of the ions in the compound) greatly INCREASES solubility.
- * The presence of a common ion greatly DECREASES solubility.