To predict whether a salt at a given concentration will precipitate out, calculate the reaction quotient Q and compare it to the Ksp

Example: $A_gCI(s) = A_g^{\dagger}(a_g) + CI^{\dagger}(a_g) j k_{sp} = 1.8 \times 10^{-10}$ $Q = CA_g^{\dagger}[CI^{\dagger}]$

IF...

- Q < Ksp; the reaction proceeds to produce more products (dissolved ions), so more solid is able to dissolve: NO PRECIPITATION
- Q > Ksp; the reaction proceeds to produce more reactants (solid), so solid falls out of solution: PRECIPITATION OCCURS
- ¥ Q = Ksp; the reaction is at equilibrium. PRECIPITATION IS JUST BEGINNING

Would a solution with (Ag+) = 0.014 M and (Cl-) = 0.00042 M precipitate?

$$Q = [Ag^{\dagger}][(1]^{-}] = (0.014)(0.00042) = 5.88410^{-6}$$

 $5.88410^{-6} > 1.8410^{-10}$
 $Q > Ksp, so precipitation occurs.$

The "common ion effect" affects the solubility of a compound in solution. The presence of one of the ions in a salt in the solution will REDUCE THE SOLUBILITY of that salt!

Silver chloride is much less soluble in a solution of 0.1 M NaCl than it is in distilled water. Why? The presence of CHLORIDE ION forces the solubility equilibrium back to the left, meaning less silver chloride can dissolve!

Solubility can also be affected by pH - depending on the acidic or basic properties of the salt!

A second example: Salicylic acid in the characterization lab

Calculate the solubility of AgCI (FW = 143.35 g/mol) in distilled water. Then, calculate the solubility of AgCI in 0.10 M NaCI solution. Report both answers in parts per million (mg/L)

Ag (I(s)
$$\rightleftharpoons Ag^{\dagger}(aq) + C[(aq)] Ksp = CAg^{\dagger}(C[-] = 1.8 \times 10^{-10})$$

For distilled water...

Let "x" equal the change in dissolved Ag+ concentration.

$$(\chi)(\chi) = 1.8 \times 10^{-10}$$

$$\chi^{2} = 1.8 \times 10^{-10}$$

$$\chi = 1.341640786 \times 10^{-5} M = CAg^{+}] = CAg(1) \lambda issolved$$

Convert answer's units to ppm (mg/L)...

1.341640786 x 10 - Smol fy (1 x
$$\frac{143.35g \text{ Ag(1)}}{\text{mol fy (1)}} \times \frac{\text{mg}}{\text{10}^{-3}g} = 1.4 \text{ ppm fy (1)}$$

converts mol to g

converts g to mg

For solubility in 0.10 M NaCl solution...

Species	[Initial]	Δ	[[Gavilibrium]
Ag+	O	ΤX	X
C1-	0.10	+ X	D,10+x
$(\chi)(0.10 + \chi) = 1.8 \times 10^{-10}$ We know that x << 0.10, so 0.10 + $\chi \approx 0.10$			
$(X)(D.10) = 1.8 \times 10^{-10}$ $X = 1.8 \times 10^{-9} M = CAg^{+} = CAgC Jissolve$			

$$\frac{1.8 \times 10^{-9} \text{ mol Ag Cl}}{\text{Compare:}} \times \frac{143.35 \text{ gAgCl}}{\text{mol Ag Cl}} \times \frac{\text{mg}}{16^{-3} \text{g}} = \frac{12.6 \times 10^{-4} \text{ ppm Ag Cl in 0.10 M NaCl}}{(0.00026 \text{ ppm})}$$

1.9 ppm AgCl in distilled water vs 0.00026 ppm AgCl in 0.10 M NaCl

Conclusion: The presence of a common ion greatly decreases solubility

ph and solubility

This compound'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