SOLUTION: Homogeneous mixture of substances Solutions contain:

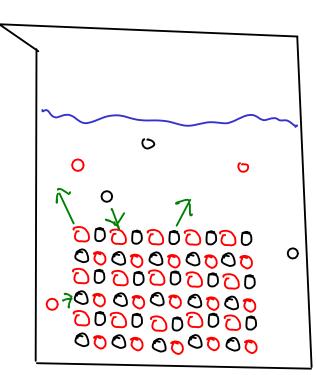
SOLUTE: Component(s) of a solution present in small amount SOLVENT: Component of a solution present in greatest amount

> We usually call water the solvent in aqueous mixtures, even if the water is present in smaller amount than another component

SOLUBILITY: The amount of a solute that will dissolve in a given volume of solvent

SATURATED SOLUTION: Contains the maximum amount of solute that it is possible to dissolve in a given volume of solvent!

A SATURATED SOLUTION is a solution where dissolved solute exists in an EQUILIBRIUM with undissolved solute!



Example: Consider a saturated solution of silver chloride:

$$A_g(I(s) \rightleftharpoons A_g^+(a_q) + CI^-(a_q))$$

At equilibrium, the rate of dissolving equals the rate of crystallization!

$$A_{g}(I(s) \rightleftharpoons A_{g}^{+}(a_{q}) + CI^{-}(a_{q}))$$

$$K_{c} = \left[I_{g}^{+}\right]\left[CI^{-}\right] = \left[I_{c} \otimes XIO^{-10}\right]$$

... What does this equilibrium constant tell us? That silver chloride isn't very soluble!

Remember, Ksp is an equilibrium constant, so everything that applies to equilibrium constants applies to the solubility constant - including what to do with coefficients:

What is the solubility product constant expression for calcium phosphate? $\binom{\alpha_3(PO_4)_2(s)}{\longrightarrow} \frac{3(a^{2t}(a_4) + 2PO_4^3(a_4))}{(a_5)^2}$ $\binom{\beta_5}{\beta_5} = \binom{(a^{2t})^3}{(PO_4)^2} \binom{PO_4^3}{2}$

Solubility calculations and Ksp

You can calculate the solubility of a compound if you know Ksp!

Calculate the solubility (in g/L) of lead(II) iodide at 25C. FW = 461.0 g/mol (See Appendix J in OpenStax for solubility product information!)

$$P_{b} I_{2}(s) \rightleftharpoons P_{b}^{2+}(aq) + 2 I^{-}(aq); K_{sp} = [P_{b}^{2+}][I^{-}]^{2} = [.4x/0^{-8}]$$

Set up an equilibrium chart and solve.

$$\frac{\text{Speciles} [\text{Initial}] \text{A} [\text{Cequilibrium}]}{\text{Pb}^{24}} \qquad \text{Let "x" equal the change in lead(II) ion concentration}} \\ \frac{\text{Pb}^{24}}{\text{I}^{-}} \qquad \frac{\text{H} \times \text{X}}{(\chi)(2\chi)^{2} = 1.4\chi10^{-8}} \\ \text{H} \chi^{3} = 1.4\chi10^{-8} \\ \text{H} \chi^{3} = 1.4\chi10^{-8} \\ \text{X} = 0.001\text{S}[8294\text{S} \text{M} = [\text{Pb}^{24}] = [\text{Pb}\text{J}_2]_{\text{dissolved}} \\ \text{The concentration of lead(II) ion equals the concentration of dissolved lead(II) iodide. But our current units are mol/L. We need to convert to g/L.} \\ \frac{0.601\text{S}[8294\text{S} \text{mol}] \text{Pb}\text{J}_2}{\text{L}} \times \frac{461.09}{\text{mol}} \text{Pb}\text{J}_2} = [0.709/L] \text{Pb}\text{J}_2}{\text{L}}$$

Optional: What's this in ppm (parts per million)? $700 \text{ ppm} \text{ PbI}_2$ For aqueous solutions, ppm translates to mg/L

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

Example:

$$A_{g}CI(s) \rightleftharpoons A_{g}^{\dagger}(c_{q}) + CI(a_{q}); K_{sp} = 1.6 \times 10^{-10}$$

$$Q = CA_{g}^{\dagger}[CI^{-}]$$

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

 \checkmark 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?

$$Q = [A_g^{\dagger}](cl^{-}) = (0.014)(0.00042) = 5.88 \times 10^{-8}$$

5.88 \text{to}^{-6} > 1.6 \text{to}^{-10}
 $Q > K_{SP}$, PRECIPITATION

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!

$$Ag(I(s) \rightleftharpoons Ag^{\dagger}(u_{\eta}) + CI^{\dagger}(u_{\eta}))$$

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(1(s) = Ag⁺(aq) + (1⁻(aq)); Ksp = (Ag⁺)(1⁻) = 1.6 × 10⁻¹⁰
Solve equilibrium...
Spe(1^es) [Imitial]
$$\triangle$$
 [Equilibrium]
Ag⁺ \bigcirc $+ \chi$ χ
(1⁻ \bigcirc $+ \chi$ χ
(x)(x) = 1.6 × 10⁻¹⁰
x² = 1.6 × 10⁻¹⁰
x = 1.2649 11064 × 10⁻⁵ M = (Ag⁺) = (Ag(1)) dissolved

$$\frac{1.26441064 \times 10^{-5} \text{ mol} A_g()}{L} \times \frac{143.35 \text{ g} A_g(1)}{\text{mol} A_g(1)} = 0.0018 \text{ g} A_g(1) = 1.8 \text{ ppm}}{L}$$

Ag(1(s) = Ag⁺ caq) + (1⁻ caq); Ksp = [Ag⁺] [(1⁻] = 1.6 × 10⁻¹⁰
Now solve for the AgCl dissolved in 0.10 M NaCl
Specifies [Initial]
$$\triangle$$
 [C(quilibrium]
Ag⁺ 0 + X X
(1⁻ 0.10 + X 0.10 + X
(x) (0.10 + x) = 1.6 × 10⁻¹⁰
Assume × cc 0.10
(x) (0.10) = 1.6 × 10⁻¹⁰
x = 1.6 × 10⁻⁹ = [Ag⁺] = [Ag(1] dissolved
Lossolved
Lossolved = 0.00023 ppm Agent

Compare: Solubility in DI water was 1.8 ppm. Solubility in 0.10 M NaCl (common ion, Cl-) is 0.00023 ppm.

Conclusion: The presence of a common ion GREATLY decreases solubility.

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