## Find Ksp Given Solubility - Question 1

Find the Ksp for calcium carbonate given that the **solubility** of calcium carbonate is found to be 6.934 p.p.m.(note p.p.m. = mg/L)

Must convert p.p.m. to mol/L first

$$\frac{6.934 \text{ mg CaCO}_3}{1 \text{ L}} \propto \frac{1 \text{ g}}{1000 \text{ mg}} \propto \frac{1 \text{ mol CaCO}_3}{100.09 \text{ g CaCO}_3} = \frac{6.93 \times 10^{-5} \text{ mol CaCO}_3}{1 \text{ L}}$$

$$CaCO_3(s) \rightleftharpoons Ca^{2+} + CO_3^{2-}$$

$$[Ca^{2+}] = 6.93 \ge 10^{-5} M$$
  
 $[CO_3^{2-}] = 6.93 \ge 10^{-5} M$ 

$$\begin{split} K_{sp} &= \ [Ca^{2+}][CO_3{}^{2-}] \\ &= \ (6.93 \ x \ 10^{-5})^2 \\ &= \ 4.8 \ x \ 10^{-9} \end{split}$$

Please note that solubility must be in mol/L in order to find ion concentrations that can be used to solve for a Ksp value.

## Find Solubility Ksp Given - Question 2

Find equilibrium ion concentrations for a saturated aluminum hydroxided solution and the mass of aluminum hydroxide in mg required to saturate 2500 mL of solution.

$$Al(OH)_3(s) \Longrightarrow Al^{3+} + 3 OH^{1-}$$
  $Ksp = 3 \ge 10^{-33}$ 

Let s represent the solubility of  $Al(OH)_3$ 

$$[Al^{3+}] = s$$
  
 $[OH^{1-}] = 3s$ 

$$\begin{split} \mathrm{K_{sp}} &= \ [\mathrm{Al^{3+}}] [\mathrm{OH^{1-}}]^3 \\ 3 \ge 10^{-33} &= \ (\mathrm{s}) (3\mathrm{s})^3 \\ 3 \ge 10^{-33} &= \ (\mathrm{s}) (27\mathrm{s^3}) \\ 3 \ge 10^{-33} &= \ 27\mathrm{s^4} \\ \mathrm{s} &= \ 3.25 \ge 10^{-9} \ \mathrm{mol/L} \end{split}$$

$$[Al^{3+}] = 3.25 \times 10^{-9} M$$
$$[OH^{1-}] = 3 \times (3.25 \times 10^{-9} M)$$
$$= 9.74 \times 10^{-9} M$$

$$2500 \text{ mL x} \frac{1 \text{ L}}{1000 \text{ mL}} \text{ x} \frac{3.25 \text{ x} 10^{-9} \text{ mol}}{1 \text{ L}} \text{ x} \frac{78.01 \text{ g} \text{ Al}(\text{OH})_3}{1 \text{ mol} \text{ Al}(\text{OH})_3} \text{ x} \frac{1000 \text{ mg}}{1 \text{ g}} = 6.34 \text{ mg} \text{ Al}(\text{OH})_3$$

## Find Maximum Ion Concentration Given a Fixed Ion Concentration - Question 3

Find the maximum concentration of  $Pb^{2+}$  in p.p.m. that can co-exist in equilibrium with a chloride ion concentration of 0.05 M.

$$Pb(Cl)_2 \rightleftharpoons Pb^{2+} + 2Cl^{1-} \qquad Ksp = 1.6 \ge 10^{-5}$$

$$\begin{split} K_{sp} &= \ [Pb^{2+}][Cl^{1-}]^2 \\ [Pb^{2+}] &= \frac{K_{sp}}{[Cl^{1-}]^2} \\ [Pb^{2+}] &= \frac{1.6 \ge 10^{-5}}{(0.05)^2} \\ [Pb^{2+}] &= 6.4 \ge 10^{-3} \ M \end{split}$$

$$\frac{6.4 \text{ x } 10^{-3} \text{ mol Pb}^{2+} \text{ L}}{1 \text{ L}} \text{ x } \frac{207.20 \text{ g Pb}^{2+}}{1 \text{ mol Pb}^{2+}} \text{ x } \frac{1000 \text{ mg}}{1 \text{ g}} = \frac{1326 \text{ mg Pb}^{2+}}{1 \text{ L}}$$

$$\therefore [Pb^{2+}] = 1326 \text{ p.p.m.}$$

## Find Ion Concentration in a Solution of a Soluble Salt (another fixed ion problem) - Question 4

what is the maximum possible concentration of magnesium ion in p.p.m. that can exist in a 1.0 M  $K_2 C_2 O_4$  solution

$$\begin{split} & \operatorname{K_2C_2O_4} \longrightarrow 2\operatorname{K^{1+}} + \operatorname{C_2O_4}^{2-} \\ & \operatorname{MgC_2O_4} \Longrightarrow \operatorname{Mg^{2+}} + \operatorname{C_2O_4}^{2-} \\ & \operatorname{MgC_2O_4} \Longrightarrow \operatorname{Mg^{2+}} + \operatorname{C_2O_4}^{2-} \\ & \operatorname{Ksp} = 8.6 \ge 10^{-5} \end{split}$$

$$K_{sp} = [Mg^{2+}][C_2O_4^{2-}]$$
$$[Mg^{2+}] = \frac{K_{sp}}{[C_2O_4^{2-}]^2}$$
$$[Mg^{2+}] = \frac{8.6 \times 10^{-5}}{1.0}$$
$$[Mg^{2+}] = 8.6 \times 10^{-5} M$$

 $\frac{8.6 \text{ x } 10^{-5} \text{ mol Mg}^{2+} \text{ L}}{1 \text{ L}} \text{ x } \frac{24.31 \text{ g Mg}^{2+}}{1 \text{ mol Mg}^{2+}} \text{ x } \frac{1000 \text{ mg}}{1 \text{ g}} = \frac{2.090 \text{ mg Mg}^{2+}}{1 \text{ L}}$ 

$$\therefore [Mg^{2+}] = 2.090 \text{ p.p.m.}$$

Ksp Problem : Common lon Effect Determin the solubility of lead iodide in a 0.5M KI solution common ion  $\begin{array}{cccc} KI \longrightarrow & K^{+} & + & I^{-} \\ \rho_{b}I_{2} & \Rightarrow & \rho_{b}^{2+} & + & 2I^{-} \\ \end{array}$ Let s represent the solubility  $\therefore [Pb^{2+}] = s$ [ I -] =25 + 0.5 contribution from KI Contribution from PbI2 Ksp = [Pb2+][I-]2  $7.1 \times 10^{-9} = 5(25+0.5)^2$ \* use methods to solve a culic or !! assume 25 <<< 0.5 .: 25 + 0.5 2 0.5  $7.1 \times 10^{-9} = 5(0.5)^2$ 5 = 2.84×10 -8 mol/ : 2,84,10 -8 <<< 0.5 (loso than 12) . assumption was valid Note the solubility will be lower than in

pure water

Complex Ksp problem - determine mass of ppte formed 400 mL of 0.0001 M Ba(NO3), is mixed with 500 mL of 0.000 3M Na2 CO3. What mass of ppte forms in mg? \* Na" + NO31are highly soluble  $Ba(NO_3)_2(5) \rightarrow Ba^{2+} + 2NO_3^{1-}$ and hence will Na2 (03 (5) > 2Na+ + (032not be involved ppte  $\Rightarrow Ba (O_3(5)) = Ba^{2+} + CO_3^{2-}$ in a pote Initial N.A.) 400mL + 500mL = 900mLτJ Total Volume: 0.9L Initial n=CV n=CV 0 n = 0.0001M2044 n=0,0003 M20.51 amount Let x represent n=4x10-5mol n=1.5x10-4mol Final the amount of 4×10-5-1 1.5×10-4-1 *I*X ∙ amount BaCO3 that forms Final 1.5 x 10-4-x 4×10-5-1 N. A. 0.9 0.9 [ ] extraneous  $K_{5p} = [Ba^{27}][CO_3^{27}]$ 1x= 1,78×10-4 mol (too big) "quadrations"  $\frac{4.9 \times 10^{-9}}{1} = \left(\frac{4 \times 10^{-5} - x}{0.9}\right) \left(\frac{1.5 \times 10^{-4} - x}{0.9}\right)$  $x = 1.137 \times 10^{-5} \text{ mol}$ (from table)

1.137×10 mol Baco3 × 197,34g Baco3 × 1000mg = 2.244 mg BaCO3 Imol Baco3 lg

Complexe Ksp problem - find Ksp value eg 400 mL of 0.0001M Ba (NO3)2 is mised with 500 m L of 0.0003M Na CO3. 2.244 mg of poto forms. Find KSp for the ppte produced  $B_{\alpha}(NO_{3})_{2} \rightarrow B_{\alpha}^{2+} + 2NO_{3}^{1-}$ Na, CO3 -> 2Na++ CO32-Ba (03(5) ₹ Ba<sup>2+</sup> + CO3<sup>2-</sup> Initial 400 mL + 500 mL = 900 mL. TJ. : Total Volume: 0.92 Initial n+CV n=CU 0 n=0.0001 Mx 0.4L n= 0.0003 Mx 0.51 Amount  $n = 4 \times 10^{-5} \text{ mol}$ n = 1.5x10-4mol Final 1.5×10-4 4x10 -5 1,137×10-5mol -1.137×107 -1,137x10-5 amount = 1,386x10 4 mol = 2.863×10-5 nul Final N.A. 3.181×10-5M 1.540×10-4M E]2.244 mg BaCO3 x 1g x 1 mol BaCO3 1000 mg 197.34 g BaCO3 = 1.137 × 10-5 mol Bacoz  $Ksp = [Ba^{2+}][CO_3^{2+}]$ Ksp = (3.181 × 10-5 \(1.540×10-4)  $k_{sp} = 4.9 \times 10^{-9}$ 

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