

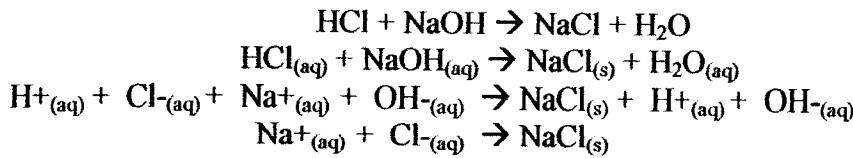
UNIT 6

EQUILIBRIUM APPLICATIONS

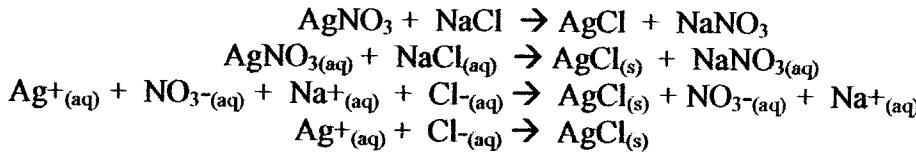
ANSWERS

Types of Equations

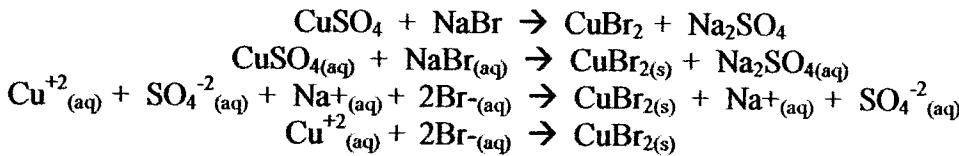
(1)



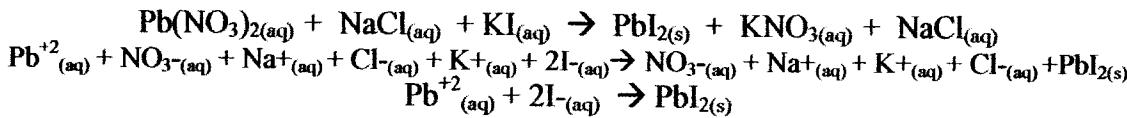
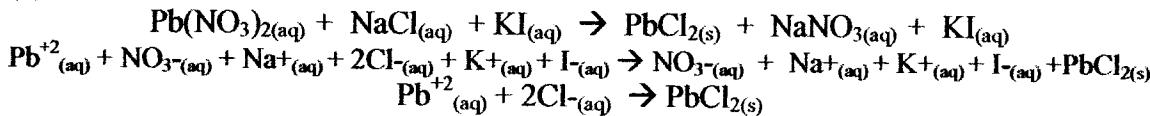
(2)



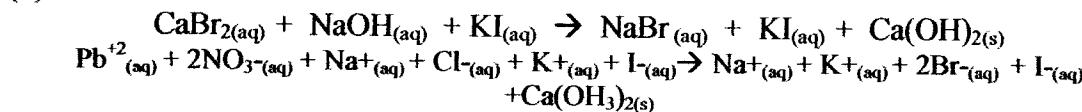
(3)



(4)



(5)



Solubility and Solubility Product Problems

① $120 \text{ mL} = 0.12 \text{ L}$

$$\frac{0.12 \text{ L} * 6 \text{ mol/L}}{2.0 \text{ L}} = 0.36 \text{ mol/L}$$

② $5.0 \text{ L} * 0.1 \text{ mol/L} = 0.5 \text{ mol}$ needed

$$\begin{aligned} 1 \text{ L} &\rightarrow 12 \text{ mol} & x &= 0.5 \text{ mol} / 12 \text{ mol} \\ x \text{ L} &\rightarrow 0.5 \text{ mol} & &= 0.04167 \text{ L} \end{aligned}$$

∴ you need to add 41.67 mL of 12M HCl

③ Original amounts:

$$\begin{aligned} 2 \text{ L} * 0.6 \text{ mol/L FeCl}_3 &\rightarrow 1.2 \text{ mol Fe}^{+3} \\ &\rightarrow 1.2 * 3 = 3.6 \text{ mol Cl}^- \end{aligned}$$

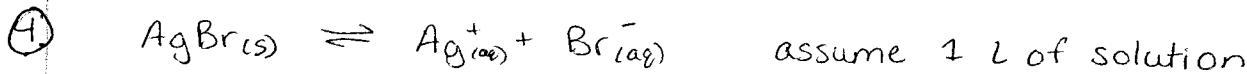
$$\begin{aligned} 1 \text{ L} * 0.9 \text{ mol/L BaCl}_2 &\rightarrow 0.9 \text{ mol Ba}^{+2} \\ &\rightarrow 0.9 * 2 = 1.8 \text{ mol Cl}^- \end{aligned}$$

...but now all of them are in 3 L ($2 \text{ L} + 1 \text{ L}$)...

$$\frac{1.2 \text{ mol Fe}^{+3}}{3 \text{ L}} = 0.4 \text{ M Fe}^{+3}$$

$$\frac{5.4 \text{ mol Cl}^-}{3 \text{ L}} = 1.8 \text{ M Cl}^-$$

$$\frac{0.9 \text{ mol Ba}^{+2}}{3 \text{ L}} = 0.3 \text{ M Ba}^{+2}$$



I	excess	0	0
E	excess -	8.8×10^{-7}	8.8×10^{-7}

$$= 244 \times 10^{-7} \text{ mol/L}$$

$$K_{sp} = [\text{Ag}^{+}] [\text{Br}^{-}] = (8.8 \times 10^{-7})^2 = 7.744 \times 10^{-14} (\text{mol/L})^2$$



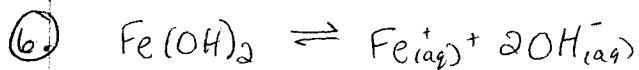
I	excess	0	0
E	excess - x	x	x

$$K_{sp} = [\text{Ca}^{+}] [\text{CO}_3^{-}]$$

$$4.8 \times 10^{-9} = x^2$$

$$x = 0.00006928$$

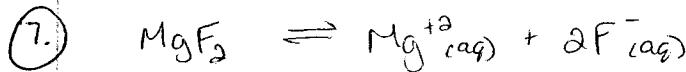
∴ the solubility of CaCO_3 is $6.928 \times 10^{-5} \text{ mol/L}$



I	excess	0	0
E	excess - 3.76×10^{-5}	1.67×10^{-5}	3.34×10^{-5}

$$\begin{cases} 1.3 \times 10^{-3} \text{ g} = x \text{ mol} \\ 89.85 \text{ g} = 1 \text{ mol} \\ \therefore \text{solubility is } 1.67 \times 10^{-5} \text{ mol/L} \end{cases}$$

$$\begin{aligned} K_{sp} &= [\text{Fe}^{+}] [\text{OH}^{-}]^2 \\ &= (1.67 \times 10^{-5})(3.34 \times 10^{-5})^2 \\ &= 1.86 \times 10^{-14} \text{ mol}^3/\text{L}^3 \end{aligned}$$



I	excess	0	0
E	excess - x	x	$2x$

$$K_{sp} = [\text{Mg}^{+2}] [\text{F}^{-}]^2$$

$$6.4 \times 10^{-9} = 4x^3$$

$$x = 1.17 \times 10^{-3} \text{ mol/L}$$

$$1.17 \times 10^{-3} \text{ mol} = x \text{ g}$$

$$1 \text{ mol} = 62.3 \text{ g}$$

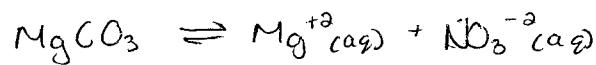
$$x = (1.17 \times 10^{-3})(62.3)$$

$$= 0.072891 \text{ g/L}$$

∴ the solubility of MgF_2 in g/L is 0.072891 g/L

$$\textcircled{8} \quad [\text{Mg}^{+2}_{(\text{aq})}] = \frac{0.04 \text{ L} * 8 \times 10^{-3} \text{ mol/L}}{0.1 \text{ L}} \\ = 0.0032 \text{ mol/L}$$

$$[\text{NO}_3^{-2}_{(\text{aq})}] = \frac{0.06 \text{ L} * 1 \times 10^{-3} \text{ mol/L}}{0.1 \text{ L}} \\ = 0.006 \text{ mol/L}$$

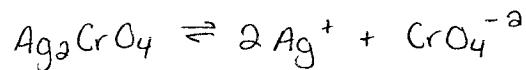


$$\text{but } K_{\text{sp}} = 2.6 \times 10^{-5} = [\text{Mg}][\text{NO}_3]$$

$$\begin{array}{ll} \text{L.S.} & \text{R.S.} \\ = 2.6 \times 10^{-5} & = (0.0032)(0.006) \\ & = 1.92 \times 10^{-7} \end{array}$$

Since RS < LS, no ppt forms.

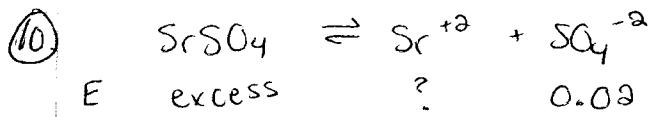
$$\textcircled{9} \quad [\text{Ag}] = \frac{0.025 \text{ L} * 4 \times 10^{-3} \text{ mol/L}}{0.1 \text{ L}} \quad [\text{CrO}_4^{-2}] = \frac{0.075 \text{ L} * 2 \times 10^{-4} \text{ mol/L}}{0.1 \text{ L}} \\ = 0.001 \text{ mol/L} \quad = 0.00015 \text{ mol/L}$$



$$\text{but } K_{\text{sp}} = 9.0 \times 10^{-13} = [\text{Ag}^+]^2 [\text{CrO}_4^{-2}]$$

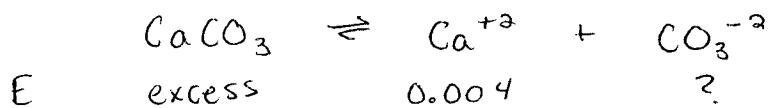
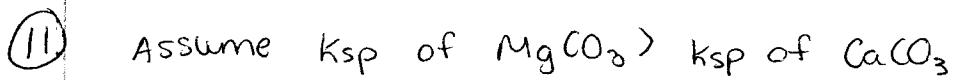
$$\text{L.S.} = 9.0 \times 10^{-13} \quad \text{R.S.} = (0.001)^2 (0.00015) \\ = 1.5 \times 10^{-10}$$

Since RS > LS, a ppt will form



$$\begin{aligned} K_{\text{sp}} &= [\text{Sr}^{+2}][\text{SO}_4^{-2}] \\ 7.6 \times 10^{-7} &= [\text{Sr}^{+2}](0.02) \\ [\text{Sr}^{+2}] &= 0.00038 \end{aligned}$$

\therefore the maximum $[\text{Sr}^{+2}]$ is $3.8 \times 10^{-5} \text{ mol/L}$ before a ppt occurs.



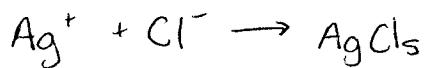
$$\begin{aligned} K_{\text{sp}} &= [\text{Ca}^{+2}][\text{CO}_3^{-2}] \\ 4.8 \times 10^{-9} &= (0.004)[\text{CO}_3^{-2}] \\ [\text{CO}_3^{-2}]_{\text{max}} &= 1.2 \times 10^{-6} \text{ mol/L} \end{aligned}$$

Since there are 5 L of water, the max amount of $[\text{CO}_3^{-2}]$ is $(5 * 1.2 \times 10^{-6} \text{ mol/L}) = 6 \times 10^{-6} \text{ mol/L}$

$$\begin{aligned} x_g &= 6 \times 10^{-6} \text{ mol} \\ 106 \text{ g} &= 1 \text{ mol} \\ x &= 6.36 \times 10^{-4} \text{ g} \end{aligned}$$

\therefore only $6.36 \times 10^{-4} \text{ g}$ of Na_2CO_3 can be added before a ppt forms.

(12)



$\therefore \# \text{ of moles of } \text{Ag}^+ = \# \text{ of moles of } \text{Cl}^-$

of moles of Ag^+ :

$$0.02364 \text{ L} * 0.1 \text{ mol/L}$$

$$= 0.002364 \text{ mol } \text{Ag}^+ \longrightarrow \therefore 0.002364 \text{ mol } \text{Cl}^-$$

$$[\text{Cl}^-] = \frac{0.002364 \text{ mol}}{0.125 \text{ L}} = 0.018912 \text{ mol/L}$$

\therefore the $[\text{Cl}^-]$ in the sample is 0.018912 mol/L

(13)

	$\text{PbI}_2 \rightleftharpoons \text{Pb}^{+3} + 2\text{I}^-$	x is solubility of PbI_3
I excess	0	0.1
E excess - x	x	$0.1 + 2x$

$$K_{\text{sp}} = [\text{Pb}^{+3}][\text{I}^-]^2$$

$$7.9 \times 10^{-9} = (x)(0.1 + 2x)^2 \leftarrow 2x^3 \text{ assume } x \text{ is small}$$

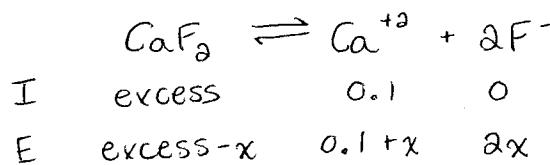
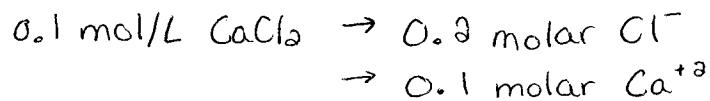
$$7.9 \times 10^{-9} = 0.01x \quad \text{use 5% rule to check}$$

$$x = 7.9 \times 10^{-7}$$

✓ check: $\frac{7.9 \times 10^{-7}}{0.1} \times 100\% = 7.9 \times 10^{-4}$

\therefore the solubility of PbI_3 in the I^- solution is $7.9 \times 10^{-7} \text{ mol/L}$

(4) CaCl_2 is soluble



$$K_{sp} = [\text{Ca}^{+2}][\text{F}^-]^2$$

$$3.9 \times 10^{-11} = (0.1 + x)(2x)^2 \leftarrow \text{assume } x \text{ is small}$$

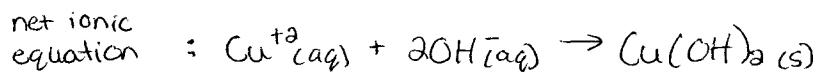
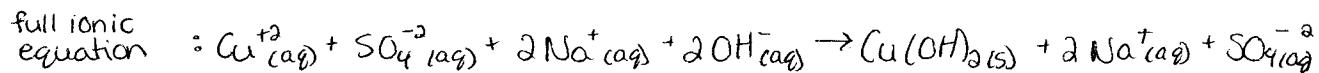
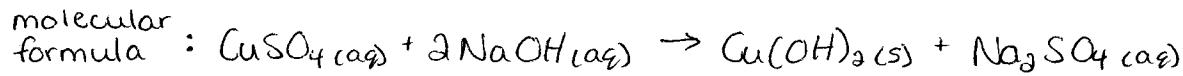
$$3.9 \times 10^{-11} = 0.4x^2$$

$$9.75 \times 10^{-11} = x^2$$

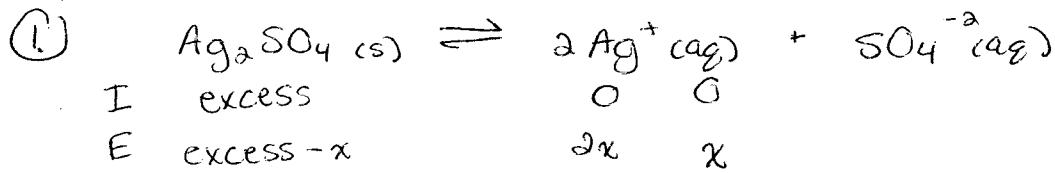
$$x = 9.874 \times 10^{-6} \text{ mol/L}$$

\therefore the maximum $[\text{F}^-]$ is $2x \Rightarrow 2(9.874 \times 10^{-6}) = 1.975 \times 10^{-5} \text{ mol/L}$

(5) Yes, a Cu(OH)_2 ppt will probably occur if the concentrations are high enough



K_{sp} Problems



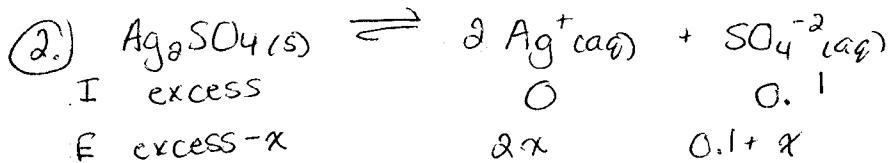
$$K_{\text{sp}} = 1.2 \times 10^{-5}$$

$$K_{\text{sp}} = [\text{Ag}^+(\text{aq})]^2 [\text{SO}_4^{2-}(\text{aq})]$$

$$1.2 \times 10^{-5} = [2x]^2 [x]$$

$$1.2 \times 10^{-5} = 4x^3$$

$$x = 0.01440$$



$$K_{\text{sp}} = 1.2 \times 10^{-5}$$

$$K_{\text{sp}} = [\text{Ag}^+(\text{aq})]^2 [\text{SO}_4^{2-}(\text{aq})]$$

$$1.2 \times 10^{-5} = [2x]^2 [0.1 + x] \leftarrow \text{assume } x \text{ is small compared to 0.1}$$

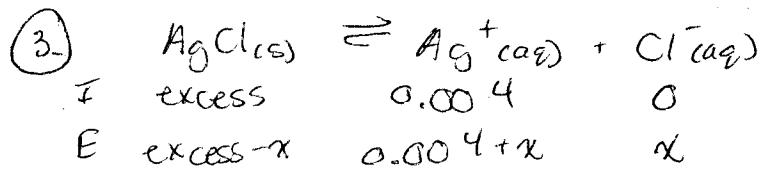
$$1.2 \times 10^{-5} = 0.4x^2$$

$$x = 0.005477$$

$$\frac{0.005477}{1 \text{ L}} = \frac{x}{0.5 \text{ L}}$$

$$x = 2.73 \times 10^{-3} \text{ mol/L}$$

∴ the solubility of Ag_2S is $2.73 \times 10^{-3} \text{ mol/L}$



$$K_{\text{sp}} = 1.8 \times 10^{-10}$$

$$K_{\text{sp}} = [\text{Ag}^+_{(\text{aq})}][\text{Cl}^-_{(\text{aq})}]$$

$$1.8 \times 10^{-10} = (0.004+x)(x) \leftarrow \text{assume } x \text{ is small compared to 0.004}$$

$$x = 4.5 \times 10^{-8} \times 0.2 \approx$$

$$= 9 \times 10^{-9}$$

$$\text{mm: } 22.98$$

$$+ \underline{35.04}$$

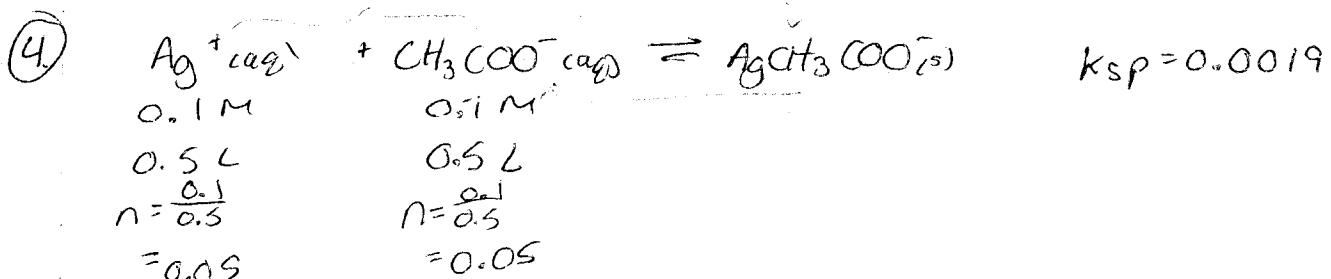
$$58.38$$

$$\text{m.} = n \times \text{mm}$$

$$= 9 \times 10^{-9} \times 58.38$$

$$= 5.25 \times 10^{-7}$$

\therefore the mass of table salt needed to precipitate the silver as silver chloride is 5.25×10^{-7} g



after mixing:

$$C = \frac{0.05}{1} \quad C = \frac{0.05}{1}$$

$$= 0.05 \text{ M} \quad = 0.05 \text{ M}$$

$$Q = [\text{Ag}^+_{(\text{aq})}][\text{CH}_3\text{COO}^-_{(\text{aq})}]$$

$$= (0.05)^2$$

$$= 0.0025$$

$$Q > K_{\text{sp}}$$

\therefore a precipitate forms

⑤ Ca^{+2} ion precipitates last because it has the greatest K_{sp} value (2.4×10^{-5})

⑥ Find the $[\text{SO}_4^{2-}]$

$$K_{\text{sp}} = [\text{Sr}^{+2}][\text{SO}_4^{2-}]$$

$$\begin{array}{l} 3.14 \times 10^{-3} \quad [\text{Sr}^{+2}] \\ \times 2.4 \times 10^{-5} \\ \hline 7.54 \times 10^{-8} \end{array}$$

Put this value into PbSO_4 to find $[\text{Pb}^{+2}]$

$$K_{\text{sp}} = [\text{Pb}^{+2}][\text{SO}_4^{2-}]$$

$$1.8 \times 10^{-5} = [\text{Pb}^{+2}][3.4 \times 10^{-6}]$$

$$[\text{Pb}^{+2}] = 5.29 \times 10^{-3}$$

$\therefore [\text{Pb}^{+2}]$ when Sr^{+2} begins to precipitate is $5.29 \times 10^{-3} \text{ mol/L}$



Bonds broken = energy absorbed

$$2(\text{C-C}) = 2(348) = 696$$

$$8(\text{C-H}) = 8(412) = 3296$$

$$5(\text{O=O}) = 5(498) = \underline{\underline{2490}} \\ + 6482$$

Bonds formed = energy released

$$6(\text{C=O}) = 6(740) = \underline{\underline{4440}}$$

$$8(\text{O-H}) = 8(464) = \underline{\underline{3712}} \\ - 8152$$

$$\therefore \text{net heat} = +6482 - 8152 \\ = -1670 \text{ KJ}$$

$$(8) \quad T = \frac{\Delta H}{\Delta S} = \frac{+44000}{118.89} \frac{\text{J/mol}}{\text{J/mol/K}} = +370.9 \text{ K} = 97.75^\circ\text{C}$$

$$\begin{aligned}\Delta H &= \sum \Delta H_f^\circ_{\text{prod}} - \sum \Delta H_f^\circ_{\text{react}} \\ &= 1(-241.8) - 1(-285.8) \\ &= +44 \text{ kJ} \\ &= +44000 \text{ J/mol}\end{aligned}$$

$$\begin{aligned}\Delta S &= \sum n S_{\text{prod}} - \sum n S_{\text{react}} \\ &= 1(188.84) - 1(69.95) \\ &= 118.89 \text{ J/mol/K}\end{aligned}$$