

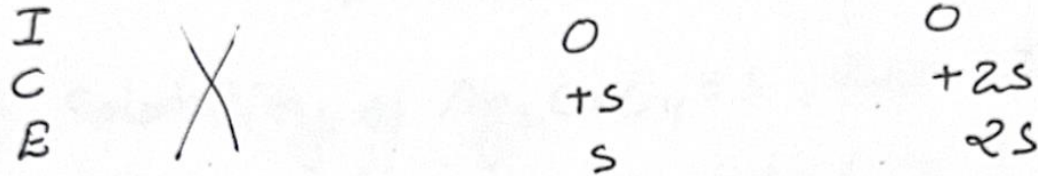
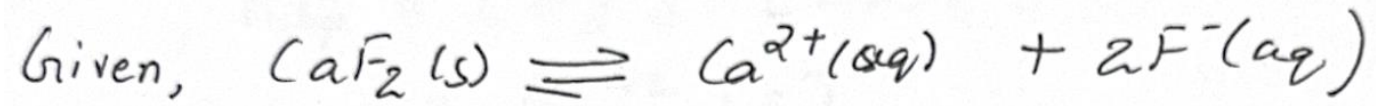
CHEM1128- EXAM 2 REVIEW

By Q-Center

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1. Calculate the solubility (in g/L) of CaF_2 in water from the known solubility product constant (3.4×10^{-11}).



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

$$K_{sp} = [s] [2s]^2$$

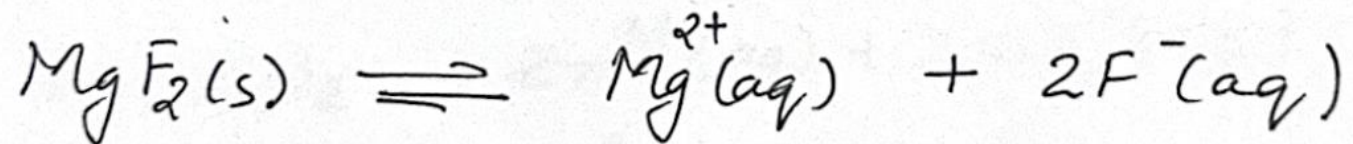
$$3.4 \times 10^{-11} = 4s^3$$

$$\Rightarrow s = 2.0 \times 10^{-4} \text{ mol of } \text{CaF}_2 \text{ in } 1 \text{ L of solution.}$$

$$\text{In g/L} \Rightarrow s = 2.0 \times 10^{-4} \frac{\text{mol}}{\text{L}} \times \frac{78.07 \text{ g}}{1 \text{ mol } \text{CaF}_2}$$

$$\boxed{s = 0.0156 \text{ g/L}}$$

2. The measured solubility of MgF_2 at 25°C is $2.6 \times 10^{-4} \text{ mol/L}$. Calculate the K_{sp} for MgF_2 .



$2.6 \times 10^{-4} \text{ mol}$ of MgF_2 dissolves.

1 mol MgF_2 splits to give 1 mol Mg^{2+} and 2 mol F^{-} .

Equilibrium concentration, $[\text{Mg}^{2+}] = 2.6 \times 10^{-4} \text{ M}$

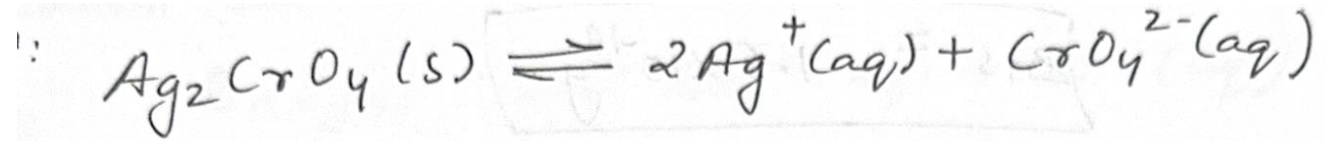
$$[\text{F}^{-}] = 2 \times (2.6 \times 10^{-4})$$

$$= 5.2 \times 10^{-4} \text{ M}$$

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

$$= [2.6 \times 10^{-4}] [5.2 \times 10^{-4}]^2 = \boxed{7.0 \times 10^{-11}}$$

3. How many grams of Silver Chromate can you dissolve in water to make one liter of solution? ($K_{sp} = 1 \times 10^{-12}$)



$$K_{sp} = [\text{Ag}^+]^2 [\text{CrO}_4^{2-}]$$
$$= [2s]^2 [s]$$

If solubility of $\text{Ag}_2\text{CrO}_4 = s$, then it forms $2s$ moles of Ag^+ and s moles of CrO_4^{2-} .

$$K_{sp} = [2s]^2 [s]$$

$$1 \times 10^{-12} = 4s^3$$

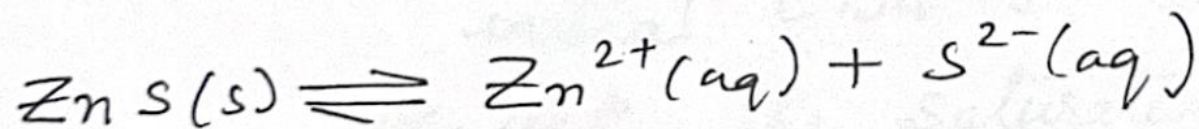
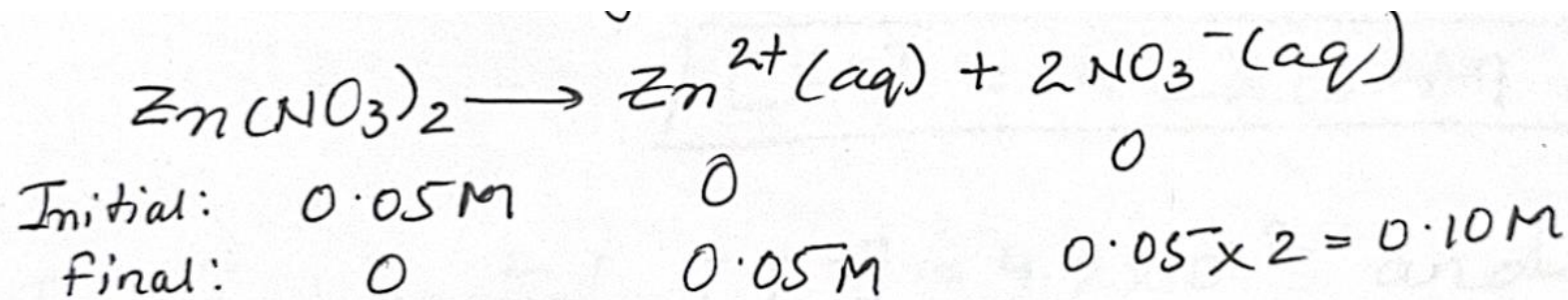
$$\boxed{6 \times 10^{-5} \text{ M} = s}$$

This means that 6×10^{-5} moles of Ag_2CrO_4 can be dissolved in enough water to make one liter of solution.

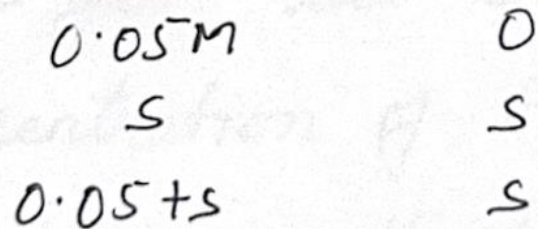
$$\frac{6 \times 10^{-5} \text{ mol}}{1 \text{ L}} \times \frac{331.7 \text{ g Ag}_2\text{CrO}_4}{1 \text{ mol Ag}_2\text{CrO}_4} = \frac{2 \times 10^{-2} \text{ g Ag}_2\text{CrO}_4}{1 \text{ L}}$$

4. What mass of ZnS ($K_{sp} = 2.5 \times 10^{-22}$) will dissolve in 300mL of 0.050M $\text{Zn}(\text{NO}_3)_2$?

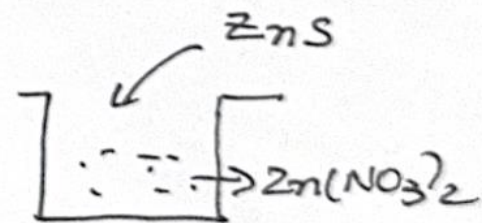
Solu:



- (I)
- (C)
- (E)

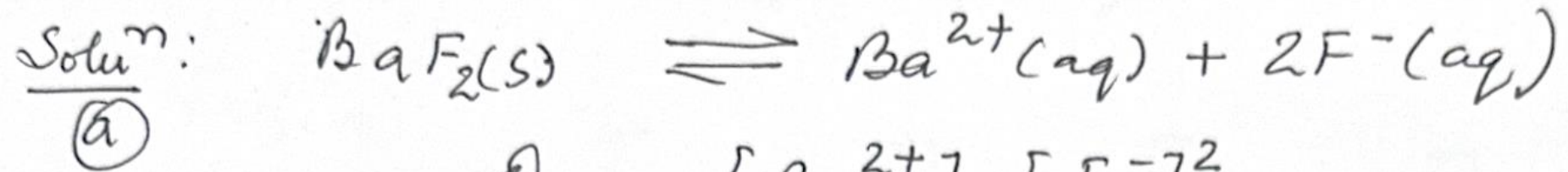


$$K_{sp} = [\text{Zn}^{2+}][\text{S}^{2-}] = (0.05 + s)(s) = 2.5 \times 10^{-22} \text{ mol/L}$$



5. A solution containing 0.020M Fluoride ions is added to a solution in which the original concentration of Barium ions is 1.0×10^{-4} M.

a) Will a precipitate form?



$$Q = [\text{Ba}^{2+}] [\text{F}^{-}]^2$$

$$Q = [1 \times 10^{-4}] [2 \times 10^{-2}]^2 = 4.0 \times 10^{-8}$$

$$K_{sp} = 1.8 \times 10^{-7}$$

Since, $K_{sp} > Q \Rightarrow$ No precipitation.

5. A solution containing 0.020M Fluoride ions is added to a solution in which the original concentration of Barium ions is 1.0×10^{-4} M.

b) What concentration of Fluoride ion is needed to get the saturated solution?

Soluⁿ: Saturated solution $\Rightarrow Q = K_{sp}$

We calculate $[F^-]$ in the K_{sp} expression by substituting values for the K_{sp} of BaF_2 and writing in the original $[Ba^{2+}]$ as equilibrium concentration.

$$K_{sp} = [Ba^{2+}] [F^-]^2$$

$$1.8 \times 10^{-7} = (1.0 \times 10^{-4}) [F^-]^2$$

$$[F^-]^2 = 1.8 \times 10^{-7} / 1.0 \times 10^{-4}$$

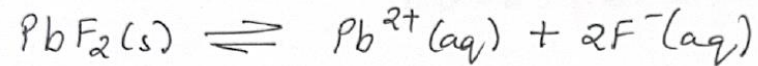
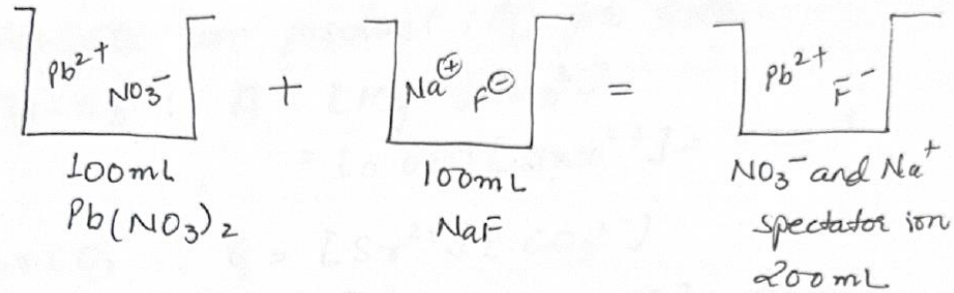
$$\boxed{[F^-] = 4.2 \times 10^{-2} M}$$

If the initial $[F^-] = 4.2 \times 10^{-2}$ and
initial $[Ba^{2+}] = 1.0 \times 10^{-4}$
we get the saturated solution.

5. A solution containing 0.020M Fluoride ions is added to a solution in which the original concentration of Barium ions is 1.0×10^{-4} M.
c) What concentration of Fluoride ion gives a precipitate?

Soluⁿ: Any solution with $[F^-] > 4.2 \times 10^{-2} M$ gives ppt.
Because then $Q > K_{sp}$.

6. A solution is prepared by mixing 100mL of $1.0 \times 10^{-2}M$ $Pb(NO_3)_2$ and 100mL of $1.0 \times 10^{-3}M$ NaF. Will $PbF_2(s)$ precipitate ($K_{sp} = 4 \times 10^{-8}$)?



Initial Concentration

$$[F^-] = 0.100\text{L} \times \frac{1.0 \times 10^{-3}\text{mol}}{1\text{L}} \times \frac{1}{0.200\text{L}} = 5.0 \times 10^{-4}M$$

(final volume)

$$[Pb^{2+}] = 0.100\text{L} \times \frac{1.0 \times 10^{-2}\text{mol}}{1\text{L}} \times \frac{1}{0.200\text{L}} = 5.0 \times 10^{-3}M$$

$$\begin{aligned}
 Q &= [Pb^{2+}] [F^-]^2 \\
 &= [5.0 \times 10^{-3}] [5.0 \times 10^{-4}]^2
 \end{aligned}$$

$$Q = 1.3 \times 10^{-9}$$

$$K_{sp} = 4 \times 10^{-8}$$

$\therefore K_{sp} > Q \Rightarrow$ NO precipitate.

7. A solution contains 0.02M Mg^{2+} and 0.02M Sr^{2+} . Sufficient carbonate is added so that the carbonate ion concentration is $2 \times 10^{-7}\text{M}$. Will a precipitate form and so, what is it?

Soluⁿ: Calculate ion product (Q) for both.

$$\text{MgCO}_3 : Q = [\text{Mg}^{2+}][\text{CO}_3^{2-}] \\ = [0.02][2 \times 10^{-7}] = 4 \times 10^{-9}$$

$$\text{SrCO}_3 : Q = [\text{Sr}^{2+}][\text{CO}_3^{2-}] \\ = [0.02][2 \times 10^{-7}] = 4 \times 10^{-9}$$

$$K_{sp}(\text{MgCO}_3) = 6.8 \times 10^{-6} > Q(\text{MgCO}_3) = 4 \times 10^{-9}$$

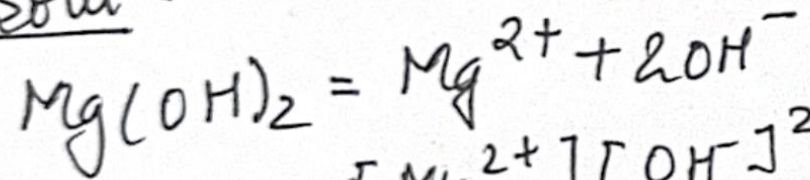
No PPT

$$K_{sp}(\text{SrCO}_3) = 5.6 \times 10^{-10} < Q(\text{SrCO}_3) = 4 \times 10^{-9}$$

PPT

8. A solution consists of 0.10 M MgCl_2 and 0.10 M ZnCl_2 . To separate the two cations from each other, solid NaOH is added. The volume of the solution does not change. Which salt will precipitate?

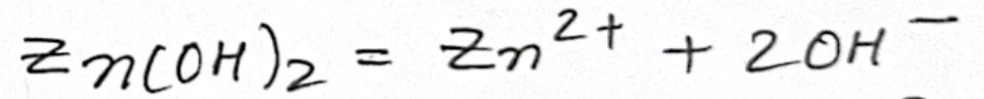
Solnⁿ:



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

$$6 \times 10^{-12} = [0.10][\text{OH}^-]^2$$

$$8 \times 10^{-6} = [\text{OH}^-]$$



$$K_{sp} = [\text{Zn}^{2+}][\text{OH}^-]^2$$

$$4 \times 10^{-17} = [0.1][\text{OH}^-]^2$$

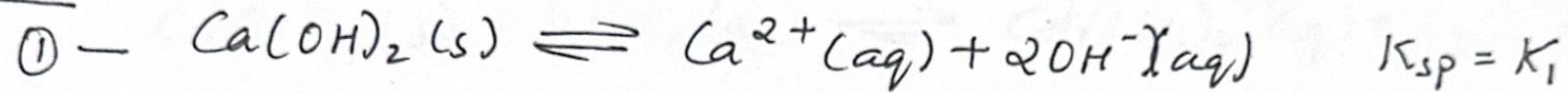
$$[\text{OH}^-] = 2 \times 10^{-8}$$

Small amount of NaOH is required to precipitate Zn^{2+} .

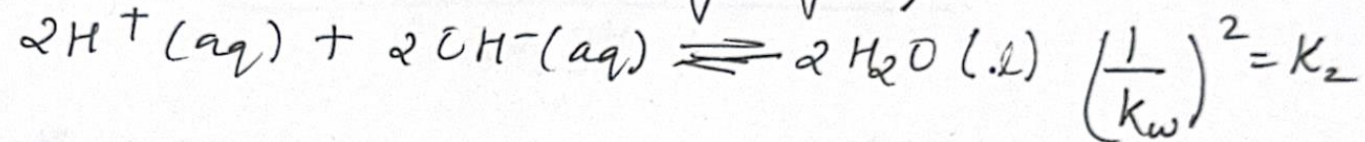
So, $\text{Zn}(\text{OH})_2$ precipitates first.

9. Write the overall equation and calculate K for the reaction in which Ca(OH)_2 is dissolved in HNO_3 . $K_{sp} = 4 \times 10^{-6}$ and $K_w = 1 \times 10^{-14}$.

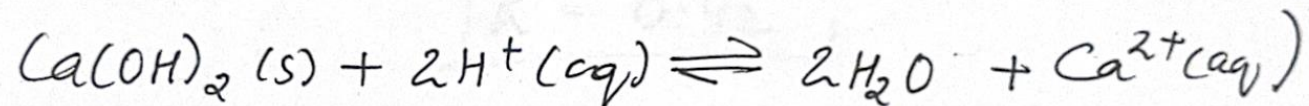
Solution:



$\textcircled{2}$ - flip and double it (multiply by 2)



Add $\textcircled{1}$ and $\textcircled{2}$,



$$K = K_1 * K_2$$

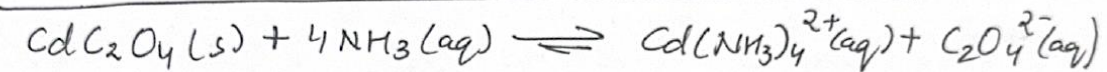
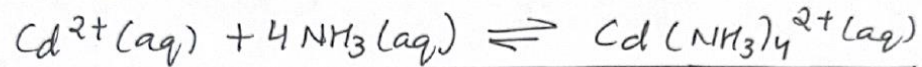
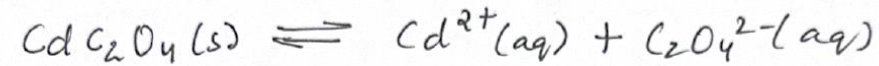
$$= K_{sp} * \left(\frac{1}{K_w}\right)^2$$

$$= (4 \times 10^{-6}) \left(\frac{1}{1 \times 10^{-14}}\right)^2$$

$$\boxed{K = 4.0 \times 10^{22}}$$

10. Cadmium(II) oxalate, CdC_2O_4 has a $K_{sp} = 1.5 \times 10^{-8}$. Cadmium ions readily combine with ammonia to form the complex ion $\text{Cd}(\text{NH}_3)_4^{2+}$ ($K_f = 2.8 \times 10^7$). Calculate the molar solubility of CdC_2O_4 in 0.010 M NH_3 .

Solution:



$$K = \frac{[\text{Cd}(\text{NH}_3)_4^{2+}][\text{C}_2\text{O}_4^{2-}]}{[\text{NH}_3]^4}$$

$$K = K_f * K_{sp} = (2.8 \times 10^7)(1.5 \times 10^{-8})$$

$$\boxed{K = 0.42}$$

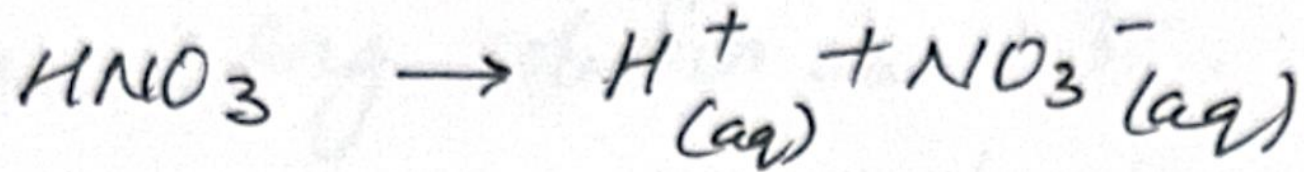
For 1 mol $\text{CdC}_2\text{O}_4 \rightleftharpoons 1 \text{ mol } \text{C}_2\text{O}_4^{2-} \rightleftharpoons 1 \text{ mol of } \text{Cd}(\text{NH}_3)_4^{2+}$ are formed.

$$s = [\text{C}_2\text{O}_4^{2-}] = [\text{Cd}(\text{NH}_3)_4^{2+}]$$

$$0.42 = \frac{(s)(s)}{(0.010)^4}$$

$$s = 6.5 \times 10^{-5} \text{ M}$$

1. Calculate the $[H^+]$ and the pH of a 0.25M solution of HNO_3 .



(strong acid)

0.25M

0.25M

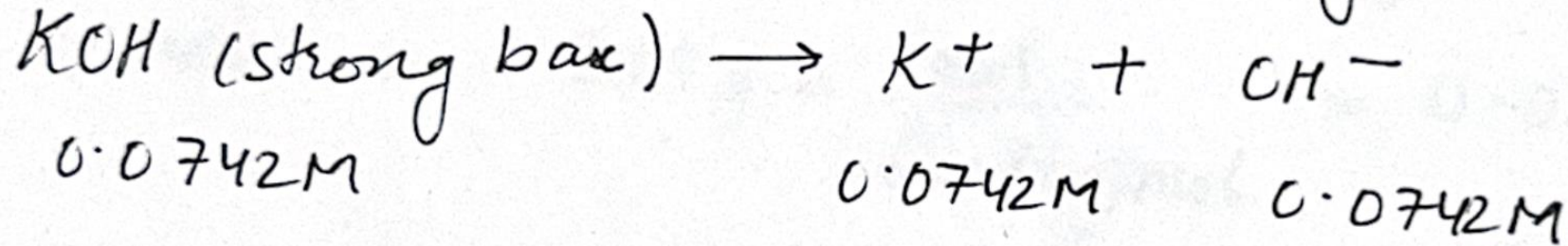
0.25M

$$pH = -\log[H^+]$$

$$= -\log[0.25] = \boxed{0.602}$$

2. Calculate the pH of a solution prepared by dissolving 2.08g KOH in enough water to make 500mL solution.

$$\text{Molarity of KOH} = \frac{2.08 \text{ g KOH}}{0.500 \text{ L}} \times \frac{1 \text{ mol KOH}}{56.1 \text{ g KOH}} = 0.0742 \text{ M}$$

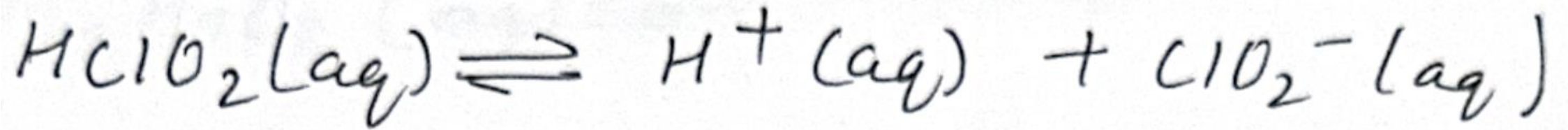


$$\text{pOH} = -\log[\text{OH}^-] = -\log(0.0742) = 1.130$$

$$\text{pH} = 14.000 - 1.130 = \boxed{12.870}$$

3. A solution of HClO_2 is prepared by dissolving 1.369g HClO_2 in enough water to make 100mL of solution. The pH of the resulting solution is 1.36.

a) Write the reaction for ionization of HClO_2 .



b) Write the expression for K_a .

$$K_a = \frac{[\text{H}^+][\text{ClO}_2^-]}{[\text{HClO}_2]}$$

3. A solution of HClO_2 is prepared by dissolving 1.369g HClO_2 in enough water to make 100mL of solution. The pH of the resulting solution is 1.36.

c) Calculate K_a .

Soln: molar mass of $\text{HClO}_2 = 68.46 \text{ g/mol}$

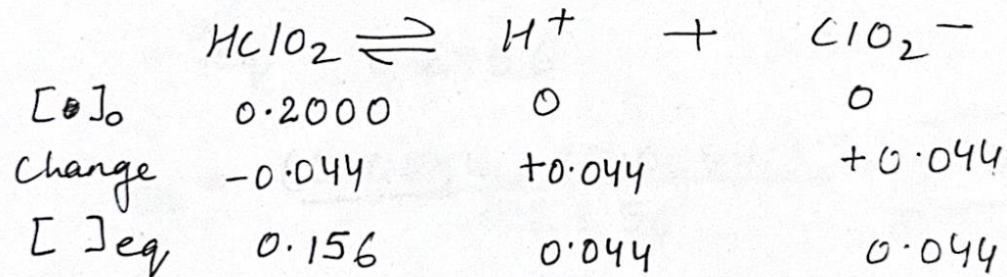
mass of $\text{HClO}_2 = \cancel{68.46 \text{ g}} \quad 1.369 \text{ g}$

moles = $1.369 \text{ g} \times \frac{1 \text{ mol}}{68.46 \text{ g/mol}} = 0.02000 \text{ mol}$

concentration (mol/L) = $\frac{0.02000}{0.1000} = 0.2000 \text{ M}$

$[\text{HClO}_2]_0 = 0.2000 \text{ M}$

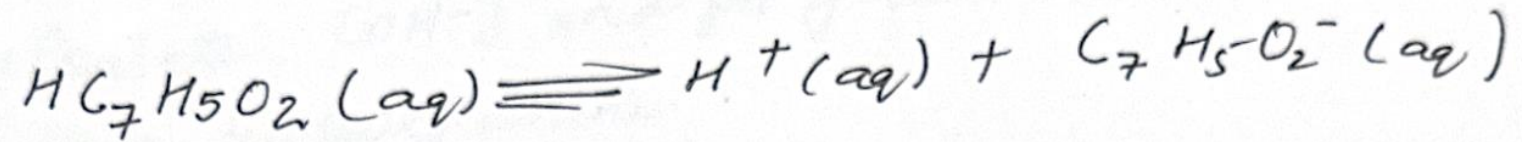
$\text{pH} = -\log [\text{H}^+] \Rightarrow [\text{H}^+]_{\text{eq}} = 10^{-1.36} = 0.044 \text{ M}$



$$K_a = \frac{[\text{H}^+] [\text{ClO}_2^-]}{[\text{HClO}_2]} = \frac{[0.044] [0.044]}{0.156}$$

$K_a = 0.012$

4. A 0.300M aqueous solution of Benzoic acid ($\text{HC}_7\text{H}_5\text{O}_2$) is 1.47% ionized.
a) Write the equation for the ionization.



- b) K_a expression for the ionization.

$$K_a = \frac{[\text{H}^+][\text{C}_7\text{H}_5\text{O}_2^-]}{[\text{HC}_7\text{H}_5\text{O}_2]}$$

4. A 0.300M aqueous solution of Benzoic acid ($\text{HC}_7\text{H}_5\text{O}_2$) is 1.47% ionized.
c) Calculate K_a for the ionization.

Solu: 1.47% ionization \Rightarrow 1.47% of $\text{HC}_7\text{H}_5\text{O}_2$ has been ionized.

$$1.47\% \text{ of } 0.300\text{M} = 0.3 \times \frac{1.47}{100} = 0.00441\text{M}$$

1 mol Benzoic acid \equiv 1 mol H^+ \equiv 1 mol benzoate ion

$$\Delta [\text{HC}_7\text{H}_5\text{O}_2] = \Delta [\text{H}^+] = \Delta [\text{C}_7\text{H}_5\text{O}_2^-] = 0.00441\text{M}$$

Table:

	$\text{HC}_7\text{H}_5\text{O}_2$	\rightleftharpoons	H^+	+	$\text{C}_7\text{H}_5\text{O}_2^-$
$[\]_0$	0.3000		0		0
Change	-0.00441		+0.00441		+0.00441
$[\]_{\text{eq}}$	0.2956		0.00441		0.00441

$$K_a = \frac{(0.00441)(0.00441)}{0.2956} = \boxed{6.58 \times 10^{-5}}$$

5. Hydroxylamine ionizes in water according to the rxn.



$$K_b = 9.1 \times 10^{-9}$$

Calculate the $[\text{OH}^-]$ and pH of one-liter solution containing 0.20M Hydroxylamine.

Soln:

$$\text{HONH}_2 + \text{H}_2\text{O} \rightleftharpoons \text{OH}^- + \text{HONH}_3^+$$

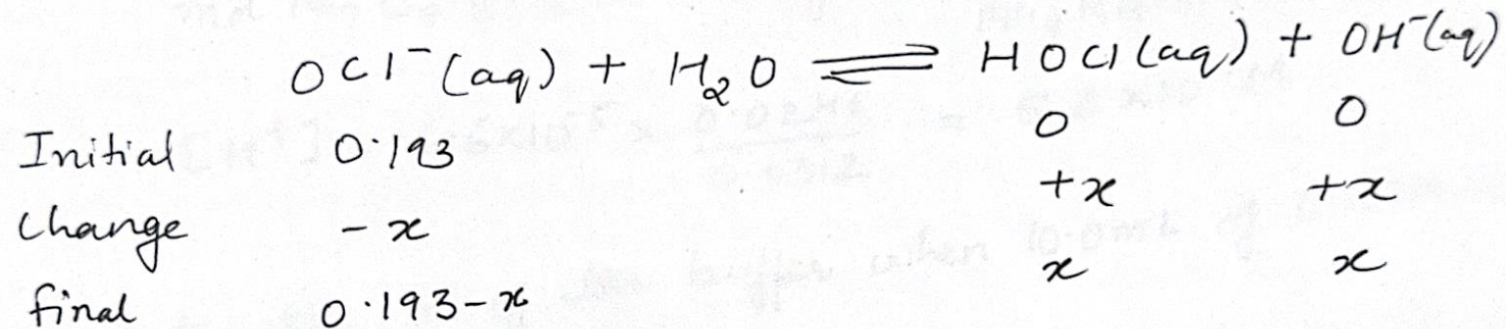
	0	0	0
$[\text{O}]_0$	0.2	0	0
Change	-x	+x	+x
$[\text{I}]_{\text{eq}}$	0.2-x	x	x

$$K_b = \frac{[\text{OH}^-][\text{HONH}_3^+]}{[\text{HONH}_2]}$$
$$9.1 \times 10^{-9} = \frac{(x)(x)}{0.20 - x} \quad \text{Ignore } -x$$
$$9.1 \times 10^{-9} = \frac{x^2}{0.2}$$
$$x = 4.3 \times 10^{-5} = [\text{OH}^-]$$
$$\text{pOH} = -\log [\text{OH}^-] = -\log [4.3 \times 10^{-5}] = 4.37$$
$$\text{pH} = 14 - \text{pOH} = 14 - 4.37 = \boxed{9.63}$$

6. Consider Sodium hypochlorite (NaOCl), household bleach. The OCl⁻ has $K_b = 3.6 \times 10^{-7}$. A solution is prepared by dissolving 12.0g of NaOCl (m.m= 74.45g/mol) in enough water to make 835mL of solution.

a) What is the pH of the solution?

Soln a) $[]_0 \text{ NaOCl} = \frac{12.0 \text{ g}}{0.835 \text{ L}} \times \frac{1 \text{ mol}}{74.45 \text{ g}} = 0.193 \text{ M}$



$$K_b = \frac{[\text{OH}^-][\text{HOCl}]}{[\text{OCl}^-]} \Rightarrow 3.6 \times 10^{-7} = \frac{(x)(x)}{0.193 - x}$$

Ignore $(-x)$, so, $x = 2.6 \times 10^{-4} \text{ M} = [\text{OH}^-]$

$$\text{p}[\text{H}^+] = \frac{1 \times 10^{-14}}{2.6 \times 10^{-4}} = 3.9 \times 10^{-11}$$

$\text{pH} = 10.41$

6. Consider Sodium hypochlorite (NaOCl), household bleach. The OCl^- has $K_b = 3.6 \times 10^{-7}$. A solution is prepared by dissolving 12.0g of NaOCl (m.m = 74.45g/mol) in enough water to make 835mL of solution.

b) Household bleach is 5.25% NaOCl by mass. Assuming that its density is 1.00g/mL is household bleach more alkaline than the prepared solution?

(b) Assuming 100g of bleach; we have 5.25g NaOCl in ~~100g~~
100 mL solution.

$$[\text{NaOCl}] = [\text{OCl}^-] = \frac{5.25\text{g}}{0.100\text{L}} \times \frac{1\text{mol}}{74.45\text{g}} = 0.705\text{M}$$

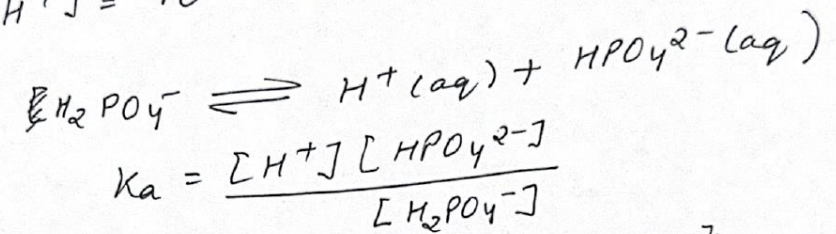
$$K_b = \frac{[\text{HOCl}][\text{OH}^-]}{[\text{OCl}^-]} \Rightarrow x = 5.0 \times 10^{-4}$$

~~% ionization~~ $[\text{H}^+] = \frac{1 \times 10^{-14}}{5 \times 10^{-4}} = 2.0 \times 10^{-11} \Rightarrow \boxed{\text{pH} = 10.70}$

BLEACH IS MORE ALKALINE

7. Suppose that you want to prepare an $\text{H}_2\text{PO}_4^- - \text{HPO}_4^{2-}$ buffer with a pH of 7. Taking the K_a of H_2PO_4^- to be 6.2×10^{-8} , how many grams of NaH_2PO_4 and Na_2HPO_4 should you add to water to make this buffer?

Solⁿ: $[\text{H}^+] = 10^{-7} = 1.0 \times 10^{-7} \text{ M}$



$$\frac{[\text{H}_2\text{PO}_4^-]}{[\text{HPO}_4^{2-}]} = \frac{[\text{H}^+]}{K_a} = \frac{1.0 \times 10^{-7}}{6.2 \times 10^{-8}} = 1.6$$

For every mole of weak base (HPO_4^{2-}), we need 1.6 mol of weak acid (H_2PO_4^-)

$$\text{mass of } \text{Na}_2\text{HPO}_4 = 1 \text{ mol} \times \frac{142 \text{ g}}{1 \text{ mol}} = 1.4 \times 10^2 \text{ g}$$

\therefore We need 1.6 mol weak acid, ~~so~~ we also need 1.6 mol NaH_2PO_4 .

$$\text{mass } \text{NaH}_2\text{PO}_4 = 1.6 \text{ mol} \times \frac{120 \text{ g}}{1 \text{ mol}} = 1.9 \times 10^2 \text{ g}$$

For buffer of $\text{pH} = 7 \Rightarrow$ Dissolve $1.4 \times 10^2 \text{ g}$ Na_2HPO_4 and $1.9 \times 10^2 \text{ g}$ of NaH_2PO_4 .

Doesn't matter how much water you use.