- 1. You have a 0.50 mM solution of sodium phosphate.
 - A) Determine the concentration of sodium ions and phosphate ions in this solution. Sodium phosphate is Na_3PO_4 , so $1 \mod Na_3PO_4 \rightarrow 3 \mod Na^+ + 1 \mod PO_4^{3^-}$. Therefore:

 $[Na^{+}] = 0.50 \text{ mM } Na_{3}PO_{4} \times \frac{1 \text{ M}}{1000 \text{ mM}} \times \frac{3 \text{ mol Na}^{+}}{1 \text{ mol Na}_{3}PO_{4}} = 1.5 \times 10^{-3} \text{ M}$ $[PO_{4}^{3-}] = 0.50 \text{ mM } Na_{3}PO_{4} \times \frac{1 \text{ M}}{1000 \text{ mM}} \times \frac{1 \text{ mol PO}_{4}^{3-}}{1 \text{ mol Na}_{3}PO_{4}} = 5.0 \times 10^{-4} \text{ M}$

B) If you want to dilute 1.0 mL of the 0.50 mM sodium phosphate solution to 5.0 μ M, how much water must be added to the original solution? We want to convert the 0.50 mM (5.0×10^{-4} M) solution to 5.0 μ M (5.0×10^{-6} M). We can set up the following expression, which tells us what volume of the 5.0 μ M solution would contain the same number of moles as 1.0 mL of the 0.50 mM solution.

 $\begin{array}{c} M_1V_1 = M_2V_2\\ (5.0 \times 10^{-4} \text{ M})(0.0010 \text{ L}) = (5.0 \times 10^{-6} \text{ M})V_2\\ V_2 = 0.10 \text{ L} (\text{ or } 100 \text{ mL})\\ \text{To get a } V_2 \text{ of 100 mL, we need to add: } 100 \text{ mL} - 1.0 \text{ mL} = 99 \text{ mL water.} \end{array}$

2. Consider the following four aqueous solutions:

 $KNO_3 Mg(NO_3)_2 Ca(NO_3)_2 NH_4NO_3$

A) Which of the aqueous solutions would produce a precipitate when added to an aqueous solution of sodium sulfate?

 $Ca(NO_3)_2$

B) Write the balanced net ionic equation for the precipitation reaction in part A. Start by writing the molecular equation: Ca(NO₃)₂ (aq) + Na₂SO₄ (aq) → CaSO₄ (s) + 2 NaNO₃ (aq)

Then, write the complete ionic equation: $2 \operatorname{Na}^+(aq) + 2 \operatorname{NO}_3^-(aq) + \operatorname{Ca}^{2+}(aq) + \operatorname{SO}_4^{2-}(aq) \rightarrow \operatorname{Ca}\operatorname{SO}_4(s) + 2 \operatorname{Na}^+(aq) + 2 \operatorname{NO}_3^-(aq)$

Finally, write the net ionic equation by eliminating all spectator ions: $Ca^{2+}(aq) + SO_4^{2-}(aq) \rightarrow CaSO_4(s)$

3. Determine the mass of solid precipitate formed when an excess lead(II) nitrate solution is mixed with 0.0800 mol sodium chloride.

Write the molecular equation, then the net ionic equation by eliminating all spectator ions: $Pb(NO_3)_2(aq) + 2 NaCl(aq) \rightarrow PbCl_2(s) + 2 NaNO_3(aq)$ $Pb^{2+}(aq) + 2 Cl^-(aq) \rightarrow PbCl_2(s)$

Because the Pb²⁺ is in excess, the Cl⁻ is limiting. Hence, $0.0800 \text{ mol NaCl} \times \frac{1 \text{ mol Cl}^{-}}{1 \text{ mol NaCl}} \times \frac{1 \text{ mol PbCl}_2}{2 \text{ mol Cl}^{-}} \times \frac{278.1 \text{ g}}{1 \text{ mol PbCl}_2} = 11.1 \text{ g PbCl}_2$ 4. Consider the following unbalanced reaction:

$$\operatorname{Cr}_{2}O_{7^{2^{-}}}(aq) + \operatorname{SO}_{3^{2^{-}}}(aq) \to \operatorname{Cr}^{3^{+}}(aq) + \operatorname{SO}_{4^{2^{-}}}(aq)$$

A) Determine the oxidation states for the specified atoms.

$Cr in Cr_2O_7^{2-} +6$	$S in SO_3^{2-} +4$	$S in SO_4^{2-} +6$
O in $Cr_2O_7^{2-}$ 2	O in SO_3^{2-} 2	O in SO_4^{2-}
B) Balance the above reaction using t1. Set up the two half-reactions:	he half-reaction method in ac $SO_3^{2^-} \rightarrow SO_4^{2^-}$ (reduct $Cr_2O_7^{2^-} \rightarrow Cr^{3^+}$ (oxidat)	cidic aqueous solution. ion: $S^{4+} \rightarrow S^{6+}$) ion: $Cr^{6+} \rightarrow Cr^{3+}$)
2. Balance elements (except O & H):	$\begin{array}{c} \mathrm{SO}_3^{2-} \rightarrow \mathrm{SO}_4^{2-} \\ \mathrm{Cr}_2 \mathrm{O}_7^{2-} \rightarrow 2 \ \mathrm{Cr}^{3+} \end{array}$	
3. Balance O atoms with H_2O :	$\begin{array}{l} \mathrm{SO_3^{2-} + H_2O \rightarrow SO_4^{2-}} \\ \mathrm{Cr_2O_7^{2-} \rightarrow 2 \ Cr^{3+} + 7 \ H_2O} \end{array}$	
4. Balance H atoms with H+:	$\begin{array}{l} \mathrm{SO_{3}^{2-} + H_{2}O \rightarrow SO_{4}^{2-} + 2 \ H^{+}} \\ \mathrm{Cr_{2}O_{7}^{2-} + 14 \ H^{+} \rightarrow 2 \ Cr^{3+} + 7 \ H_{2}O} \end{array}$	
5. Balance charges with e⁻:	$\begin{array}{l} \mathrm{SO_{3}^{2-} + H_{2}O \rightarrow SO_{4}^{2-} + 2 \ H^{+} + 2 \ e^{-} \\ \mathrm{Cr_{2}O_{7}^{2-} + 14 \ H^{+} + 6 \ e^{-} \rightarrow 2 \ \mathrm{Cr^{3+}} + 7 \ \mathrm{H_{2}O} \end{array}$	
5. Balance electrons:	$3 \times [SO_3^{2-} + H_2O \rightarrow SO_4^{2-} + 2H^+ + 2e^-]$ Cr ₂ O _{7²⁻ + 14H⁺ + 6e⁻ \rightarrow 2Cr³⁺ + 7H₂O}	
6. Sum up the two half-reactions:	$Cr_2O_7^{2-} + 3 SO_3^{2-} + 8 H^+ \rightarrow 2 Cr^{3+} + 3 SO_4^{2-} + 4 H_2O$	

5. A titration is performed where 48.0 mL of 1.00 M HCl is needed to react completely with 20.0 mL of a LiOH solution with unknown concentration.

What is the concentration of the unknown LiOH solution?

First, determine how many moles of HCl or H⁺ ions (mole ratio between HCl:H⁺ is 1:1) were required to reach the end-point:

$$n_{HCl} = 48.0 \text{ mL HCl} \times \frac{1 \text{ L}}{1000 \text{ mL}} \times \frac{1.00 \text{ mol HCl}}{1 \text{ L}} = 0.0480 \text{ mol HCl} \qquad n_{H^+} = 0.0480 \text{ mol H}^+$$

Second, recognize that the end-point occurs when $n_{acid} = n_{base}$, which means we have 0.0480 mol OH⁻ at the end-point. All of these OH⁻ ions come from solution B (diluted LiOH), which means we must have started with 0.0480 mol LiOH since the LiOH:OH⁻ mole ratio is 1:1.

Finally, we can now find the concentration of the LiOH solution:

$$M_{\text{LiOH}} = \frac{0.0480 \text{ mol LiOH}}{20.0 \text{ mL} \times \frac{1 \text{ L}}{1000 \text{ mL}}} = 2.40 \text{ M}$$

Note: An alternative solution would be to start with this equivalent expression:

$$\begin{pmatrix} \text{M}_{acid} \text{V}_{acid} = \text{M}_{base} \text{V}_{base} \\ \left(1.00 \ \frac{\text{mol HCl}}{1 \text{ L}}\right) \left(48.0 \text{ mL HCl} \times \frac{1 \text{ L}}{1000 \text{ mL}}\right) = \text{M}_{base} \left(20.0 \text{ mL HCl} \times \frac{1 \text{ L}}{1000 \text{ mL}}\right) \\ \mathbf{M}_{base} = \mathbf{2.40 \text{ M}}$$