Station 1:
It is found that $1.892 \times 10^{-13}$ grams of the compound cadmium (II) sulphide will dissolve in 350.0 mL of water to form a saturated solution. Using this data, calculate the value for the $\mathrm{K}_{\text {sp }}$ of CdS

$$
\begin{aligned}
& \operatorname{CdS} \underset{S}{ } C_{S} d^{2+}+S_{S}^{2-} \quad \text { (one source) } \\
& S=\frac{1.892 \times 10^{-13} \mathrm{~g}}{0.3500 \mathrm{~L}} \times \frac{1 \mathrm{~mol}}{144.5 \mathrm{~g}}=3.741 \times 10^{-15} \mathrm{~m} \\
& K_{s \rho}=\left[C d^{2+}\right]\left[S^{2-}\right]=S^{2} \\
& =\left(3.741 \times 10^{-15}\right)^{2} \quad=1.399 \times 10^{-29}
\end{aligned}
$$

How did you do?

| "I don't get it yet, <br> but I'm trying." | "I'm starting to get <br> it." | "I get it." | "I really get it and <br> can teach others <br> how to do it." |
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Station 2:
Up to 15.0 g of barium chloride can be dissolved in $2.5 \mathrm{~L}^{\text {of }} \mathrm{Al}_{2}\left(\mathrm{SO}_{4}\right)_{3}$ solution without forming a precipitate. Find the mass of aluminum in the solution.

$$
\begin{aligned}
& \frac{15.0 \mathrm{~g}}{2.5 \mathrm{~L}} \times \frac{1 \mathrm{~mol}}{208.3 \mathrm{~g}}=0.0288 \mathrm{M} \\
& =\left[\mathrm{Ba}^{2+}\right] \\
& \mathrm{Al}_{2}\left(\mathrm{SO}_{4}\right)_{3} \stackrel{\mathrm{r}}{ } \mathrm{Al}^{3+}+3 \mathrm{SO}_{4}{ }^{2-} \\
& 2.6 \times 10^{-9} \mathrm{M} \quad 3.8 \times 10^{-9} \mathrm{M} \\
& v_{\frac{2}{3}} \\
& \text { ?. } \mathrm{Al}^{3+}=2.5 \mathrm{~L} \times \frac{2.6 \times 10^{-9} \mathrm{~mol}}{1 \mathrm{~L}} \times \frac{27.0 \mathrm{~g}}{1 \mathrm{~mol}}=1.7 \times 10^{-7} \mathrm{~g}
\end{aligned}
$$

How did you do?

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Station 3:
Calculate the mass of $\mathrm{Na}_{2} \mathrm{CO}_{3}$ that must be added to 2.50 L of $0.00085 \mathrm{M} \mathrm{MgCl}_{2}$ in order to just start precipitation

$$
\begin{gathered}
\mathrm{Na}_{2} \mathrm{CO}_{3} \\
{[\mathrm{MgCl}]_{2} \mathrm{MgCl}_{2}} \\
=0.00085
\end{gathered}
$$

How did you do?

| "I don't get it yet,, <br> but I'm trying." | "I'm starting to get <br> it." | "I get it." | "I really get it and <br> can teach others <br> how to do it." |
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Station 4:
A sample of a saturated solution of $\mathrm{MgF}_{2}$ was evaporated and the following data table was constructed:

Mass of empty evaporating dish: 78.5418 g
Mass of evaporating dish and $\mathrm{MgF}_{2}$ residue after evaporation: 78.5434 g
Volume of saturated $\mathrm{MgF}_{2}: 100.00 \mathrm{~mL}=0.10000 \mathrm{~L}$
Temperature: $25.0^{\circ} \mathrm{C}$
Use this data to calculate the value of $\mathrm{K}_{\text {sp }}$ for $\mathrm{MgF}_{2}$ at $25^{\circ} \mathrm{C}$

$$
\begin{aligned}
& M g F_{2} \text { residue }=78.5434 \mathrm{~g}-78.5418 \mathrm{~g}=0.0016 \mathrm{~g} \\
& \\
& \frac{0.0016 \mathrm{~g}}{0.10000 \mathrm{~L}} \times \frac{1 \mathrm{~mol}}{62.3 \mathrm{~g}}=2.6 \times 10^{-4} \mathrm{M}=\mathrm{s}
\end{aligned}
$$

$$
M g F_{2} \gtrless \underset{(S)}{M g^{2+}}+2 F^{-}
$$

$$
K_{s p}=4 s^{3}=4\left(2.6 \times 10^{-4}\right)^{3}=6.8 \times 10^{-11}
$$

How did you do?

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Station 5:
What is the maximum mass of copper (II) chloride you can add 100.0L of a 0.025 M solution of sodium iodate without causing precipitation?

$$
\begin{aligned}
& \mathrm{CuCl}_{2}+\underset{\left[\mathrm{NaIO}_{3}\right]=}{2 \mathrm{NaIO}_{3}} \geqslant \mathrm{CH}_{2}\left(\mathrm{IO}_{3}\right)_{2}(\mathrm{ss})+2 \mathrm{NaCl} \\
& {\left[\mathrm{IO}_{3}{ }^{-}\right]=0.025 \mathrm{M} \longrightarrow \quad \mathrm{~K}_{s p}=\left[\mathrm{Cu}^{2+}\right]\left[10_{3}{ }^{-}\right]^{2}=6.9 \times 10^{-8}} \\
& {\left[\mathrm{Cu}^{2+}\right]=\frac{6.9 \times 10^{-8}}{(0.025)^{2}}=1.1 \times 10^{-4} \mathrm{M}} \\
& \mathrm{CuCl}_{2} \geq \mathrm{Cu}^{2+}+2 \mathrm{Cl}^{-} \\
& 1.1 \times 10^{-4} \sim^{1.1 \times 10^{-4}} \\
& ? \mathrm{~g} \mathrm{cu}^{2+}=100.0 \mathrm{~L} \times \frac{1.1 \times 10^{-4} \mathrm{~mol}}{1 \mathrm{~L}} \times \frac{134.5 \mathrm{~g}}{1 \mathrm{~mol}} \\
& =1.48 \mathrm{~g}
\end{aligned}
$$

How did you do?

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Station 6:
What is the maximum volume of 0.0350 M sodium sulphate solution required to obtain a saturated solution of strontium sulphate with 1.25 g of strontium nitrate?

$$
\begin{aligned}
& \mathrm{Na}_{2} \mathrm{SO}_{4}+\underset{\underline{\mathrm{Sc}}\left(\mathrm{NO}_{3}\right)_{2} \rightleftharpoons 2 \mathrm{NaNO}_{3}+\underset{\mathrm{SrSO}}{4}(\mathrm{~s})}{ } \\
& {\left[\mathrm{Na}_{2} \mathrm{SO}_{4}\right]=\left[\mathrm{SO}_{4}^{2-}\right]} \\
& =0.0350 \mathrm{M} \\
& K_{5 p}=\left[\mathrm{Sr}^{2+}\right]\left[\mathrm{SO}_{4}^{2-}\right]= \\
& 3.4 \times 10^{-7} \\
& \begin{array}{l}
\mathrm{Sr}\left(\mathrm{NO}_{3}\right)_{2} \approx \mathrm{Sc}^{2 t}+2 \mathrm{NO}_{3}^{-} . \\
9.7 \times 10^{-6}
\end{array} \\
& {\left[\mathrm{Sr}^{2+}\right]=\frac{3.4 \times 10^{-7}}{0.0350 \mathrm{~m}}} \\
& =9.7 \times 10^{-6} \mathrm{M}
\end{aligned}
$$

How did you do?

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Station 7:
Calculate the $\left[\mathrm{Ag}^{+}\right]$required to just start precipitation of $\mathrm{Ag}_{2} \mathrm{CO}_{3}$ in a 0.0030 M solution of $\left(\mathrm{NH}_{4}\right)_{2} \mathrm{CO}_{3}$

$$
\begin{aligned}
& \left(\mathrm{NH}_{4}\right)_{2} \mathrm{CO}_{3}+\mathrm{Ag}^{+}
\end{aligned} \begin{aligned}
& \begin{array}{l}
\left.\left.\mathrm{K} \mathrm{NH}_{4}\right)_{2} \mathrm{CO}_{3}\right]= \\
{\left[\mathrm{CO}_{3}^{2-}\right]}
\end{array} \\
& =0.0030 \mathrm{M} \\
& {\left[\mathrm{Ag}^{+}\right]=\sqrt{\frac{8.5 \times 10^{-12}}{2}\left[\mathrm{CO}_{3}^{2-}\right]}=8.5 \times 10^{-12}}
\end{aligned}
$$

How did you do?

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Station 8:
If 250.0 mL of $3.40 \times 10^{-4} \mathrm{M} \mathrm{Cu}\left(\mathrm{NO}_{3}\right)_{2}$ is mixed with 350.0 mL of $3.12 \times 10^{-4} \mathrm{M} \mathrm{KIO}{ }_{3}$, will a precipitate form?

$$
\begin{aligned}
& \underline{\underline{\mathrm{Cu}}\left(\mathrm{NO}_{3}\right)_{2}+\mathrm{KIO}} \\
& \rightleftharpoons\{\underbrace{\mathrm{Cu}\left(\mathrm{IO}_{3}\right)_{2(5)}}+\mathrm{KNO}_{3} \\
& C_{2}=\frac{\left(3.12 \times 10^{-4}\right)(350.0)}{(600.0)} \\
& =1.82 \times 10^{-4} \\
& =1.42 \times 10^{-4} \\
& =\left[\mathrm{Cu}^{2+}\right] \\
& =\left[\mathrm{IO}_{3}-\right] \\
& \text { Trial } K_{s p}=\left[\mathrm{Cu}^{2+}\right]\left[\mathrm{IO}_{3}\right]^{2} \\
& =\left(1.42 \times 10^{-4}\right)\left(1.82 \times 10^{-4}\right)^{2} \\
& \text { Actual isp } \\
& =6.9 \times 10^{-8} \\
& =4.70 \times 10^{-12}
\end{aligned}
$$

How did you do?

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