## Buffer Solutions

1. What is a buffer solution?

A buffer solution is one that is able to resist (or minimize) changes in pH . In order to identify if you have a buffer solution you are looking for a solution that contains a weak acid/base conjugate pair in solution.

For example:
If a solution contained both HOCl and NaOCl - it would be a buffer as it meets both criteria - a weak acid ( HOCl ) and it's conjugate base ( ${ }^{-} \mathrm{OCl}$ ). Keep in mind that the base will be added in the form of a conjugate salt, in this cae NaOCl . The $\mathrm{Na}^{+}$is just a spectator ion and its presence does not make or break the formation of the buffer.
2. When solving problems for a buffer solution you can bypass the ICE chart and use the Henderson-Hasselbalch equation instead. What is the Henderson-Hasselbalch equation?

$$
p H=p K_{a}+\log \frac{[b a s e]}{[a c i d]}
$$

Remember that " $p$ " means "- log of ..."
3. Identify the buffer?

When doing these problems you are looking to see

1. Are all substances in solution weak?
2. Is there a conjugate pair?
a. HCl and $\mathrm{NaClO}_{4}$

Answer the questions

1. No. HCl is a strong acid.

This automatically disqualifies this solution as being a buffer.
b. HClO and $\mathrm{LiClO}_{2}$

1. Yes. Both HClO and $\mathrm{ClO}_{2}{ }^{-}$are weak.
2. No. The conjugate base of HClO is $\mathrm{ClO}^{-}$not $\mathrm{ClO}_{2}^{-}$.

This disqualifies this solution as being a buffer.
c. HF and KF

1. Yes. Both HF and $\mathrm{F}^{-}$are weak.
2. Yes. The conjugate base of HF is $\mathrm{F}^{-}$

Because both criteria are met, this solution is a buffer.

## d. $\mathrm{CH}_{3} \mathrm{NH}_{2}$ and $\mathrm{CH}_{3} \mathrm{NH}_{3} \mathrm{Cl}$

1. Yes. Both $\mathrm{CH}_{3} \mathrm{NH}_{2}$ and $\mathrm{CH}_{3} \mathrm{NH}_{3}{ }^{+}$are weak.

2 Yes. The conjugate acid of $\mathrm{CH}_{3} \mathrm{NH}_{2}$ is $\mathrm{CH}_{3} \mathrm{NH}_{3}^{+}$

Because both criteria are met, this solution is a buffer.
4. Determine the pH of the following solutions
a. 1.00 L solution of $1.00 \mathrm{M} \mathrm{HNO}_{2}$ and $1.50 \mathrm{M} \mathrm{NaNO}_{2}$. $\left(K_{a}=4 \times 10^{-4}\right)$

This is a buffer solution. $\mathrm{HNO}_{2}$ is a weak acid and the salt contains its conjugate base $\mathrm{NO}_{2}{ }^{-}$. This means that we can is the Henderson-Hasselbalch equation to solve for the pH .

$$
\begin{gathered}
p H=p K_{a}+\log \frac{[b a s e]}{[a c i d]} \\
p H=-\log \left(4 \times 10^{-4}\right)+\log \frac{(1.50 \mathrm{M})}{(1.00 \mathrm{M})} \\
p H=3.57
\end{gathered}
$$

## b. 25.0 g of $\mathrm{NH}_{3}$ and $40.0 \mathrm{~g} \mathrm{NH}_{4} \mathrm{NO}_{3}$ in 501.0 mL of water. $\left(K_{a}=5.6 \times 10^{-10}\right)$

This is a buffer solution. $\mathrm{NH}_{3}$ is a weak base and the salt contains its conjugate acid $\mathrm{NH}_{4}{ }^{+}$. This means that we can use the Henderson-Hasselbalch equation to solve for the pH. We will just need to convert to the correct units first.

$$
\begin{aligned}
& 25.0 \mathrm{~g} \mathrm{NH} 33 \frac{1 \mathrm{~mol}}{17.04 \mathrm{~g}}=1.467 \mathrm{~mol} \mathrm{NH} 3 \\
& 40.0 \mathrm{~g} \mathrm{NH} \mathrm{~N}_{4} \mathrm{NO}_{3} \frac{1 \mathrm{~mol}}{80.06 \mathrm{~g}}=0.50 \mathrm{~mol} \mathrm{NH} \mathrm{NO}_{3}
\end{aligned}
$$

Because the $\mathrm{NO}_{3}{ }^{-}$is just a spectator ion, we are only really interested in the amount of conjugate base this salt provides, we convert from moles of $\mathrm{NH}_{4} \mathrm{NO}_{3}$ to moles of $\mathrm{NH}_{4}{ }^{+}$.

$$
\begin{gathered}
p H=-\log \left(5.6 \times 10^{-10}\right)+\log \frac{\left(\frac{1.47 \mathrm{~mol}}{0.501 \mathrm{~L}}\right)}{\left(\frac{0.50 \mathrm{~mol}}{0.501 \mathrm{~L}}\right)}=-\log \left(5.6 \times 10^{-10}\right)+\log \frac{(1.47 \mathrm{~mol})}{(0.50 \mathrm{~mol})} \\
p H=9.72
\end{gathered}
$$

As you can see the volumes cancel out, since the volumes will ultimately cancel out, it is valid to simply plug in moles rather than molarity when using this formula.

## 5. What is buffer capacity?

The amount of acid or base a buffer can absorb without having a significant change in pH .
6. Consider:
0.1 M NaF and 0.1M HF
1.0M NaF and 1.0M HF
0.01M NaF and 0.01M HF
a. Which has the highest pH ?

They all have the same pH because they all have the same $\mathrm{K}_{\mathrm{a}}$ value and ratio of [base]: [acid]. So in a buffer the pH doesn't depend on how concentrated the acid and base are; it depends on their ratios to one another.
b. Which has the highest buffer capacity?
1.0 M NaF and 1.0 M HF has the highest buffer capacity as it has the greatest concentration of acid/base.
7. Consider a solution that contains both $\mathrm{NH}_{3}$ and $\mathrm{NH}_{4} \mathrm{NO}_{3}$. If a solution has a $\mathrm{pH}=8.5$, calculate the ratio of

$$
\begin{gathered}
\frac{\left[\mathrm{NH}_{3}\right]}{\left[\mathrm{NH}_{4}{ }^{+}\right]} \\
\mathrm{K}_{\mathrm{a}}=5.6 \times 10^{-10}
\end{gathered}
$$

Looking at the solution you will notice that it contains a weak base and the salt contains its conjugate acid. That means that this solution is a buffer and we can use the Henderson - Hasselbalch equation.

$$
p H=p K_{a}+\log \frac{[b a s e]}{[a c i d]}
$$

All we have to do is plug the information provided into the equation and solve for the $\frac{[\text { basel }]}{[\text { acid }]}$.

$$
\begin{gathered}
8.5=-\log \left(5.6 \times 10^{-10}\right)+\log \frac{[\text { base }]}{\text { [acid] }]} \\
\frac{[\text { base }]}{[\text { acid] }]}=0.177
\end{gathered}
$$

8. What volumes of $0.22 \mathrm{M} \mathrm{CH}_{3} \mathrm{COOH}$ and $0.46 \mathrm{M} \mathrm{NaCH} 3 \mathrm{COO}^{2}$ must be mixed to prepare a 1.00 L solution buffered at $\mathrm{pH}=5.00$ ? $\left(\mathrm{K}_{\mathrm{a}}=\right.$ $1.76 \times 10^{-5}$ )

Once again we start by determining whether or not this solution is a buffer. As it contains $\mathrm{CH}_{3} \mathrm{COOH}$ ( a weak acid) and a salt that contains its conjugate base $\left(\mathrm{CH}_{3} \mathrm{COO}^{-}\right)$, this solution would be a buffer. This means that we can use the Henderson-Hasselbalch equation - plugging in all the relevant data.

$$
\begin{gathered}
p H=p K_{a}+\log \frac{[\text { base }]}{[\text { acid }]} \\
5.00=-\log \left(1.76 \times 10^{-5}\right)+\log \frac{[\text { base }]}{[\text { acid }]} \\
\frac{[\text { base }]}{[\text { [acid }]}=1.76
\end{gathered}
$$

because they are in the same volume of solution

$$
\frac{\text { moles base }}{\text { moles acid }}=1.76
$$

Remember:

$$
\text { Molarity } x \text { Volume }=\text { moles }
$$

Since the volume of the solution is equal to 1.00 L we know that the sum of the volume of base added plus the volume of acid added is equal to 1.00 L .

$$
\begin{gathered}
\text { Let } x=\text { volume of base added } \\
\text { Let }(1.00-x)=\text { volume of acid added. }
\end{gathered}
$$

Thus we can set up the following equation

$$
\begin{gathered}
\frac{(x)(0.46 \mathrm{M})}{(1.00-x)(0.22)}=1.76 \\
0.46 x=0.387-0.387 x \\
0.847 x=0.387 \\
x=0.457 L \rightarrow \text { volume of base added } \\
1.00-0.457=0.543 L \rightarrow \text { volume of acid added }
\end{gathered}
$$

