This is BCE\#17.

I recommend you print out this page and bring it to class. Click here to show a set of five BCE17 student responses randomly selected from all of the student responses thus far in a new window.

John, here are your responses to the BCE and the Expert's response.
In this BCE you will measure the $\mathbf{p H}$ of several aqueous solutions of acids and bases. You will watch a movie of a ph meter simulation developed at Iowa State University by Dr. Greenbowe and his students. Play the movie below. In the movie the concentrtion of each solution is set to 0.100 M . The beaker will be filled with $\mathbf{7 5} \mathrm{mLs}$ of each solution in turn. You are to record the measured pH and then to calculate the $\left[\mathrm{H}^{+}\right]$for each measurement in the table below the movie.


1a. Complete the $\mathbf{p H}$ column in the following table.

| Solution | $\mathbf{p H}$ | Equilibrium $\left[\mathrm{H}^{+}\right]$or $\left[\mathrm{OH}^{-}\right.$ <br> $]$ |
| :---: | :---: | :---: |
| 0.100 M HCl | 0.99 <br> 1.00 <br>  | $\left[\mathrm{H}^{+}\right]=1.0 \times 10^{-1} \mathrm{M}$ |
| $79 \%$ |  |  |


|  |  |  |
| :---: | :---: | :---: |
| 0.100 M HNO3 | $\begin{aligned} & 0.99 \\ & 1.00 \\ & 100 \% \end{aligned}$ | $\begin{gathered} 1 \mathrm{E}-1 \\ {\left[\mathrm{H}^{+}\right]=} \\ 1.0 \times 10^{-1} \mathrm{M} \\ 780 / 0 \end{gathered}$ |
| $\mathbf{0 . 1 0 0 ~ M ~ H 2 ~}{ }_{2} \mathrm{SO}_{4}$ | $\begin{aligned} & \hline 0.88 \\ & 0.88 \\ & 95 \% \\ & \hline \end{aligned}$ | $\left[\begin{array}{l} 0.131 \\ {\left[\mathrm{H}^{+}\right]=} \\ 1.31 \times 10^{-1} \mathrm{M} \\ 68 \% \end{array}\right.$ |
| $0.100 \mathrm{M} \mathrm{HC}_{2} \mathrm{H}_{3} \mathrm{O}_{2}$ | $\begin{aligned} & 2.87 \\ & 2.87 \\ & 98 \% \\ & \hline \end{aligned}$ | $\begin{gathered} 1.31 \mathrm{E}-3 \\ {\left[\mathrm{H}^{+}\right]=1.35 \times 10^{-3} \mathrm{M}} \\ 68 / / 0 \\ \hline \end{gathered}$ |
| 0.100 M NaOH | $\begin{gathered} 13 \\ 13.00 \\ 100 \% \\ \hline \end{gathered}$ | $\begin{gathered} 1 \mathrm{E}-1 \\ {\left[\mathrm{OH}^{-}\right]=1.0 \times 10^{-1} \mathrm{M}} \end{gathered}$ |
| $\mathbf{0 . 1 0 0} \mathrm{M} \mathrm{NH}_{3}$ | 11.12 <br> 11.12 <br> $95 \%$ | $\begin{gathered} 1.32 \mathrm{E}-3 \\ {\left[\mathrm{OH}^{-}\right]=1.32 \times 10^{-3} \mathrm{M}} \\ 47 \% \\ \hline \end{gathered}$ |

Sample Calculation of $\left[\mathrm{H}^{+}\right]$:
We know that $\mathrm{pH}=-\log \left[\mathrm{H}^{+}\right]$, to solve for the $\left[\mathrm{H}^{+}\right]$the negative sign must be moved to the side with pH
$-\mathbf{p H}=\log \left[\mathrm{H}^{+}\right]$
then both sides must be raised to the power of 10,
or $10^{-\mathrm{pH}}=10^{\log \left[\mathrm{H}^{+}\right]}$
but $10^{\log \left[\mathrm{H}^{+}\right]}=\left[\mathrm{H}^{+}\right]$
therefore $\left[\mathrm{H}^{+}\right]=10^{-\mathrm{pH}}$
To solve for the $\left[\mathrm{H}^{+}\right]$for 0.100 MHCl we use the measured pH of the solution $(\mathrm{pH}=1.00)$ in the equation $\left[\mathrm{H}^{+}\right]=10^{-\mathrm{pH}}$ to determine the equilibrium $\left[\mathrm{H}^{+}\right]$.
$\left[\mathrm{H}^{+}\right]=10^{-1}$ so $\left[\mathrm{H}^{+}\right]=1 \times 10^{-1} \mathrm{M}$
To solve for the $\left[\mathrm{H}^{+}\right]$for $0.100 \mathrm{M} \mathrm{H}_{2} \mathrm{SO}_{4}$ we use the measured pH of the solution ( $\mathrm{pH}=0.88$ ) in the equation $\left[\mathrm{H}^{+}\right]=10^{-\mathrm{pH}}$ to determine the equilibrium $\left[\mathrm{H}^{+}\right]$.
$\left[\mathrm{H}^{+}\right]=10^{-0.88}$ so $\left[\mathrm{H}^{+}\right]=1.3 \times 10^{-1} \mathrm{M}$
To solve for the $\left[\mathrm{H}^{+}\right]$for $0.100 \mathrm{M} \mathrm{HC}_{2} \mathrm{H}_{3} \mathrm{O}_{2}$ we use the measured pH of the solution $(\mathrm{pH}=2.87)$ in the equation $\left[\mathrm{H}^{+}\right]=10^{-\mathrm{pH}}$ to determine the equilibrium $\left[\mathrm{H}^{+}\right]$.
$\left[\mathrm{H}^{+}\right]=10^{-2.87}$ so $\left[\mathrm{H}^{+}\right]=1.3 \times 10^{-3} \mathrm{M}$
To solve for the $\left[\mathrm{OH}^{-}\right]$for 0.100 M NaOH we use the measured pH of the solution $(\mathrm{pH}=13.00)$ in the equation $\left[\mathrm{H}^{+}\right]=10^{-\mathrm{pH}}$ to determine the equilibrium $\left[\mathrm{H}^{+}\right]$.
$\left[\mathrm{H}^{+}\right]=10^{-13}$ so $\left[\mathrm{H}^{+}\right]=1.0 \times 10^{-13} \mathrm{M}$ and
and then recall that $1.0 \times 10^{-14}=\left[\mathrm{H}^{+}\right]\left[\mathrm{OH}^{-}\right]$
or $\left[\mathrm{OH}^{-}\right]=1.0 \times 10^{-14} /\left[\mathrm{H}^{+}\right]$
so for $0.100 \mathrm{M} \mathrm{NaOH}:\left[\mathrm{OH}^{-}\right]=1.0 \times 10^{-14} /\left[1.0 \times 10^{-13}\right]=1.0 \times 10^{-1} \mathrm{M}$
To solve for the $\left[\mathrm{OH}^{-}\right]$for $0.100 \mathrm{M} \mathrm{NH}_{3}$ we use the measured pH of the solution ( $\mathrm{pH}=11.12$ ) in the equation $\left[\mathrm{H}^{+}\right]=10^{-\mathrm{pH}}$ to determine the equilibrium $\left[\mathrm{H}^{+}\right]$.
$\left[\mathrm{H}^{+}\right]=10^{-11.12}$ so $\left[\mathrm{H}^{+}\right]=7.59 \times 10^{-12} \mathrm{M}$
$\left[\mathrm{OH}^{-}\right]=1.0 \times 10^{-14} /\left[\mathrm{H}^{+}\right]$
so for $0.100 \mathrm{M} \mathrm{NH}_{3}:\left[\mathrm{OH}^{-}\right]=1.0 \times 10^{-14} /\left[7.59 \times 10^{-12}\right]=1.3 \times 10^{-3} \mathrm{M}$
b) Complete the Equilibrium $\left[\mathrm{H}^{+}\right]$or $\left[\mathrm{OH}^{-}\right]$column by using the pH to calculate the $\left[\mathrm{H}^{+}\right]$or $\left[\mathrm{OH}^{-}\right]$for each acid or base.

2a. Each of the acids in the table above has the same concentration, 0.100 M . Did all of the acids have the same $\mathbf{p H}$ ? No No , two of the acids ( HCl and $\mathrm{HNO}_{3}$ ) had the same pH . However, both HCl and $\mathrm{HNO}_{3}$ had different pH compared to $\mathrm{H}_{2} \mathrm{SO}_{4}$ and $\mathrm{HC}_{2} \mathrm{H}_{3} \mathrm{O}_{2}$ (which differed between themselves).
b) What do you think might be an explanation for the experimental pHs you observed for this set of acids.
with the exception of HCl and HNO each acid dissociated to a different extent
HCl and $\mathrm{HNO}_{3}$ are both monoprotic strong acids. $\mathrm{H}_{2} \mathrm{SO}_{4}$ is a diprotic acid, so we would not be surprised that its pH differs from a monoprotic strong acid. However, the $\mathbf{p H}$ of $\mathrm{H}_{2} \mathrm{SO}_{4}$ is even more interesting. The $\left[\mathrm{H}^{+}\right]$suggests that the second proton is not completely dissociated. If it were we would have expected $a\left[\mathrm{H}^{+}\right]=\mathbf{0 . 2 0 0} \mathrm{M}$. So something interesting is going on there. Finally the $\mathbf{p H}$ of the $\mathbf{H C}_{\mathbf{2}} \mathbf{H}_{\mathbf{3}} \mathrm{O}_{\mathbf{2}}$ solution differs from everyone. $\mathrm{HC}_{2} \mathrm{H}_{3} \mathrm{O}_{2}$ is a monoprotic acid, but the calculated [ $\mathrm{H}^{+}$] suggests that the proton in $\mathrm{HC}_{2} \mathbf{H}_{3} \mathrm{O}_{2}$ does not completely dissociate as was observed in HCl and $\mathrm{HNO}_{3}$. This affirms that $\mathrm{HC}_{2} \mathrm{H}_{3} \mathrm{O}_{2}$ is a weak acid.

3a. Each of the bases in the table above has the same concentration, 0.100 M . Did all of the bases have the same $\mathbf{p H}$ ? No No , two of the bases $\left(\mathbf{N a O H}\right.$ and $\left.\mathrm{NH}_{3}\right)$ have different pHs .
b) What do you think might be an explanation for the experimental pHs you observed for this set of bases.
each base dissociated to a different extent
NaOH is a strong base, while $\mathrm{NH}_{3}$ is a weak base. A strong base completely dissociates. A weak base only partially dissociates.
2. Is there anything about the questions that you feel you do not understand? List your concerns/questions.
nothing
3. If there is one question you would like to have answered in lecture, what would that question be?
nothing

