## DATA SHEETS AND CALCULATIONS FOR ACIDS \& BASES

| Name |
| :--- |
| Partner's Name |
| Grade and Instructor Comments |

Part 1: Experimental Measurement—Determining a Numerical Value for $K_{w}$
Experimental pH of $\mathbf{0 . 0 1 0} \mathbf{~ M ~ N a O H ~ = ~}$ $\qquad$

## Part 1: Calculations_—Determining a Numerical Value for $\mathrm{K}_{\mathrm{w}}$

What is your measured pH ? $\qquad$
Based on the measured pH , what is the hydronium ion concentration? $\left[\mathrm{H}_{3} \mathrm{O}^{+}\right]=$ $\qquad$ M
Knowing that in a 0.010 M NaOH solution, $\left[\mathrm{OH}^{-}\right]=0.010 \mathrm{M}$, calculate a value for $\mathrm{K}_{\mathrm{w}}$ from your experimental value of the measured hydronium ion concentration and the known $\mathrm{OH}^{-}$concentration of the 0.010 M NaOH .

$$
\mathrm{K}_{\mathrm{w}}=\left[\mathrm{H}_{3} \mathrm{O}^{+}\right]\left[\mathrm{OH}^{-}\right]=
$$

and $\mathrm{pK}_{\mathrm{w}}=-\log \mathrm{K}_{\mathrm{w}}=$

Compare your results with the data taken from the scientific literature:

| $\mathbf{T}\left({ }^{\circ} \mathbf{C}\right)$ | $\mathbf{K}_{\mathbf{w}}$ | $\mathbf{p K}_{\mathbf{w}}$ |
| :--- | :--- | :--- |
| 20 | $0.68 \times 10^{-14}$ | 14.17 |
| 25 | $1.01 \times 10^{-14}$ | 14.00 |

$\mathrm{K}_{\mathrm{w}}$, as is the case for all equilibrium constants, varies with temperature. However, at a given temperature, $\mathrm{K}_{\mathrm{w}}$ is a constant for an aqueous solution. This means that at $25^{\circ} \mathrm{C}$ in any aqueous solution, regardless of solute, the value of $\mathrm{K}_{\mathrm{w}}\left\{=\left[\mathrm{H}_{3} \mathrm{O}^{+}\right]\left[\mathrm{OH}^{-}\right]\right\}$is $1.01 \times 10^{-14}$.

What is the pH of pure water at $25^{\circ} \mathrm{C}$ ?

## Part 2. Determination of $\mathrm{K}_{\mathrm{a}}$ for the Ammonium Ion-Experimental Measurements

## Data Table for Solutions of Ammonium Ion and Ammonia

Enter the experimental $p H$ values you determine in the lab in the last column. Complete the open areas of the table.

* Be very careful to rinse your glass electrode thoroughly with water before an after making this measurement.

| Solution | $\left[\mathrm{NH}_{4} \mathrm{Cl}\right], \mathrm{M}$ | $\left[\mathrm{NH}_{3}\right], \mathrm{M}$ | Solute Type <br> (Acid, Base, or Acid + Conj. Base | Enter Your <br> Experimental pH |
| :--- | :--- | :--- | :--- | :--- |
|  |  |  |  |  |
| A | 0.10 | 0 | Acid |  |
|  | 1.0 | 0 |  |  |
| C |  |  |  |  |
|  | 0.050 | 0.050 |  |  |
| D |  |  |  |  |

## Part 2: Calculating $\mathrm{K}_{\mathrm{a}}$ for the Ammonium Ion

a) Write the balanced, net ionic equation for the reaction of ammonium ion with water.
b) Write the equilibrium constant expression for $\mathrm{K}_{\mathrm{a}}$ for aqueous $\mathrm{NH}_{4}{ }^{+}$
c) Enter your experimental information for $\left[\mathrm{NH}_{3}\right]$ and $\left[\mathrm{NH}_{4}{ }^{+}\right]$(from the previous page) into the table below. Use your measured pH values for each solution (A-E) to calculate $\left[\mathrm{H}_{3} \mathrm{O}^{+}\right]$and enter these values in the table below. Finally, calculate $\mathrm{K}_{\mathrm{a}}$ for the ammonium ion and enter the values in the table. Show one representative calculation here.

Average calculated $\mathrm{K}_{\mathrm{a}}$ value $=$ $\qquad$ and average $\mathrm{pK}_{\mathrm{a}}=$ $\qquad$

| Solution | $\left[\mathrm{H}_{3} \mathrm{O}^{+}\right], \mathrm{M}$ | $\left[\mathrm{NH}_{3}\right], \mathrm{M}$ | $\left[\mathrm{NH}_{4}^{+}\right], \mathrm{M}$ | Calculated $\mathrm{K}_{\mathrm{a}}$ for $\mathrm{NH}_{4}^{+}$ | Calculated $\mathrm{pK}_{\mathrm{a}}$ for $\mathrm{NH}_{4}{ }^{+}$ |  |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| A |  |  |  |  |  |  |
| - |  |  |  |  |  |  |
| C |  |  |  |  |  |  |
| D |  |  |  |  |  |  |
|  |  |  |  |  | Not a required calcu- <br> lation | Not a required calcu- <br> lation |

## Part 3. Properties of $\mathrm{NH}_{4}{ }^{+} / \mathrm{NH}_{3}$ Buffer Solutions-Experimental Measurements

- Your experimental readings from page 70 are entered in the column labeled "Initial pH."
- NOTE: make a pH measurement on pure water before adding acid or base.
- Data for the pH after the addition of excess acid or base is entered into the columns marked " pH on adding $\mathrm{H}^{+}$" and " pH adding $\mathrm{OH}^{-\prime}$ " as appropriate.
- Fill in the boxes marked " $\Delta \mathrm{pH}$ " with calculated numbers.

| Solution | $\left[\mathrm{NH}_{4}^{+}\right], \mathrm{M}$ | $\left[\mathrm{NH}_{3}\right], \mathrm{M}$ | $\Delta \mathrm{pH}$ on adding $\mathrm{H}^{+}$ | pH on adding $\mathrm{H}^{+}$ | Initial pH | pH on adding $\mathrm{OH}^{-}$ | $\Delta \mathrm{pH}$ <br> adding $\mathrm{OH}^{-}$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| A | 0.10 | 0 |  |  |  |  |  |
| -B | 1.0 | 0 |  |  |  |  |  |
| C | 0.050 | 0.050 |  |  |  |  |  |
| D | 0.50 | 0.50 |  |  |  |  |  |
| E* | 0 | 0.10 |  |  |  |  |  |
| -Pure water |  |  |  |  |  |  |  |

## Part 3: Properties of $\mathrm{NH}_{4}{ }^{+} / \mathrm{NH}_{3}$ Buffer Solutions-Questions and Calculations

## Effect of Dilution on the $\mathbf{p H}$ of a Buffer:

If the solution is diluted more than 10 -fold, which solution - $1.0 \mathrm{M} \mathrm{NH}_{4} \mathrm{Cl}$ (solution B) or $0.50 \mathrm{M} \mathrm{NH}_{4} \mathrm{Cl}+$ $0.50 \mathrm{M} \mathrm{NH}_{3}$ (solution D) —does the pH change more? (Base your answer on the data in the "Initial pH" column on page 76.)

Explain, on the basis of the $\mathrm{K}_{\mathrm{a}}$ expression, why dilution has less effect on the pH of a buffer solution than on the pH of a solution containing only the acid as a solute (here $\mathrm{NH}_{4}{ }^{+}$).

## Effect of Added $\mathrm{H}_{3} \mathrm{O}^{+}$and $\mathrm{OH}^{-}$on a Buffer

Compare the values of $\Delta \mathrm{pH}$ (the changes in pH ) for solutions C and D (in the table on page 15) with those for solutions of the acid along (A and B) or conjugate base alone (E).
a) Which solutions show a buffering action?
b) Write balanced chemical equations for reactions that prevent larger changes in pH .

## Part 4. Titration Curves

The change in indicator color in an acid-base titration is a signal that the equivalence point is very near. Here you test two indicators that change colors in two different pH ranges.

| Indicator | Color in <br> Acidic Solution | Color in <br> Basic Solution |
| :--- | :--- | :--- |
| Bromcresol green |  |  |
| Phenolphthalein |  |  |

See Chemistry Ev Chemical Reactivity, page 872, Figure 18.10 for indicator colors.

## Titration Results: Option (a)- $\mathbf{H C l}+\mathbf{N a O H}$

The volumes of NaOH in the table are suggested values. Enter your actual volumes of NaOH used in the table (second and fifth columns). Enter experimental data in every cell in the table.

## Deductions from the HCl Titration Curve

| Suggested $\mathrm{V}_{\mathrm{NaOH}}, \mathrm{mL}$ | Actual <br> $V_{\mathrm{NaOH}}, \mathrm{mL}$ | Measured pH | Indicator Color | Suggested $\mathrm{V}_{\mathrm{NaOH}}, \mathrm{mL}$ | Actual <br> $V_{\mathrm{NaOH}^{\prime}} \mathrm{mL}$ | Measured pH | Indicator Color |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 0 |  |  |  | 22 |  |  |  |
| 3 |  |  |  | 23 |  |  |  |
| 6 |  |  |  | 24 |  |  |  |
| 8 |  |  |  | 24.5 |  |  |  |
| 10 |  |  |  | 25 |  |  |  |
| 12 |  |  |  | 25.5 |  |  |  |
| 14 |  |  |  | 26 |  |  |  |
| 16 |  |  |  | 28 |  |  |  |
| 18 |  |  |  | 30 |  |  |  |
| 20 |  |  |  | 32 |  |  |  |

a) Write a balanced, net ionic equation for the reaction that occurs during the titration.
b) How many equivalence points can you detect? Explain the connection between the number of equivalence points and the reaction occurring.

Be sure to attach to your report form a carefully drawn plot of pH versus volume of base added. Be sure your name appears on the plot.
c) CLEARLY LABEL on your titration curve the formulas for the species present at:
i) before adding NaOH
(ii) after 15 mL of NaOH has been added
ii) at the equivalence point
d) What is the connection between the indicator colors and the equivalence point?

## Titration Results: Option (b) $-\mathrm{H}_{3} \mathrm{PO}_{4}+\mathbf{N a O H}$

The volumes of NaOH in the table are suggested values. Enter your actual volumes of NaOH used in the table (second and fifth columns). Enter experimental data in every cell in the table.

| Suggested $\mathrm{V}_{\mathrm{NaOH}}, \mathrm{~mL}$ | Actual $\mathrm{V}_{\mathrm{NaO} \mathrm{H}}, \mathrm{~mL}$ | Measured pH | Indicator <br> Color | Suggested $\mathrm{V}_{\mathrm{NaO} \mathrm{H}^{\prime}} \mathrm{mL}$ | Actual <br> $\mathrm{V}_{\mathrm{NaOH},} \mathrm{mL}$ | Measured pH | Indicator <br> Color |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 0 |  |  |  | 19 |  |  |  |
| 3 |  |  |  | 19.5 |  |  |  |
| 6 |  |  |  | 20 |  |  |  |
| 8 |  |  |  | 20.5 |  |  |  |
| 9 |  |  |  | 21 |  |  |  |
| 9.5 |  |  |  | 22 |  |  |  |
| 10 |  |  |  | 23 |  |  |  |
| 10.5 |  |  |  | 24 |  |  |  |
| 11 |  |  |  | 26 |  |  |  |
| 12 |  |  |  | 28 |  |  |  |
| 13 |  |  |  | 30 |  |  |  |
| 15 |  |  |  | 35 |  |  |  |
| 17 |  |  |  | 40 |  |  |  |

## Deductions from the Phosphoric Acid Titration Curve

a) Write balanced, net ionic equations for the three possible successive reactions that occur during the titration.
1.
2.
3.
b) How many equivalence points can you detect?
c) CLEARLY LABEL on your titration curve the formulas for the species present at:
i) the equivalence points
ii) between the equivalence points
d) For which reaction or reactions (in a above) did you NOT see an equivalence point?
e) Write the equilibrium constant expressions for $\mathrm{K}_{1}$ and $\mathrm{K}_{2}$ of $\mathrm{H}_{3} \mathrm{PO}_{4}$
f) Determine the pK values for $\mathrm{H}_{3} \mathrm{PO}_{4}$ from your curve.

$$
\begin{array}{ll}
\mathrm{pK}_{1}\left(\mathrm{H}_{3} \mathrm{PO}_{4}\right)=\ldots & \text { and so } \mathrm{K}_{1}\left(\mathrm{H}_{3} \mathrm{PO}_{4}\right)= \\
\mathrm{pK}_{2}\left(\mathrm{H}_{3} \mathrm{PO}_{4}\right)= & \text { and so } \mathrm{K}_{2}\left(\mathrm{H}_{3} \mathrm{PO}_{4}\right)=
\end{array}
$$

$\qquad$
$\qquad$
g) Calculate the ratio of experimental $\mathrm{K}_{1}$ and $\mathrm{K}_{2}$ values:
$\mathrm{K}_{1} / \mathrm{K}_{2}=$ $\qquad$
h) What are the values of pH at the first and second equivalence point on your pH titration curve for phosphoric acid?
pH at 1st equivalence point $\qquad$
pH at 2 nd equivalence point $\qquad$
Explain why bromcresol green and phenolphthalein are suitable indicators for determining the concentration of a phosphoric acid solution. (See Figure 18.10 on page 872 of Chemistry $\mathcal{E}$ Chemical Reactivity.)

