

- **How Strong is a Weak Acid?**

- The equation for dissociation: $\text{HA (aq)} + \text{H}_2\text{O(l)} \rightleftharpoons \text{H}_3\text{O}^+(\text{aq}) + \text{A}^-(\text{aq})$
- The equilibrium constant for this reaction is called the **acid-dissociation constant**, K_a : $K_a = [\text{H}_3\text{O}^+][\text{A}^-] / [\text{HA}]$

- **Calculating K_a from the pH**

- The pH of a 0.10 M solution of formic acid, HCOOH , at 25 °C is 2.38. Calculate K_a for formic acid at this temperature.
 - $K_a = [\text{H}_3\text{O}^+][\text{HCOO}^-] / [\text{HCOOH}]$
 - Assume that $[\text{H}_3\text{O}^+] = [\text{HCOO}^-]$, $[\text{H}_3\text{O}^+] = 10^{-2.38} = 4.2 \times 10^{-3} \text{ M}$

	$[\text{HCOOH}], \text{ M}$	$[\text{H}_3\text{O}^+], \text{ M}$	$[\text{HCOO}^-], \text{ M}$
I	0.10	0	0
C	-4.2×10^{-3}	$+4.2 \times 10^{-3}$	$+4.2 \times 10^{-3}$
E	$0.10 - 4.2 \times 10^{-3} = 0.0958$	4.2×10^{-3}	4.2×10^{-3}

- $K_a = [4.2 \times 10^{-3}][4.2 \times 10^{-3}] / [0.0958] = 1.8 \times 10^{-4}$

- **Calculating Percent Ionization**

- Percent ionization = $([\text{H}_3\text{O}^+]_{\text{eq}} / [\text{HA}]_{\text{initial}}) \times 100$
- e.g. 1
 - $[\text{H}_3\text{O}^+]_{\text{eq}} = 4.2 \times 10^{-3} \text{ M}$
 - $[\text{HCOOH}]_{\text{initial}} = 0.10 \text{ M}$
 - Percent ionization = $(4.2 \times 10^{-3} / 0.10) \times 100 = 4.2\%$
- e.g. 2
 - In 0.100 M HF, the percent ionization is 8.1%. Calculate K_a .
 - $\text{HF (aq)} + \text{H}_2\text{O(l)} \rightleftharpoons \text{H}_3\text{O}^+(\text{aq}) + \text{F}^-(\text{aq})$
 - $K_a = [\text{H}_3\text{O}^+][\text{F}^-] / [\text{HF}]$
 - Amount dissociated: $(8.1/100) \times 0.100\text{M} = 8.1 \times 10^{-3}\text{M} = [\text{H}_3\text{O}^+][\text{F}^-]$
 - $K_a = (8.1 \times 10^{-3})^2 / (0.100 - 8.1 \times 10^{-3}) = 7.14 \times 10^{-4}$

- **Calculating pH Using K_a**

- Procedure:
 - Write the equilibrium constant expression from the ionization equilibrium.
 - Set up an ICE table to determine equilibrium concentrations as a function of x .
 - Substitute equilibrium concentrations into the expression for K_a and solve for x .
- e.g. What is the pH of 0.00100M acetic acid? $K_a = 1.8 \times 10^{-5}$
 - $\text{HA (aq)} + \text{H}_2\text{O(l)} \rightleftharpoons \text{H}_3\text{O}^+(\text{aq}) + \text{A}^-(\text{aq})$
 $K_a = [\text{H}_3\text{O}^+][\text{A}^-] / [\text{HA}]$
 $K_a = x^2 / 0.001 - x$ (ICE table)
 - option 1: quadratic function
 - option 2: assume $x \ll 0.00100$, $x^2 / 0.001 = 1.8 \times 10^{-5}$
 - $x = 1.34 \times 10^{-4}$, so pH = 3.873
 - Handy rule of thumb: If $[\text{HA}]_o \gg 100K_a$, assume $[\text{HA}]_{\text{eq}} = [\text{HA}]_o$
 - 5% rule: $x / [\text{HA}]_o \times 100 < 5\%$
 $x / [\text{HA}] \times 100 = 13\% > 5\%$ **(NO)**
 - option 3: successive approximation
 - First approx.: $x = 1.34 \times 10^{-4}$ ($x^2 / 0.001$)
 - Second approx.: $x = 1.25 \times 10^{-4}$ ($x^2 / (0.001 - 1.34 \times 10^{-4})$)
 - Third approx.: $x = 1.26 \times 10^{-4}$ ($x^2 / (0.001 - 1.25 \times 10^{-4})$)
 - **$x = 1.26 \times 10^{-4}$, pH = 3.900**

- **Polyprotic Acids**

- Polyprotic acids have more than one acidic proton.
- Generally, $K_{a1} \gg K_{a2} \gg K_{a3}$ so you can ignore the 2nd (and 3rd) ionizations.