

## Calculations Involving Acids

Section 8.4

Homework

Pg. 516 #2  
Pg. 520 #1, 2  
Pg. 521 #1, 2  
Pg. 524 #1, 2  
Pg. 525 #1-10a

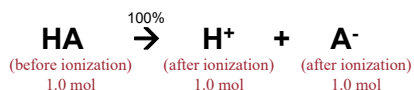
## Types of Calculations

- $K_a$
- pH
- % ionization

...or some combination of all of the above!

## Solutions of Strong Acids

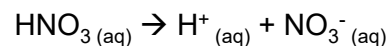
Complete ionization occurs:



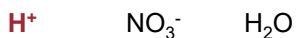
Therefore,  $[\text{HA}] = [\text{H}^+] = [\text{A}^-]$

**Example 1.** Find the  $[\text{H}^+]$ ,  $[\text{OH}^-]$ , pH and pOH of a 0.042 mol/L  $\text{HNO}_3(\text{aq})$  solution.

This reaction proceeds according to the equation:



Before getting started, consider the MAJOR entities in solution:



There are actually TWO processes that can contribute to  $[\text{H}^+]$ :

(1) ionization of  $\text{HNO}_3$   $K_a = \text{infinitely large}$   
(2) autoionization of water:  $\text{H}_2\text{O} \rightarrow \text{H}^+ + \text{OH}^-$   $K_w = 1.0 \times 10^{-14}$

Ignore process (2) because  
 $K_a$  of  $\text{HNO}_3$  is so much larger  
 $\therefore \text{HNO}_3$  is a much stronger acid;  
contribution of  $\text{H}_2\text{O}$  is negligible

**Example 1.** Find the  $[\text{H}^+]$ ,  $[\text{OH}^-]$ , pH and pOH of a 0.042 mol/L  $\text{HNO}_3(\text{aq})$  solution.

Strategy for strong acid calculations:

1. Assume that  $[\text{H}^+] = [\text{HA}]$
2. Perform necessary calculations using  $[\text{H}^+]$

**Example 1.** Find the  $[H^+]$ ,  $[OH^-]$ , pH and pOH of a 0.042 mol/L  $HNO_3$  (aq) solution.

**Strategy for strong acid calculations:**

1. Assume that  $[H^+] = [HA]$
2. Perform necessary calculations using  $[H^+]$

1 Find  $[H^+]$

Since  $HNO_3$  is a strong acid,  
 $[H^+] = [HNO_3] = 0.042\text{ M}$

3 Find pOH

$$\begin{aligned} \text{pH} + \text{pOH} &= 14.00 \\ \text{pOH} &= 14.00 - \text{pH} \\ \text{pOH} &= 14.00 - 1.38 \\ \text{pOH} &= 12.62 \end{aligned}$$

2 Find pH

$$\begin{aligned} \text{pH} &= -\log[H^+] = -\log(0.042) \\ \text{pH} &= 1.38 \end{aligned}$$

4 Find  $[OH^-]$

$$\begin{aligned} [OH^-] &= 10^{-\text{pOH}} \\ &= 10^{-12.62} \\ [OH^-] &= 2.4 \times 10^{-13} \text{ mol/L} \end{aligned}$$

**Practice.**

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1. Find  $[OH^-]$  of a 0.0700 mol/L HCl solution. Ans:  $1.43 \times 10^{-13}\text{ M}$
2. A 2.00-L HBr solution contains 0.070 mol of acid. Find pH and pOH. Ans: pH = 1.46, pOH = 12.54

## Solutions of Weak Acids

Weak acids only partially ionize in water.

- **Percent ionization** describes how much of the original acid ionizes to produce  $H^+$ .

For the general weak acid ionization reaction  $HA \rightleftharpoons H^+ + A$

$$\% \text{ ionization} = \frac{[H^+]}{[HA]_0} \times 100\% \quad \text{where } [H^+] \text{ is the concentration of ionized acid, and } [HA]_0 \text{ is the initial acid concentration.}$$

### Example 2. Finding % Ionization Using pH

Calculate the percent ionization of propanoic acid,  $HC_3H_5O_2$ , if a 0.050 mol/L solution has a pH of 2.78.

- 1 Write the ionization equation  $HC_3H_5O_2 \rightleftharpoons H^+ + C_3H_5O_2^-$
- 2 Use pH to find  $[H^+]$ 

$$[H^+] = 10^{-2.78}$$

$$[H^+] = 1.7 \times 10^{-3} \text{ mol/L}$$

← the quantity of  $H^+$  from the ionization of  $HC_3H_5O_2$ .
- 3 Calculate % ionization
 
$$\begin{aligned} \% \text{ ionization} &= \frac{[H^+]}{[HC_3H_5O_2]_0} \times 100\% \\ &= \frac{1.7 \times 10^{-3}}{0.050} \times 100\% \\ \% \text{ ionization} &= 3.3\% \end{aligned}$$

### Example 3. Finding $K_a$ using % Ionization

Calculate the  $K_a$  of hydrofluoric acid, HF, if a 0.100 mol/L solution at equilibrium has a percent ionization of 7.8%.

- 1 Write the ionization equation  $HF \xrightarrow{7.8\%} H^+ + F^-$
- 2 Write  $K_a$  expression
 
$$K_a = \frac{[H^+][F^-]}{[HF]}$$

Requires all concentrations at equilibrium

3 Set up ICE table

	HF	$H^+$	$F^-$
I	0.100	0	0
C	-x	+x	+x
E	0.100 - x	x	x

Given: % ionization = 7.8%

RTE:  $K_a$

	HF	$H^+$	$F^-$
I	0.100	0	0
E	0.100 - x	x	x

0.0922 mol/L    0.0078 mol/L    0.0078 mol/L

- 4 Use % ionization to find  $[H^+]$  and  $[F^-]$ 

$$\begin{aligned} x &= 0.078 \times 0.100 \text{ mol/L} \\ x &= 0.0078 \text{ mol/L} = [H^+] = [F^-] \end{aligned}$$
- 5 Use x to find  $[HF]$ 

$$[HF] = 0.100 - x = 0.0922 \text{ mol/L}$$
- 6 Plug values into  $K_a$ 

$$K_a = \frac{[H^+][F^-]}{[HF]} = \frac{(0.0078)(0.0078)}{(0.0922)}$$

$$K_a = 6.6 \times 10^{-4}$$

### Example 4. Calculating the pH of a weak acid solution

Calculate the hydrogen ion concentration and pH of a 0.10 mol/L acetic acid solution.  $K_a$  is  $1.8 \times 10^{-5}$ .

- Write the ionization equation  

$$\text{HC}_2\text{H}_3\text{O}_2 \rightleftharpoons \text{H}^+ + \text{C}_2\text{H}_3\text{O}_2^-$$
- Consider all major entities, and reactions that contribute to pH.  
 Major entities in solution:  
 $\text{HC}_2\text{H}_3\text{O}_2 \quad \text{H}^+ \quad \text{C}_2\text{H}_3\text{O}_2^- \quad \text{H}_2\text{O}$   
 Contributing reactions:  
 (1)  $\text{HC}_2\text{H}_3\text{O}_2 \rightleftharpoons \text{H}^+ + \text{C}_2\text{H}_3\text{O}_2^- \quad K_a = 1.8 \times 10^{-5}$   
 (2)  $\text{H}_2\text{O} \rightleftharpoons \text{H}^+ + \text{OH}^- \quad K_w = 1.0 \times 10^{-14}$

	$\text{HC}_2\text{H}_3\text{O}_2$	$\text{H}^+$	$\text{C}_2\text{H}_3\text{O}_2^-$
I	0.10	0	0
C	-x	+x	+x
E	0.10 - x	x	x

$$K_a = \frac{[\text{H}^+][\text{C}_2\text{H}_3\text{O}_2^-]}{[\text{HC}_2\text{H}_3\text{O}_2]}$$

$$1.8 \times 10^{-5} = \frac{(x^2)}{(0.10-x)} \quad \leftarrow \frac{0.10}{K_a} \gg 100$$

$$1.8 \times 10^{-5} \cong \frac{(x^2)}{(0.10)}$$

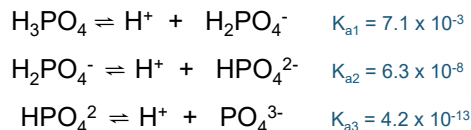
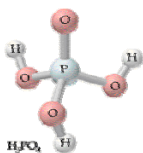
$$x^2 \cong 1.8 \times 10^{-6}$$

$$x \cong 1.3 \times 10^{-3} \text{ mol/L} \quad \leftarrow \frac{x}{0.10} = 1.3\% \text{ Assumption was valid!}$$

- Use  $[\text{H}^+]$  to find pH  
 $\text{pH} = -\log [\text{H}^+]$   
 $\text{pH} = -\log (1.3 \times 10^{-3}) = \mathbf{1.87}$

### Polyprotic acids have more than one ionisable proton.

e.g., phosphoric acid,  $\text{H}_3\text{PO}_4$



Only the **FIRST** ionization step contributes a considerable amount of  $\text{H}^+$ .

Treat polyprotic acid problems like monoprotic acids, where you only consider the first step!

What trend do you notice for the value of each successive  $K_a$ ?

**Example 5.** Calculate the pH a 0.20 mol/L solution of ascorbic acid,  $\text{H}_2\text{C}_6\text{H}_6\text{O}_6$ .  
 $K_{a1} = 7.9 \times 10^{-5}$   
 $K_{a2} = 1.6 \times 10^{-12}$

- Compare the  $K_a$  values:  
 Does the second ionization contribute much  $\text{H}^+$ ?  
 $K_{a2} \ll K_{a1}$   
 Only the **first step** contributes an appreciable amount of  $\text{H}^+$
- Write the ionization equation  

$$\text{H}_2\text{C}_6\text{H}_6\text{O}_6 \rightleftharpoons \text{H}^+ + \text{HC}_6\text{H}_6\text{O}_6^-$$
- Consider all major entities, and reactions that contribute to pH.  
 Major entities in solution:  
 $\text{H}_2\text{C}_6\text{H}_6\text{O}_6 \quad \text{H}^+ \quad \text{HC}_6\text{H}_6\text{O}_6^- \quad \text{H}_2\text{O}$   
 Contributing reactions:  
 (1)  $\text{H}_2\text{C}_6\text{H}_6\text{O}_6 \rightleftharpoons \text{H}^+ + \text{HC}_6\text{H}_6\text{O}_6^- \quad K_a = 7.9 \times 10^{-5}$   
 (2)  $\text{H}_2\text{O} \rightleftharpoons \text{H}^+ + \text{OH}^- \quad K_w = 1.0 \times 10^{-14}$

- Set up ICE table

	$\text{H}_2\text{C}_6\text{H}_6\text{O}_6$	$\text{H}^+$	$\text{HC}_6\text{H}_6\text{O}_6^-$
I	0.20	0	0
C	-x	+x	+x
E	0.20 - x	x	x

- Use  $K_a$  to solve for x

$$K_a = \frac{[\text{H}^+][\text{HC}_6\text{H}_6\text{O}_6^-]}{[\text{H}_2\text{C}_6\text{H}_6\text{O}_6]}$$

$$7.9 \times 10^{-5} = \frac{(x^2)}{(0.20-x)} \cong \frac{(x^2)}{(0.20)} \quad \leftarrow \frac{0.20}{K_a} \gg 100$$

$$x \cong 4.0 \times 10^{-3} \text{ mol/L} \quad \leftarrow \frac{x}{0.20} = 2\% \text{ Assumption was valid!}$$

- Use  $[\text{H}^+]$  to find pH

$$\text{pH} = -\log [\text{H}^+]$$

$$\text{pH} = -\log (4.0 \times 10^{-3}) = \mathbf{2.40}$$

## Summary

- Strong acids ionize completely, so  $[\text{H}^+]$  can be obtained directly from  $[\text{HA}]$ .
- For weak acids, % ionization describes the percentage of the original acid that ionizes to produce  $\text{H}^+$ .
- Weak acid problems are equilibrium problems. Set up an ICE table, and apply simplifying assumptions where appropriate.
- Polyprotic acids can be treated as weak monoprotic acids, since the contribution of the second/third ionizations is so miniscule.

## Homework

Pg. 516 #2

Pg. 520 #1, 2

Pg. 521 #1, 2

Pg. 524 #1, 2

Pg. 525 #1-10a