

Lecture 21 CH102 A1 (MWF 9:05 am) Spring 2019 Copyright © 2019 Dan Dill dan@bu.edu

[TP] At 25 °C, titration of 1.0 L of 0.010 M weak acid with OH^- has $\text{pH} = 8.5$ at the equivalence point, whereas titration of 1.0 L of 0.010 M strong acid with OH^- has $\text{pH} = 7.0$ at the equivalence point. This means that, compared to the moles of OH^- needed to reach equivalence in the weak acid titration, the moles of OH^- needed in the strong acid titration must be ...

25% 1. less
 25% 2. the same
 25% 3. more
 25% 4. Further information needed

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 Wednesday, March 20, 2019

- Titration: Adding successive amounts of base

Next: Buffers: $[\text{H}_3\text{O}^+]$ when different amounts of “not enough” base added;
 Practice: Too little, just enough, too much?; Ch15: Solubility, precipitation, and complexation

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Titration: Adding successive amounts of base

When successive amounts of base added, for each addition carry out step 1 by “going back to the beginning.”

Say 0.02 mol of OH^- is added to 0.10 mol of weak acid HA.

Then an additional 0.01 mol of OH^- is added.

The net result is 0.03 mol of OH^- added to 0.10 mol of weak acid.

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Titration of weak acid with hydroxide

Let's see how this works “titrating” 1.0 L of 0.01 M HA, $K_a = 1.0 \times 10^{-5}$, at 25 °C.

Calculate $[\text{H}_3\text{O}^+]$ and pH.

$[\text{H}_3\text{O}^+] = 3.2 \times 10^{-4}$ and $\text{pH} = 3.5$

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Titration of weak acid with hydroxide

Let's see how this works "titrating" 1.0 L of 0.01 M HA, $K_a = 1.0 \times 10^{-5}$, at 25 °C.

Now, **add 0.004 mol of OH^-** , and assume no volume change.

Calculate $[\text{H}_3\text{O}^+]$ and pH.

$[\text{H}_3\text{O}^+] = 1.5 \times 10^{-5}$ and pH = 4.8



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Titration of weak acid with hydroxide

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Now, **add another 0.001 mol of OH^-** , and assume no volume change.

Calculate $[\text{H}_3\text{O}^+]$ and pH.

$[\text{H}_3\text{O}^+] = 1.0 \times 10^{-5}$ and pH = 5.0 = **pKa! (half-equivalence point)**



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Titration of weak acid with hydroxide

Let's see how this works "titrating" 1.0 L of 0.01 M HA, $K_a = 1.0 \times 10^{-5}$, at 25 °C.

Now, **add another 0.001 mol of OH^-** , and assume no volume change.

Calculate $[\text{H}_3\text{O}^+]$ and pH.

$[\text{H}_3\text{O}^+] = 6.7 \times 10^{-6}$ and pH = 5.2



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Titration of weak acid with hydroxide

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Now, **add another 0.004 mol of OH^-** , and assume no volume change.

Calculate $[\text{H}_3\text{O}^+]$ and pH.

$[\text{H}_3\text{O}^+] = 3.2 \times 10^{-9}$ and pH = 8.5 (**equivalence point**)



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Titration of weak acid with hydroxide

Let's see how this works "titrating" 1.0 L of 0.01 M HA, $K_a = 1.0 \times 10^{-5}$, at 25 °C.

Now, **add another 0.001 mol of OH^-** , and assume no volume change.

Calculate $[\text{H}_3\text{O}^+]$ and pH.

$[\text{H}_3\text{O}^+] = 1.0 \times 10^{-11}$ and pH = 11.0 (**excess base**)



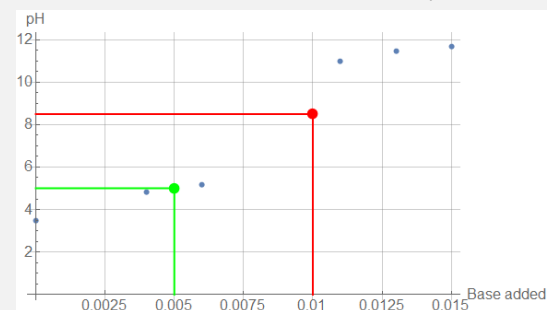
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Titration of weak acid with hydroxide

Let's see how this works "titrating" 1.0 L of 0.01 M HA, $K_a = 1.0 \times 10^{-5}$, at 25 °C.



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Titration of **strong** acid with hydroxide

Let's see how this works "titrating" 1.0 L of 0.01 M HA, $K_a = 1.0 \times 10^{+5}$, at 25 °C.

Calculate $[\text{H}_3\text{O}^+]$ and pH.

$[\text{H}_3\text{O}^+] = 1.0 \times 10^{-2}$ and pH = 2



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Titration of **strong** acid with hydroxide

Let's see how this works "titrating" 1.0 L of 0.01 M HA, $K_a = 1.0 \times 10^{+5}$, at 25 °C.

Now, **add 0.004 mol of OH^-** , and assume no volume change.

Calculate $[\text{H}_3\text{O}^+]$ and pH.



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Titration of **strong** acid with hydroxide

Add 0.004 mol of OH^- to 1.0 L of 0.01 M HA, $K_a = 1.0 \times 10^{+5}$, at 25 °C, assuming no volume change.

Step 1 results in 0.006 mol HA and 0.004 mol A^- .

Step 2

	[HA]	$[\text{H}_3\text{O}^+]$	$[\text{A}^-]$	Q
Initial	0.006/V	1×10^{-7}	0.004/V	$\ll K_a$
Change				
Equilibrium				



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Titration of **strong** acid with hydroxide

Add 0.004 mol of OH^- to 1.0 L of 0.01 M HA, $K_a = 1.0 \times 10^{+5}$, at 25 °C, assuming no volume change.

Step 1 results in 0.006 mol HA and 0.004 mol A^- .

Step 2

	[HA]	$[\text{H}_3\text{O}^+]$	$[\text{A}^-]$	Q
Initial	0.006/V	1×10^{-7}	0.004/V	$\ll K_a$
Change	$\approx -0.006/V$	$\approx +0.006/V$	$\approx +0.006/V$	
Equilibrium				



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Titration of **strong** acid with hydroxide

Add 0.004 mol of OH^- to 1.0 L of 0.01 M HA, $K_a = 1.0 \times 10^{+5}$, at 25 °C, assuming no volume change.

Step 1 results in 0.006 mol HA and 0.004 mol A^- .

Step 2

	[HA]	$[\text{H}_3\text{O}^+]$	$[\text{A}^-]$	Q
Initial	0.006/V	1×10^{-7}	0.006/V	$\ll K_a$
Change	$\approx -0.006/V$	$\approx +0.006/V$	$\approx +0.006/V$	
Equilibrium	≈ 0	$\approx 0.006/V$	$\approx 0.010/V$	$\approx K_a$

$[\text{H}_3\text{O}^+] = 6.0 \times 10^{-3}$ and pH = 2.2



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Titration of **strong** acid with hydroxide

Let's see how this works "titrating" 1.0 L of 0.01 M HA, $K_a = 1.0 \times 10^{+5}$, at 25 °C.

Now, **add another 0.001 mol of OH^-** , and assume no volume change.

Calculate $[\text{H}_3\text{O}^+]$ and pH.

$[\text{H}_3\text{O}^+] = 5.0 \times 10^{-3}$ and pH = 2.3 (**half-equivalence point**)



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Titration of strong acid with hydroxide

Let's see how this works "titrating" 1.0 L of 0.01 M HA, $K_a = 1.0 \times 10^{+5}$, at 25 °C.

Now, add another 0.001 mol of OH^- , and assume no volume change.

Calculate $[\text{H}_3\text{O}^+]$ and pH.

$[\text{H}_3\text{O}^+] = 4.0 \times 10^{-3}$ and pH = 2.4



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Titration of strong acid with hydroxide

Let's see how this works "titrating" 1.0 L of 0.01 M HA, $K_a = 1.0 \times 10^{+5}$, at 25 °C.

Now, add another 0.004 mol of OH^- , and assume no volume change.

Calculate $[\text{H}_3\text{O}^+]$ and pH.



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Strong acid titration: Equivalence point

Add 0.010 mol of OH^- to 1.0 L of 0.01 M HA, $K_a = 1.0 \times 10^{+5}$, at 25 °C, assuming no volume change.

Step 1 results in 0.000 mol HA and 0.010 mol A^- .

Step 2, since $K_b = K_w/K_a \ll K_w$...

	$[\text{A}^-]$	$[\text{HA}]$	$[\text{OH}^-]$	Q
Initial	0.010/V	0	1×10^{-7}	$\ll K_b$
Change				
Final				



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Strong acid titration: Equivalence point

Add 0.010 mol of OH^- to 1.0 L of 0.01 M HA, $K_a = 1.0 \times 10^{+5}$, at 25 °C, assuming no volume change.

Step 1 results in 0.000 mol HA and 0.010 mol A^- .

Step 2, since $K_b = K_w/K_a \ll K_w$...

	$[\text{A}^-]$	$[\text{HA}]$	$[\text{OH}^-]$	Q
Initial	0.010/V	0	1×10^{-7}	$0 \ll K_b$
Change	$-x$	$+x$	$+x$	
Final				



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Strong acid titration: Equivalence point

Add 0.010 mol of OH^- to 1.0 L of 0.01 M HA, $K_a = 1.0 \times 10^{+5}$, at 25 °C, assuming no volume change.

Step 1 results in 0.000 mol HA and 0.010 mol A^- .

Step 2, since $K_b = K_w/K_a \ll K_w$...

	$[\text{A}^-]$	$[\text{HA}]$	$[\text{OH}^-]$	Q
Initial	$0.010/V$	0	1×10^{-7}	$\ll K_b$
Change	$-x$	$+x$	$+x$	
Equilibrium	$\approx 0.010/V$	x	$\approx 1 \times 10^{-7}$	$\approx K_b$

$[\text{H}_3\text{O}^+] = [\text{OH}^-] = 1.0 \times 10^{-7}$ and pH = 7.0 (equivalence point)



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Titration of strong acid with hydroxide

Let's see how this works "titrating" 1.0 L of 0.01 M HA, $K_a = 1.0 \times 10^{+5}$, at 25 °C.

Now, add another 0.001 mol of OH^- , and assume no volume change.

Calculate $[\text{H}_3\text{O}^+]$ and pH.



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Strong acid titration: Excess base

Add 0.011 mol of OH^- to 1.0 L of 0.01 M HA, $K_a = 1.0 \times 10^{+5}$, at 25 °C, assuming no volume change.

Step 1 results in 0.000 mol HA, 0.001 mol OH^- , and 0.010 mol A^- .

Step 2, since $K_b = K_w/K_a \ll K_w$...

	$[\text{A}^-]$	$[\text{HA}]$	$[\text{OH}^-]$	Q
Initial	$0.010/V$	0	$0.001/V$	$\ll K_b$
Change	$-x$	$+x$	$+x$	
Equilibrium	$\approx 0.010/V$	x	$\approx 0.001/V$	$\approx K_b$

$[\text{H}_3\text{O}^+] = \frac{K_w}{0.001/V} = 1.0 \times 10^{-11}$ and pH = 11.0 (excess base)



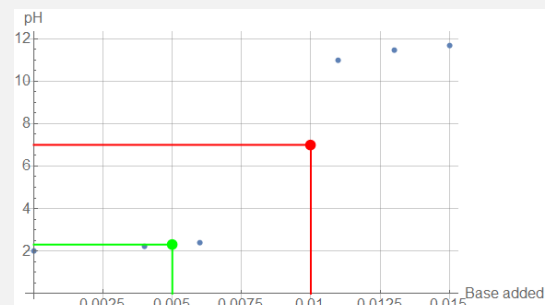
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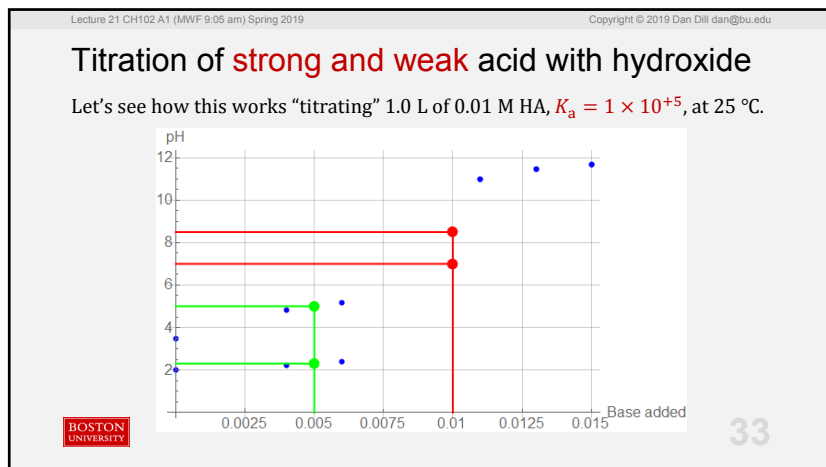
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Titration of strong acid with hydroxide

Let's see how this works "titrating" 1.0 L of 0.01 M HA, $K_a = 1 \times 10^{+5}$, at 25 °C.



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[TP] At 25 °C, titration of 1.0 L of 0.010 M weak acid with OH^- has $\text{pH} = 8.5$ at the equivalence point, whereas titration of 1.0 L of 0.010 M strong acid with OH^- has $\text{pH} = 7.0$ at the equivalence point. This means that, compared to the moles of OH^- needed to reach equivalence in the weak acid titration, the moles of OH^- needed in the strong acid titration must be ...

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