## This is BCE#23.

I recommend you print out this page and bring it to class. <u>Click here</u> to show a set of five BCE23 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.

Consider a titration where 0.250 M NaOH is added to 50.0 mL of 0.400 M HC<sub>2</sub>H<sub>3</sub>O<sub>2</sub>.

**1.** How many moles of HC<sub>2</sub>H<sub>3</sub>O<sub>2</sub> are in the original **50.0** mL sample?

0.0200 mol 76% 12% 0.400 M for moles.

 $(0.0500 \text{ L})^*(0.400 \text{ mol}/1 \text{ L}) = 0.0200 \text{ mol} \text{ HC}_2\text{H}_3\text{O}_2$ 

2. Calculate the volume required to reach the equivalence point when 0.250 M NaOH is added to 50.0 mL of 0.400 M  $HC_2H_3O_2$ .

80.0 mL 65%

At the equivalence point the moles of  $HC_2H_3O_2$  are equal to the moles of NaOH. In Q1 we calculated the moles of  $HC_2H_3O_2$  to be 0.0200 mol. So we must add 0.0200 moles of NaOH to reach the equivalence point. The volume is,

0.0200 mol NaOH\*(1 L/0.250 mol) = 0.0800 L (80.0 mL)

3. What volume of 0.250 M NaOH is required to react with exactly one half of the moles of  $HC_2H_3O_2$ ?

40.0 mL 53%

80.0 mL/2 = 40.0 mL

4. Calculate  $[H^+]$  concentration when exactly half of the  $HC_2H_3O_2$  has been neutralized by the NaOH.

[H<sup>+</sup>] = 1.8e-5 M

Reworded this question is asking you to calculate the  $[H^+]$  when 40.0 mL of 0.250 M NaOH are added to 50.0 mL of 0.400 M HC<sub>2</sub>H<sub>3</sub>O<sub>2</sub>. To solve this we must write the neutralization equation and set up the IF table.

 $mol HC_2H_3O_2 = 0.0500 L * (0.400 mol/1 L) = 0.0200 mol$ 

mol NaOH= 0.0400 L \*(0.250 mol/1 L) = 0.0100 mol

	HC <sub>2</sub> H <sub>3</sub> O <sub>2</sub> (aq) +	NaOH(l)	₽	$C_2H_3O_2^-(aq) +$	H <sub>2</sub> O(l)
Ι	0.0200 mol	0.0100 mol		0	-
F	0.0100 mol	0		0.0100 mol	-

After completing the IF table we look at the Final row to determine what type of system we have. Looking at the Final row we have 0.0100 mol of a weak acid and 0.0100 moles of its conjugate base. This is a common ion system. So now we must set up an ICE table to calculate the [H<sup>+</sup>] for this common ion system.

We need to calculate the  $[HC_2H_3O_2]$  and the  $[C_2H_3O_2^-]$  first.

 $[HC_2H_3O_2] = 0.0100 \text{ mol}/0.0900 \text{ L} = 0.111 \text{ M}$ 

 $[C_2H_3O_2] = 0.0100 \text{ mol}/0.0900 \text{ L} = 0.111 \text{ M}$ 

	HC <sub>2</sub> H <sub>3</sub> O <sub>2</sub> (aq) +	₹	$C_2H_3O_2(aq) +$	H <sup>+</sup> (aq)
Ι	0.111		0.111	~0
С	-X		+x	+x

I	E	<b>0.111 - x</b>	(	<b>).111 + x</b>	+x

 $K_a = [C_2H_3O_2^-][H^+]/[HC_2H_3O_2]$ 

 $1.75 \ge 10^{-5} = [0.111 + x][x]/[0.111 - x]$ 

assume 0.111 - x = 0.111

 $1.75 \ge 10^{-5} = [0.111][x]/[0.111]$ 

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1.75 \ge 10^{-5} = [x] = [H^+]
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5. What is interesting about the concentration of [H<sup>+</sup>] in Q4?

The [H<sup>+</sup>] at the half-equivalence point is the same as the magnitude of the equilibrium constant.

At the half equivalence point the  $[H^+] = K_a$  for the weak acid.....cool!

6. Is there anything about the questions that you feel you do not understand? List your concerns/questions.

## nothing

7. If there is one question you would like to have answered in lecture, what would that question be?

## nothing