

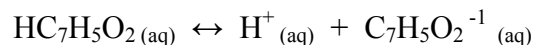
AP CHEMISTRY

TOPIC 7: ACIDS & BASES, PART E, PART II

Day 88:

- Buffers

CLEARLY SHOW THE METHOD USED AND THE STEPS INVOLVED IN ARRIVING AT YOUR ANSWERS.



1. Benzoic acid, $\text{HC}_7\text{H}_5\text{O}_2$, is a monoprotic acid that dissociates in an aqueous solution, as represented in the equation above. Benzoic acid is 3.29 percent dissociated in 0.85 M $\text{HC}_7\text{H}_5\text{O}_2(\text{aq})$ at 298 K. For parts (a) through (d) below, assume the temperature remains at 298 K.

- (a) Write the expression for the acid-dissociation constant, K_a , for benzoic acid and calculate its value.

$$K_a = \frac{[\text{H}^+][\text{C}_7\text{H}_5\text{O}_2^{-1}]}{[\text{HC}_7\text{H}_5\text{O}_2]}$$

$$0.85 \text{ M} \times 0.0329 = 0.027965 \text{ M} = x$$

	[$\text{HC}_7\text{H}_5\text{O}_2$]	\leftrightarrow	[H^+]	+	[$\text{C}_7\text{H}_5\text{O}_2^{-1}$]
I	0.85 M		0		0
C	0.85 - 0.027965		+ 0.027965		+ 0.027965
E	0.822035		0.027965		0.027965

$$K_a = \frac{(0.027965)^2}{0.822035} = 9.51 \times 10^{-4}$$

- (b) Calculate the pH of 0.85 M $\text{HC}_7\text{H}_5\text{O}_2$.

$$\text{pH} = -\log [\text{H}^+] = -\log (0.027965) = \mathbf{1.55}$$

- (c) Calculate the pH of a solution formed by dissolving 0.067 mol of solid sodium benzoate, $\text{NaC}_7\text{H}_5\text{O}_2$. in 250 mL of 0.85 M $\text{HC}_7\text{H}_5\text{O}_2$. Assume the volume changes is negligible.

$$M = \frac{0.067 \text{ mol NaC}_7\text{H}_5\text{O}_2}{0.250 \text{ L}} = 0.268 \text{ M NaC}_7\text{H}_5\text{O}_2$$



	$[\text{HC}_7\text{H}_5\text{O}_2]$	\leftrightarrow	$[\text{H}^{+1}]$	+	$[\text{C}_7\text{H}_5\text{O}_2^{-1}]$
I	0.85 M		~ 0		0.268 M
C	+ x		- x		- x
E	0.85 + x		$[\text{H}^{+1}]$		0.268 - x

$$K_a = \frac{[\text{H}^{+1}][\text{C}_7\text{H}_5\text{O}_2^{-1}]}{[\text{HC}_7\text{H}_5\text{O}_2]} = \frac{[\text{H}^{+1}](0.268 - x)}{(0.85 + x)} = \frac{[\text{H}^{+1}](0.268)}{(0.85)} = 9.51 \times 10^{-4}$$

$$[\text{H}^{+1}] = \frac{(9.51 \times 10^{-4})(0.85)}{(0.268)} = 3.02 \times 10^{-3}$$

$$\text{pH} = -\log [\text{H}^{+1}] = -\log (3.02 \times 10^{-3}) = \mathbf{2.52}$$

OR

$$\text{pH} = pK_a + \log \frac{[\text{A}^{-1}]}{[\text{HA}]} = -\log(9.51 \times 10^{-4}) + \log\left(\frac{0.268}{0.85}\right) = 2.52$$

- (d) 5.33 grams of solid HCl is added to 250 mL of the solution created in part (c). Calculate the pH, in the resulting solution.

$$\frac{5.33 \text{ g HCl}}{36.458 \text{ g}} \times \frac{1 \text{ mol HCl}}{1} = 0.146 \text{ mol HCl}$$

$$\frac{0.146 \text{ mol HCl}}{0.250 \text{ L}} = 0.585 \text{ M HCl}$$

	[HC ₇ H ₅ O ₂]	←	[H ⁺]	+	[C ₇ H ₅ O ₂ ⁻¹]
I	0.850 M		0.585 M		0.268 M
C	+ 0.268		- 0.268		- 0.268
E	0.850 + 0.268 = 1.118		0.585 - 0.268 = 0.317		~ 0

This is not a “good” buffer solution.

The solution could not handle the additional H⁺ ions (from the strong acid). Since this solution is unable to handle the additional strong acid, we KNOW the concentration of the H⁺ at equilibrium. Therefore:

$$\mathbf{pH} = -\log [\text{H}^{+}] = -\log (0.317) = \mathbf{0.50}$$

2. A buffered solution (at equilibrium) contains 3.75 M acetic acid, ($\text{HC}_2\text{H}_3\text{O}_2$, $K_a = 1.80 \times 10^{-5}$) and 2.43 M potassium acetate, $\text{KC}_2\text{H}_3\text{O}_2$.

a) Calculate the pH of this solution.



	$[\text{HC}_2\text{H}_3\text{O}_2]$	\leftrightarrow	$[\text{H}^+]$	+	$[\text{C}_2\text{H}_3\text{O}_2^{-1}]$
E	3.75 M		~ 0		2.43 M

$$K_a = \frac{[\text{H}^+][\text{C}_2\text{H}_3\text{O}_2^{-1}]}{[\text{HC}_2\text{H}_3\text{O}_2]} = \frac{[\text{H}^+](2.43)}{(3.75)} = 1.80 \times 10^{-5}$$

$$[\text{H}^+] = \frac{(1.80 \times 10^{-5})(3.75)}{(2.43)} = 2.78 \times 10^{-5}$$

$$\text{pH} = -\log [\text{H}^+] = -\log (2.78 \times 10^{-5}) = \mathbf{4.56}$$

OR

$$\text{pH} = \text{p}K_a + \log \left[\frac{A^{-1}}{HA} \right] = -\log(1.80 \times 10^{-5}) + \log \left(\frac{2.43}{3.75} \right) = 4.56$$

b) Calculate the pH after 34.3 grams of $\text{NaOH}_{(s)}$ is added to 1.0 L of this solution (volume change is negligible).

$$\frac{34.3 \text{ g NaOH}}{39.998 \text{ g}} \times \frac{1 \text{ mol NaOH}}{1} = 0.858 \text{ mol NaOH}$$

$$\frac{0.858 \text{ mol NaOH}}{1 \text{ L}} = 0.858 \text{ M NaOH}$$

	$[\text{C}_2\text{H}_3\text{O}_2^{-1}]$	+	$[\text{H}_2\text{O}]$	\leftarrow	$[\text{OH}^{-1}]$	+	$[\text{HC}_2\text{H}_3\text{O}_2]$
I	2.43 M		-		0.858 M		3.75 M
C	+ 0.858		-		- 0.858		- 0.858
E	3.288		-		~ 0		2.892

$$\text{pH} = \text{p}K_a + \log \left[\frac{A^{-1}}{HA} \right] = -\log(1.80 \times 10^{-5}) + \log \left(\frac{3.288}{2.892} \right) = 4.80$$

OR

$$K_b = \frac{K_w}{K_a} = \frac{1.00 \times 10^{-14}}{1.80 \times 10^{-5}} = 5.56 \times 10^{-10}$$

$$K_b = \frac{[OH^{-1}][HC_2H_3O_2]}{[C_2H_3O_2^{-1}]} = \frac{[OH^{-1}](2.892)}{(3.288)} = 5.56 \times 10^{-10}$$

$$[OH^{-1}] = \frac{(5.56 \times 10^{-10})(3.288)}{(2.892)} = 6.32 \times 10^{-10}$$

$$pOH = -\log [OH^{-1}] = -\log (6.32 \times 10^{-10}) = 9.20$$

$$pH = 14 - pOH = 14 - 9.20 = \mathbf{4.80}$$